National Institute for Health and Care Excellence

Draft

Head injury: assessment and early management (update)

[H] Evidence review for CT, MRI and X-ray of the cervical spine in people with head injury – diagnostic

NICE guideline <number>

Evidence reviews underpinning recommendations x to y and research recommendations in the NICE guideline

September 2022

Draft for consultation

These evidence reviews were developed by the Guideline Development Team NGC



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1 CT, MRI and X-ray of the cervical spine in people with head injury – diagnostic

3 1.1 Review question

- What is the diagnostic accuracy of CT, MRI and X-ray of the cervical spine for initial imaging in people with head injury?
 - What is the clinical and cost effectiveness of CT, MRI and X-ray of the cervical spine for initial imaging in people with head injury?

8 1.1.1 Introduction

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9 Patients with head injury may sustain bony and/or soft tissue injuries to the cervical spine. 10 When imaging is required, the 2014 version of the NICE guideline recommended as the first line test either a series of cervical spine X-rays or a Computerised Tomography (CT) scan. 11 Depending on the clinical situation, Magnetic Resonance (MR) was also indicated in some 12 13 cases to determine injury to the ligamentous structures, intervertebral discs and spinal cord at both the cranio-cervical junction and the sub-axial cervical spine. This review includes new 14 evidence published since the last update of the guideline on the diagnostic accuracy and 15 16 clinical and cost effectiveness of CT, MRI and X-ray of the cervical spine for initial imaging in people with head injury. 17

18 **1.1.2 Summary of the protocol**

19 For full details see the review protocol in Appendix A.

20 Table 1: PICO characteristics of review question – diagnostic accuracy

Population	Infants, children and adults with head injury and suspected cervical spine injury				
	<u>Strata: C</u> -spine injury risk stratification (based on Canadian C-Spine Rule or NEXUS – two stratifications are different so to be kept separate)				
	Adults (≥16 years) at:Children + infants (0-16 years) at:• high risk• high risk• moderate risk• moderate risk				
	low risk low risk				
	Exclusion: adults and children (including infants <1 year) with superficial injuries to the eye or face without suspected or confirmed head or brain injury.				
	Cut-off of 60% will be used for assigning to strata for all age groups.				
Target condition	Cervical spine injury in patients who have experienced a head injury				
Index tests	 Computed tomography (CT) scan of cervical spine Magnetic resonance imaging (MRI) of cervical spine X-ray of cervical spine 				
Reference	Reference standard for CT:				
standards	CT and MR imaging of cervical spine Or				
	 2 weeks follow-up after CT including autopsy findings 				
	Reference standard for MR imaging:				
	CT and MR imaging of cervical spine				

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	 Or 2 weeks follow-up after MR imaging including autopsy findings
	Reference standard for X-ray:
	CT or MR imaging of cervical spine
	Or CT and MRI imaging of cervical spine
	Or
	2 weeks follow-up after X-ray including autopsy findings
	For X-ray only include children and people below 65 years.
	People >65 years are considered as high risk and will be offered CT cervical spine within 8 hours of injury (CG 176).
	Vascular injuries will be picked up by MR imaging
Statistical measures and outcomes	 Diagnostic accuracy CT, MRI and X-ray of the cervical spine for: any significant cervical spine injury
	(fracture/bony injury, soft tissue/ligament damage, spinal cord injuries, vascular injuries)
	No objective definition for significant cervical spine injury. Note definitions as reported in the papers.
	Diagnostic test accuracy to be reported by test sensitivity/specificity
	For measurement of imprecision, clinical decision thresholds for sensitivity and specificity are set at 90% and 60%.
	Sensitivity is considered to be more important than specificity. Sensitivity is more important as that will change management. Often, the decision is whether someone can be discharged from ED. A test with high sensitivity that is negative is very reassuring in ruling out an injury and allowing early discharge or mobilisation. It's unlikely that imaging will produce false positives.
Study design	Diagnostic cross-sectional studies, cohort studies (prospective and retrospective)
	Systematic reviews and meta-analyses of the above

1 Table 2: PICO characteristics of review question – diagnostic test and treat

Population	Infants, children and adults with head injury and suspected cervical spine injury
	<u>Strata:</u> C-spine injury risk stratification (based on Canadian C-Spine Rule or NEXUS – two stratifications are different so to be kept separate)
	Adults (≥16 years) at:Children + infants (0-16 years) at:• high risk• high risk• moderate risk• moderate risk• low risk• low risk
	Exclusion: adults and children (including infants <1 year) with superficial injuries to the eye or face without suspected or confirmed head or brain injury. Cut-off of 60% will be used for assigning to strata for all age groups.

Target condition	Cervical spine injury in patients who have experienced a head injury					
Interventions	 Computed tomography (CT) scan of cervical spine 					
	Magnetic resonance imaging (MRI) of cervical spine					
	X-ray of cervical spine					
Comparators	MRI of cervical spine, X-ray of cervical spine and CT of cervical spine compared to each other					
	For X-ray only include children and people below 65 years					
	People >65 years are considered as high risk and will be offered CT cervical spine within 8 hours of injury (CG 176)					
	Vascular injuries will be picked up by MR imaging					
Outcomes	All outcomes are considered equally important for decision making and therefore have all been rated as critical:					
	Mortality at 3 months					
	 Quality of life - 3 months or more 					
	 Objectively applied score of disability e.g. Glasgow Outcome Score (GOS) or extended GOS - at 3 months or more 					
	Length of hospital stay					
	 Unscheduled re-admission (28 days or longer) 					
	Neurological deterioration					
	Neurological deterioration could be because of either no imaging or no appropriate imaging					
	Spinal injuries are determined by different scales– e.g. American Spinal Injury Association (ASIA), functional independence measure (FIM). Different scales are used. Report as in the studies.					
	Vascular insult would be picked up in outcome neurological deterioration					
Study design	 Randomised controlled trials (RCTs), systematic reviews of RCTs. If no RCT evidence is available, non-randomised studies will be considered if they adjust for key confounders, starting with prospective cohort studies. 					
	Key confounders					
	Age					
	• Gender					
	GCS or pupillary response at presentation					

1 **1.1.3 Methods and process**

- 2 This evidence review was developed using the methods and process described in
- 3 <u>Developing NICE guidelines: the manual</u>. Methods specific to this review question are 4 described in the review protocol in appendix A and the methods document.
- 5 Declarations of interest were recorded according to <u>NICE's conflicts of interest policy</u>.
- 6

1 **1.1.4 Diagnostic evidence**

2 1.1.4.1 Included studies

3 Diagnostic accuracy

Forty-one studies reporting diagnostic accuracy data were included in the review; 1-7, 9, 10, 12-14, 4 ^{16-19, 21, 22, 25-31, 33, 35-44, 46-49} these are summarised in Tables 2 and 3 below. Evidence from 5 these studies is summarised in the clinical evidence summary below in Tables 9-23 and 6 references provided in 1.1.14 References . The assessment of the evidence quality was 7 conducted with emphasis on test sensitivity and specificity as this was identified by the 8 committee as the primary measure in guiding decision-making. Clinical decision thresholds of 9 sensitivity/specificity =0.9 and 0.60 above which a test would be recommended and 0.7 and 10 0.4 below which a test is of no clinical use were set by the committee. 11

12 Studies focusing on adults and children were reported separately. A total of 33 and 8 studies 13 were identified for adult and children populations, respectively. Some studies evaluated the 14 diagnostic accuracy of more than one diagnostic imaging modality. The number of studies 15 identified for each index test is given below

16

17	Adults

18	 X-ray as index test – 13 studies
19	 CT as index test – 16 studies
20	• CT and MRI as separate index tests – 9 studies
21	
22	
23	
24	Note that the bottom grouping was separated fro
25	studies where CT + MRI used as reference stand

Note that the bottom grouping was separated from other studies as these were studies where CT + MRI used as reference standard (or data available to analyse in this way), meaning only sensitivity values could be obtained (no information about specificity given both tests form part of the reference standard).

- 29 Children
- 30 X-ray as index test 3 studies
 - CT as index test 7 studies
- MRI as index test 6 studies
- 33

31

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27 28

34 **Population**

35 <u>Head injury</u>

For most studies identified in the literature search, head injury was not mentioned and the 36 37 population was described only as those with suspected cervical spine injury undergoing 38 imaging of the cervical spine. Despite it being unclear if head injury was present or the 39 proportion that had suffered head injury in these studies, these studies were included and 40 downgraded for indirectness, as it was noted that many of those with cervical spine injury are likely to have experienced head injury based on the nature of the injury, for example those 41 with whiplash are likely to have suffered a head injury as well. For studies where the 42 population was limited to those that were unconscious or obtunded, often requiring intensive 43 44 care unit admission, although head injury was not specifically mentioned in many studies, it was assumed that these groups did have at least suspected head injury given the severity of 45 46 their injuries; these studies were not downgraded based on head injury not being mentioned.

1 However, they were still downgraded for another reason: the severe nature of their injuries 2 makes them a very specific subgroup of the population that attend the emergency

3 department with suspected cervical spine injury. Results would be less applicable than those

of people who are discharged from the emergency department without admission to hospital 4

- 5 or intensive care.
- 6 Therefore, the population of included studies varies, with the following major groups 7 identified:
- 8 Studies where it is clearly stated or suggested (for example undergoing a head CT) 9 that all or a majority of patients sustained a head injury and underwent imaging of the 10 cervical spine
- 11 Studies where the presence of head injury is not mentioned and patients underwent • imaging of the cervical spine (for those that were obtunded or unconscious, based on 12 the severity of injuries head injury was assumed to have occurred) 13
- Studies where it is clear a proportion had some form of confirmed or suspected head 14 • injury but for the remaining patients it is unclear (for example, a proportion had head 15 CT or diagnosed with intracranial haemorrhage) and underwent imaging of the 16 17 cervical spine

18 The inclusion of studies where it was unclear if head injury was present or not meant that 19 there was overlap with an evidence review relating to cervical spine assessment performed as part of the NICE Spinal injury: assessment and initial management guideline: 20

- 21 For adults, 13 of the included studies had also been included in the spinal injury assessment and initial management NICE guideline 22
- For children, 3 of the included studies had also been included in the spinal injury 23 assessment and initial management NICE guideline 24

25

Other population details 26

27 Some studies included a broad population of patients with suspected cervical spine injury 28 undergoing imaging but others were more specific. For example, some studies only included those that were unconscious or those with severe traumatic injuries. This was taken into 29 account when deciding whether studies were similar enough to be grouped together. 30

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- 32 For adults, four main groupings were identified, which are presented in separate included studies tables in section 1.1.5 and separate GRADE tables in section 1.1.6: 33
 - all having index test and not limited to those that were admitted
 - only including those admitted, not those subsequently discharged following index test •
- only including those that are obtunded, unconscious and/or requiring intensive care 36 • unit admission (X-ray was excluded as an index test in this group given X-ray would not be used as the initial imaging test in this population with severe injuries)
 - other very specific populations •

40

39

- 41 For children, two main groupings were identified, which are presented in separate included studies tables in section 1.1.5 and separate GRADE tables in section 1.1.6: 42
- 43 all having index test and not limited to those that were admitted
- 44 only including those that are obtunded, unconscious and/or requiring intensive care unit admission 45

1 **Reference standards**

Reference standards used varied across studies even for those using the same index test.
Broad groups of reference standards identified in studies were as follows, though the details
may differ between studies:

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> CT scan or MRI – accepted as a reference standard where X-ray was the index test. CT scan often used as the reference standard for bony/osseous injuries (for example, fracture) and MRI for ligamentous injuries. CT also accepted as a reference standard for studies that were assessing the ability of MRI to specifically detect fractures and MRI accepted as a reference standard for studies that were assessing the ability of the ability of CT to detect ligamentous injuries, as it was acknowledged that CT would be the reference standard for bony injuries and MRI for ligamentous/soft tissue injuries.

- CT + MRI combined some studies used CT and MRI combined as a reference standard, or provided data that enabled this to be worked out, meaning the sensitivity of CT and MRI individually could be calculated (no information about specificity could be obtained using this as a reference standard for CT and MRI individually as index tests)
- 24

25 Outcome definitions

Target conditions being detected by index tests varied across studies. The protocol for this review specified 'any significant cervical spine injury'.

28 Some studies did report only more serious or significant injuries, for example cervical spine 29 injuries that were defined as unstable or requiring intervention.

Some studies only reported 'any cervical spine injury', the definition of which varied across
 studies and was sometimes poorly defined. For studies where this was the only outcome
 reported, this was accepted and included in the analysis.

Some studies reported both significant injuries and any injuries – in this case the results for
 significant injuries were included in the analysis as this was more in line with the protocol.

A further way in which outcome definitions varied across studies was the types of injuries (for example, bony or ligamentous/soft tissue) that were included. Some studies included any type of injury in the outcome/target condition whereas others focused the study on specific

- types of injuries, for example only fractures or only ligamentous injuries.
- 39

40 <u>Pooling</u>

Given the wide variation discussed above in terms of population, reference standard and

42 outcome definitions, pooling of results was not appropriate. Studies that were broadly similar

- 43 in terms of index test, population, reference standard and target condition were grouped
- 44 under the same headings but not formally pooled.
- 45

46 **Diagnostic test and treat**

- 1 All included studies provided data for the diagnostic accuracy component of this question, as 2 no diagnostic test and treat studies matching the protocol were identified.
- 3
- 4 See also the study selection flow chart in Appendix C, sensitivity and specificity forest plots in 5 Appendix E, and study evidence tables in Appendix D.
- 6

7 1.1.4.2 Excluded studies

8 See the excluded studies list in Appendix I.

2 **1.1.5 Summary of studies included in the diagnostic evidence**

3 Table 3: Summary of studies included in the evidence review – adults – all having index test and not limited to those that were admitted

Study	Population	Target condition	Index test	Reference standard	Comments
X-ray as index test					
Bailitz 2009 ³ N=1505 Conducted in USA Prospective	Adults (≥16 years) meeting one or more of NEXUS criteria and requiring cervical spine imaging following trauma Unclear if all or most had head injury as no details provided	Clinically significant cervical spine injury	Cervical spine radiographs (X-ray)	Final diagnosis in medical record at discharge	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Unclear if reference standard included a 2-week follow-up period Included in spinal assessment guideline Risk stratification: meeting at least one NEXUS criterion – separate results for high,
Duane 2010 ¹³ N=49 Conducted in USA Retrospective	Adult patients (≥18 years) following blunt trauma who had flexion-extension plain films and MRI of cervical spine	Ligamentous injury of the cervical spine	X-ray – flexion- extension plain films	MRI – gold standard for ligamentous injuries	 moderate and low risk Indirectness: Head injury not mentioned so unclear if most or all had head injury Focuses only on ligamentous injuries

Study	Population	Target condition	Index test	Reference standard	Comments
	Unclear if all or most had head injury as no details provided				Included in spinal assessment guideline Risk stratification: unclear
Gale 2005 ¹⁹ N=400 Conducted in USA Retrospective	Blunt trauma patients undergoing head CT and also having plain radiography (X-ray) of cervical spine All included patients underwent head CT	Cervical spine fracture	Plain radiography (X- ray)	CT of cervical spine – gold standard for fractures	 Indirectness: Focuses only on fractures in the outcome Risk stratification: unclear
Gharekhanloo 2021 ²⁰ N=220 Conducted in Iran Prospective	Adult trauma patients referred to an ED in Iran. They received plain radiography and CT to evaluate cervical spine injury. Low risk status based on NEXUS criteria.	Cervical spine injury	Plain radiography (X- ray)	CT of cervical spine – gold standard for cervical spine injury	 Indirectness: Head injury not mentioned so unclear how many had head injury Only 10 people had abnormal CT. Risk stratification: unclear
Griffen 2003 ²² N=1199 Conducted in USA Retrospective	Adult blunt trauma patients undergoing cervical spine assessment by X ray and CT	Cervical spine injury – poorly defined	X-ray of cervical spine	Unclear, possibly all imaging/follow-up	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Unclear if reference standard included a

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Study	Population	Target condition	Index test	Reference standard	Comments
	Unclear if all or most had head injury as no details provided				2-week follow-up period Included in spinal assessment guideline Risk stratification: unclear
Lee 2001 ²⁹ N=604 Conducted in USA Retrospective	Adult trauma patients undergoing imaging examination of the cervical spine with conventional radiography and helical CT Unclear if all or most had head injury as no details provided	Fractures	Conventional radiography (X-ray)	Helical CT scan	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Focuses only on fractures Results reported at fracture-level not patient-level (patients could have more than one fracture and these included in analysis individually) Included in spinal assessment guideline Risk stratification: described as high index of suspicion for cervical spine injury (unclear which rule based on)
Mathen 2007 ³¹	Trauma patients (average age 35.4 years) not meeting	Clinically significant cervical spine injury – requiring surgery or	Plain films (X-ray)	Final diagnosis of cervical spine injury based on all	Indirectness:

Study	Population	Target condition	Index test	Reference standard	Comments
N=667 Conducted in USA Prospective	NEXUS low-risk criteria and undergoing CT and radiography of cervical spine Unclear if all or most had head injury as no details provided	long-term stabilisation with a collar or halo		prospectively collected clinical data and imaging results	 Head injury not mentioned so unclear if most or all had head injury Unclear if reference standard included a 2-week follow-up period Included in spinal assessment guideline Risk stratification: those not meeting NEXUS low risk criteria
Nguyen 2005 ³³ N=112 Conducted in USA Prospective	Patients with blunt trauma undergoing imaging of cervical spine Unclear if all or most had head injury as no details provided	Cervical spine fractures	X-ray	Diagnosis based on final reports including all imaging	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Focuses only on fractures Unclear if reference standard included a 2-week follow-up period Risk stratification: reports data for low and high risk separately, based on NEXUS

Study	Population	Target condition	Index test	Reference standard	Comments
Takami 2014 ⁴⁶ N=179 Conducted in Japan Prospective	Patients sustaining high-energy trauma immobilised and undergoing X-ray and CT of cervical spine Proportion had concomitant head injury but unclear how many, reported to be 15% in those with fractures	Cervical spine fracture	X-ray of cervical spine	Full CT of spine	 Indirectness: Head injury present in a small proportion but unclear if remaining had head injury as part of the injury mechanism Focuses only on fractures Included in spinal assessment guideline Risk stratification: unclear, those with high- energy trauma
CT as index test					
Bailitz 2009 ³ N=1505 Conducted in USA Prospective	Adults (≥16 years) meeting one or more of NEXUS criteria and requiring cervical spine imaging following trauma Unclear if all or most had head injury as no details provided	Clinically significant cervical spine injury	CT of cervical spine	Final diagnosis in medical record at discharge	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Unclear if reference standard included a 2-week follow-up period Included in spinal assessment guideline Risk stratification: meeting at least one

Study	Population	Target condition	Index test	Reference standard	Comments
					NEXUS criterion – separate results for high, moderate and low risk
Duane 2016 ¹⁶ N=9227 Conducted in USA Retrospective	Adults (≥18 years) following trauma and undergoing assessment of cervical spine Unclear if most or all had head injury as no details provided	Fracture and/or ligamentous injury	CT scan	Later found to have cervical spine injury – poorly defined. Possibly includes any report of injury during follow-up and also results of any additional imaging performed (e.g. MRI). Likely that reference standard differs between patients and unclear follow-up duration	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Reference standard poorly defined and unclear if matches protocol Risk stratification: unclear, described as patients with criteria for trauma team alert
Griffen 2003 ²² N=1199 Conducted in USA Retrospective	Adult blunt trauma patients undergoing cervical spine assessment by X ray and CT Unclear if all or most had head injury as no details provided	Cervical spine injury – poorly defined	CT of cervical spine	Unclear, possibly all imaging/follow-up	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Unclear if reference standard included a 2-week follow-up period Included in spinal assessment guideline Risk stratification: unclear

Study	Population	Target condition	Index test	Reference standard	Comments
Inaba 2016 ²⁷ N=10,276 Conducted in USA Prospective	Adults (≥18 years) following blunt trauma undergoing CT scan of the cervical spine (failed NEXUS low- risk criteria) Unclear if all or most had head injury as no details provided	Clinically significant cervical spine fracture Defined as abnormal or equivocal finding on CT or MRI consistent with acute traumatic injury along with one of three actives interventions: surgical stabilisation, Halo Orthotic placement or use of Cervical- Thoracic Orthotic	CT of cervical spine	Final diagnosis at time of discharge, including any additional imaging and operative findings dependent on each patient	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Limits only to cervical spine fractures Reference standard does not include a 2 week follow-up Risk stratification: patients failing low-risk NEXUS criteria
Mathen 2007 ³¹ N=667 Conducted in USA Prospective	Trauma patients (average age 35.4 years) not meeting NEXUS low-risk criteria and undergoing CT and radiography of cervical spine Unclear if all or most had head injury as no details provided	Clinically significant cervical spine injury – requiring surgery or long-term stabilisation with a collar or halo	Multi-slice CT of cervical spine	Final diagnosis of cervical spine injury based on all prospectively collected clinical data and imaging results	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Unclear if reference standard included a 2-week follow-up period Included in spinal assessment guideline Risk stratification: those not meeting NEXUS low risk criteria

Study	Population	Target condition	Index test	Reference standard	Comments
Nguyen 2005 ³³ N=112 analysed by CT Conducted in USA Prospective	Patients with blunt trauma undergoing imaging of cervical spine Unclear if all or most had head injury as no details provided	Cervical spine fractures	CT of cervical spine	Diagnosis based on final reports including all imaging	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Focuses only on fractures Unclear if reference standard included a 2-week follow-up period Risk stratification: reports data for low and high risk separately, based on NEXUS
Ptak 2001 ³⁷ N=676 Conducted in USA Retrospective	Patients (mean age 47.2 years) presenting to emergency radiology division for cervical spine injury evaluation following trauma by CT Unclear if all or most had head injury as no details provided	Cervical spine fracture	CT of cervical spine	Final clinical diagnosis (including operative and discharge), possibly incorporating CT results	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Focuses only on fractures Unclear if reference standard included a 2-week follow-up period Included in spinal assessment guideline

Study	Population	Target condition	Index test	Reference standard	Comments
					Risk stratification: unclear
Vanguri 2014 ⁴⁸ N=5676 Conducted in USA Retrospective	Adult blunt trauma undergoing cervical spine assessment by CT Unclear if all or most had head injury as no details provided	Cervical spine injury – poorly defined	Cervical spine CT	Unclear, possibly including other imaging such as MRI and flexion-extension depending on patient	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Unclear if reference standard included a 2-week follow-up period Risk stratification: unclear, those meeting criteria for trauma team activation
CT alone and MRI a of CT and MRI sepa	•	ests – CT and MRI combir	ned used as reference sta	ndard (only sensitivity cou	uld be calculated for each
Friesen 2014 ¹⁸ N=206 analysed Conducted in Australia Retrospective	Adults (≥16 years) with CT and MRI performed for suspected blunt acute cervical spine trauma Likely most had a suspicion of head injury as 76% had combined cervical spine and brain CT	Unstable cervical spine injury Defined by Denis 3 column definition as well as any cases requiring urgent (within 5 days) surgery or urgent surgical immobilisation (such as halo-traction ring) and following additional injuries: flexion teardrop fracture,	Helical CT of cervical spine OR MRI of cervical spine	MRI said to be reference standard for unstable injuries in the study which is not the case according to our protocol Data presented in paper therefore analysed using combined CT and MRI as reference standard	Reference standard of MRI alone does not match protocol, but available data analysed using CT and MRI as a combined reference standard (meaning only sensitivity could be calculated, not specificity) Risk stratification: 98% met at least one NEXUS criterion for imaging

Study	Population	Target condition	Index test	Reference standard	Comments
		bilateral locked facets, hangman's fracture, Jefferson fracture and Type 2 dens fracture			
Malhotra 2018 ³⁰ N=1080 Conducted in USA Retrospective	Patients with suspected blunt cervical spine injury that underwent CT of cervical spine followed by MRI of cervical spine Unclear if most or all had head injury as no details provided	Any cervical spine injury, including osseous and ligamentous injuries	CT of cervical spine OR MRI of cervical spine	MRI said to be reference standard for cervical spine injuries in the study which is not the case according to our protocol Data presented in paper therefore analysed using combined CT and MRI as reference standard	 Indirectness: Unclear if all had head injury as no details provided Reference standard of MRI alone does not match protocol, but available data analysed using CT and MRI as a combined reference standard (meaning only sensitivity could be calculated, not specificity
Novick 2018 ³⁵ N=241 Conducted in USA Retrospective	Patients (mean age 43.9 years) undergoing both CT and MRI of cervical spine for any reason, with a history of trauma in medical records. Unclear if all or most had head injury – 17% reported to have closed head injury, but	Cervical spine injuries – ligamentous or bony injury of the cervical vertebral spine, disc injuries, or spinal cord injuries as assessed by imaging	CT of cervical spine OR MRI of cervical spine	CT and MRI as a combined reference standard – means specificity cannot be calculated (as false positives not possible when the index test forms part of the reference standard)	 Indirectness: 17% reported to have closed head injury be unclear if remaining participants suffered head injury as part of the injury mechanism. Not possible to calculate specificity using reference standard as defined in the study.

Study	Population	Target condition	Index test	Reference standard	Comments
	unclear for others if head injury was part of the injury mechanism				Risk stratification: unclear
Schoenfeld 2018 ⁴² N=668 Conducted in USA Retrospective	Adults receiving CT and MRI for evaluation of cervical spine injury following trauma Unclear if most or all had head injury as no details provided	Cervical spine injury – poorly defined	CT of cervical spine OR MRI of cervical spine	No specific reference standard mentioned but possible to calculate sensitivity of CT and MRI using CT + MRI as reference standard as specified in protocol	Indirectness: • Unclear if all had head injury as no details provided Using CT + MRI as reference standard means it is only possible to calculate sensitivity and not specificity Risk stratification: unclear
Songur 2020 ⁴⁴ N=195 Conducted in Turkey Retrospective	Patients (mean age 47.3 years) admitted to ED with diagnosis of blunt cervical spine trauma undergoing CT and MRI of cervical spine Unclear if most or all had head injury as no details provided	Unstable cervical spine injury – Based on neurological status of the patient, degree of spinal canal stenosis and degree of instability. Denis' 1983 definition of single-level ligamentous injury extending to two of three columns.	CT of cervical spine OR MRI of cervical spine	MRI said to be reference standard for cervical spine injuries in the study which is not the case according to our protocol Data presented in paper therefore analysed using combined CT and MRI as reference standard	Indirectness: • Unclear if all had head injury, but suggests all may have had CT of brain Reference standard of MRI alone does not match protocol, but available data analysed using CT and MRI as a combined reference standard (meaning only sensitivity could be calculated, not specificity) Risk stratification: unclear

2 3

Table 4: Summary of studies included in the evidence review – adults – only including those admitted, not those subsequently

discharged following index test

dmitted with blunt aumatic injury and valuated for cervical ine injury by cross- ble lateral diographs (X-ray) % had head CT as	Acute cervical spine injuries – poorly defined	Cross-table lateral radiographs (X-ray)	Reference standard unclear, possibly a final diagnosis based on any further imaging performed	 Indirectness: 50% had head CT and unclear if remaining had head injury Unclear if reference standard included a 2-
aumatic injury and valuated for cervical ine injury by cross- ble lateral diographs (X-ray) % had head CT as			unclear, possibly a final diagnosis based on any further imaging	 50% had head CT and unclear if remaining had head injury Unclear if reference
art of diagnostic sts, unclear if maining patients ad some form of ead injury as part of e injury mechanism				week follow-up period Included in spinal assessment guideline Risk stratification: unclear
ert patients (>16 ears) following blunt auma that underwent eral cervical spine m (X-ray) and ervical spine CT nclear if all or most ad head injury as no etails provided	Cervical spine fracture	Lateral cervical spine film (X-ray)	CT of cervical spine – gold standard for fractures	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Focuses only on fractures in the outcome Included in spinal assessment guideline Risk stratification: unclear
sta mad e e au e au e rv nc ad	aining patients some form of d injury as part of injury mechanism t patients (>16 rs) following blunt ma that underwent ral cervical spine (X-ray) and rical spine CT lear if all or most head injury as no	s, unclear if aining patients some form of d injury as part of injury mechanism t patients (>16 rs) following blunt ma that underwent ral cervical spine (X-ray) and rical spine CT lear if all or most head injury as no	s, unclear if aining patients some form of d injury as part of injury mechanism t patients (>16 rs) following blunt ma that underwent ral cervical spine (X-ray) and rical spine CT lear if all or most head injury as no	s, unclear if aining patients some form of d injury as part of injury mechanism t patients (>16 rs) following blunt ma that underwent ral cervical spine (X-ray) and rical spine CT lear if all or most head injury as no

Study	Population	Target condition	Index test	Reference standard	Comments
Resnick 2014 ⁴¹ N=830 Conducted in USA Prospective	Adults (>18 years) that sustained blunt trauma and underwent CT evaluation of the cervical spine Unclear if all or most had head injury as no details provided	Clinically significant cervical spine injury – those requiring surgical intervention for stabilisation or halo placement, as well as unstable injuries requiring a hard collar	Multidetector row helical CT	Final diagnosis at time of discharge (including all imaging and operative findings)	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Unclear if reference standard included a 2- week follow-up period Included in spinal assessment guideline Risk stratification: unclear

2 3

Table 5: Summary of studies included in the evidence review – adults – only including those that are obtunded, unconscious and/or requiring intensive care unit admission

Study	Population	Target condition	Index test	Reference standard	Comments			
X-ray as index test								
	Despite some studies reporting data for X-ray as an index test in this population, these were not included given X-ray would not usually be used as an initial imaging test in this population of obtunded/unconscious patients.							
CT as index test								
Adams 2006 ¹ N=97 Conducted in USA	Patients undergoing MRI cervical spine trauma protocol at high risk of axial trauma due to pain,	Cervical spine injury – poorly defined	CT of cervical spine	Final diagnosis based on MRI and CT and clinical decision-making of spinal consultants	 Indirectness: All included were at high-risk/more severely injured which may be less applicable to 			
Retrospective	neurological symptoms or obtundation after				general population of those attending ED with suspected cervical spine			

Study	Population	Target condition	Index test	Reference standard	Comments
	significant blunt trauma Unclear if all or most had head injury as no details provided				 injuryUnclear if reference standard included a 2-week follow-up period References standard possibly places focus on MRI results Included in spinal assessment guideline Risk stratification: deemed high risk for axial trauma, unclear which stratification rule used
Berne 1999 ⁴ N=58 Conducted in USA Prospective	 High-risk blunt trauma patients (age ≥17 years) where spine could not be evaluated clinically (e.g. due to head injury, shock, etc.) and need for CT of another body area and intensive care unit admission 53% had associated head injury (intracranial bleed), unclear if remaining suffered some form of head injury as part of the injury mechanism 	Unstable cervical spine injury – classified as unstable in consultation with combined neurosurgical- orthopaedic spine service and based on published guidelines	Complete cervical CT	Final diagnosis based on all imaging/studies	 Indirectness: All required ICU admission so represent more severe subgroup of injuries which may be less applicable to general population of those attending ED with suspected cervical spine injury Unclear if reference standard included a 2- week follow-up period Risk stratification: described as high-risk blunt trauma, unclear which stratification rule used

Study	Population	Target condition	Index test	Reference standard	Comments
Brohi 2005 ⁶ N=381 analysed for CT Conducted in UK Retrospective	Unconscious intubated trauma patients (median age 34 years for whole cohort) Unclear if all or most had head injury as no details provided	Unstable cervical spine injury – defined using White and Punjabi system and three- column model of Denis	Helical CT scan of cervical spine	Final diagnosis, including all imaging performed (MRI in some) and follow-up through hospital stay to identify missed injuries	 Indirectness: All included were unconscious representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injuryHead Unclear if reference standard included a 2-week follow-up period Included in spinal assessment guideline Risk stratification: not reported, but all were unconscious, intubated patients
Raza 2013 ⁴⁰ N=53 Conducted in UK Retrospective	Adult blunt trauma patients with GCS ≤14 (altered sensorium/obtunded), intoxicated with alcohol or drugs and undergoing CT of cervical spine following trauma	Clinically significant cervical spine injury - poorly defined	CT of cervical spine	Final diagnosis of injury at hospital discharge, follow-up appointments or any readmissions Possibly includes >2 weeks follow-up as readmissions and follow-up appointments taken into account	Indirectness: All included had altered sensorium/were obtunded representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury

Study	Population	Target condition	Index test	Reference standard	Comments
	Unclear if all or most had head injury as no details provided				Risk stratification: unclear
Widder 2004 ⁴⁹ N=102 Conducted in Canada Prospective	High-risk severely injured patients (average age 32.0 years) following blunt trauma Unclear if all or most had head injury as no details provided	Cervical spine abnormality – poorly defined	CT of cervical spine	Final diagnosis at discharge and any readmissions Possibly includes >2 weeks follow-up as readmissions taken into account	Indirectness: • All included were high-risk severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury Risk stratification: high-risk severely injured patients, unclear which stratification rule used
CT alone and MRI a CT and MRI separat		ests – CT and MRI combir	ied used as reference sta	ndard (only sensitivity co	uld be calculated for each of
Fisher 2013 ¹⁷ N=277 Conducted in USA Retrospective	Obtunded patients (GCS <15) following blunt trauma undergoing CT and MRI of cervical spine. Mixture of adults and children, but majority were adults ≥18 years (86%)	Clinically significant cervical spine injury CT and MRI scans considered clinically significant if detecting one of the following: ligamentous injury in two adjacent spinal columns, subluxations, cord injury, nerve root injury, disc herniations, and	CT alone OR MRI alone	CT and MRI as a combined reference standard – means specificity cannot be calculated (as false positives not possible when the index test forms part of the reference standard)	Indirectness: • All included were obtunded representing more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injuryHead Despite calculating sensitivity of the two modalities used alone, the

Study	Population	Target condition	Index test	Reference standard	Comments
	Unclear if most or all had head injury as no details provided	fractures except certain types as specified by NEXUS			study notes the intention was not to compare the accuracy of CT and MRI as a solo modality but to assess the added value of MRI to more safely clear the cervical spine. Not possible to calculate specificity using reference standard as defined in the study Risk stratification: unclear, all obtunded patients
Lau 2018 ²⁸ N=63 Conducted in Singapore Retrospective	Patients suffering blunt traumatic injuries that were mentally obtunded and evaluation of cervical spine using CT and MRI Suggests all may have undergone assessment for brain injuries (limited information)	Cervical spine injuries – poorly defined but appears to include bony and soft tissue injuries	CT scan of cervical spine OR MRI of cervical spine	MRI said to be reference standard for cervical spine injuries in the study which is not the case according to our protocol Data presented in paper therefore analysed using combined CT and MRI as reference standard	Indirectness: • All included were obtunded representing more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury Reference standard of MRI alone does not match protocol, but available data analysed using CT and MRI as a combined reference standard (meaning only sensitivity could be calculated, not specificity)

Study	Population	Target condition	Index test	Reference standard	Comments
					Risk stratification: unclear, all obtunded patients
Parmar 2018 ³⁶ N=27 analysed Conducted in Australia Prospective	Adult unconscious trauma patients that had CT and MRI of cervical spine Unclear if most or all had head injury as no details provided	Any cervical spine injury	CT of cervical spine OR MRI of cervical spine	MRI said to be reference standard for cervical spine injuries in the study which is not the case according to our protocol Data presented in paper therefore analysed using combined CT and MRI as reference standard	Indirectness: Indirectness: All included were unconcious representing more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury Reference standard of MRI alone does not match protocol, but available data analysed using CT and MRI as a combined reference standard (meaning only sensitivity could be calculated, not specificity) Risk stratification: unclear, all unconscious requiring mechanical ventilation
Tan 2014 ⁴⁷ N=83 Conducted in USA	Obtunded patients with diagnosis of intracranial haemorrhage and undergoing CT and MRI of cervical spine following non-high	Unstable cervical spine injury	CT of cervical spine OR MRI of cervical spine	MRI said to be reference standard for cervical spine injuries in the study which is not the case according to our protocol	 Indirectness: All included were obtunded representing more severely injured subgroup which may be less applicable to general population of those

Study	Population	Target condition	Index test	Reference standard	Comments
Retrospective	impact trauma (e.g. ground level falls) All had head injury (intracranial haemorrhage) to be included			Data presented in paper therefore analysed using combined CT and MRI as reference standard	attending ED with suspected cervical spine injury Reference standard of MRI alone does not match protocol, but available data analysed using CT and MRI as a combined reference standard (meaning only sensitivity could be calculated, not specificity) Risk stratification: unclear, all obtunded patients admitted to ICU with intracranial haemorrhage

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Table 6: Summary of studies included in the evidence review – adults – other very specific populations

Study	Population	Target condition	Index test	Reference standard	Comments
X-ray as index test					
Dan Lantsman 2020 ¹⁰ N=129 analysed Conducted in Israel Retrospective	Those (median age 83 years) with radiographic diagnosis of diffuse idiopathic skeletal hyperostosis following low-energy trauma and suspected spinal injury (results provided separately for cervical spine injuries)	Acute fracture - those not present in studies prior to the trauma and consisting of a radiographically depicted cortical disruption or impaction of	X-ray of spine Performed in anterior- posterior and lateral projections.	Whole spine CT (results provided separately for cervical spine injuries) Performed in axial plane on 64-slice machine.	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Limited to very specific population of those that had diffuse idiopathic skeletal hyperostosis which may not be

Study	Population	Target condition	Index test	Reference standard	Comments
	Unclear if all or most had head injury as no details provided	the trabeculae and paravertebral soft tissue infiltration.			 applicable to general population Only includes fracture in the outcome and not other types of injuries Risk stratification: unclear
Goodnight 2008 ²¹ N=379 Conducted in USA Retrospective	Adults (≥18 years) following blunt trauma that received CT of cervical spine and follow-up flexion- extension radiographs for continued cervical pain Unclear if all or most had head injury as no details provided	Ligamentous cervical spine injury	X-ray – flexion- extension radiographs	All available evidence, including MRI in some patients	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Those with confirmed fractures were excluded, meaning population may differ from those presenting without any imaging/assessment Focuses only on ligamentous injuries Unclear if reference standard included a 2- week follow-up period Included in spinal assessment guideline Risk stratification: unclear
CT as index test					
Bush 2016 ⁷	Intoxicated adults (≥18 years) with blunt	Clinically significant cervical spine injury: any injury defined as	CT scan	Cervical spine injury diagnosis at discharge/follow-up:	Indirectness:Head injury not mentioned so unclear if

Study	Population	Target condition	Index test	Reference standard	Comments
N=632 analysed (intoxicated subgroup) Conducted in USA Prospective	trauma undergoing CT of the cervical spine Unclear if most or all had head injury as no details provided	unstable or potentially unstable injury that required surgical stabilisation or prolonged immobilisation.		 includes composite endpoint, which included MRI findings, operative findings and clinical status at discharge. Components of reference standard likely differ between patients. Also mentions identification of missed clinically significant injuries from outpatient notes following discharge. Unclear how long this follow up was for and whether the same in all patients. 	 most or all had head injury Limited to very specific population of those that are intoxicated Unclear if reference standard included a 2- week follow-up period Risk stratification: unclear, but all intoxicated adults
Goodnight 2008 ²¹ N=379 Conducted in USA Retrospective	Adults (≥18 years) following blunt trauma that received CT of cervical spine and follow-up flexion- extension radiographs for continued cervical pain Unclear if all or most had head injury as no details provided	Ligamentous cervical spine injury	Helical CT of cervical spine	All available evidence, including MRI in some patients	 Indirectness: Head injury not mentioned so unclear if most or all had head injury Those with confirmed fractures were excluded, meaning population may differ from those presenting without any imaging/assessment Focuses only on ligamentous injuries

Study	Population	Target condition	Index test	Reference standard	Comments
					 Unclear if reference standard included a 2- week follow-up period
					Included in spinal assessment guideline
					Risk stratification: unclear

Table 7: Summary of studies included in the evidence review – children – all having index test and not limited to those that were admitted

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Study	Population	Target condition	Index test	Reference standard	Comments		
X-ray as index test							
Rana 2009 ³⁹ N=54 Conducted in USA Retrospective	Paediatric patients (<18 years) following trauma and undergoing cervical spine imaging by plain radiography and CT Unclear if all or most had head injury as no details provided	Cervical spine injury – poorly defined	X-ray	CT Unclear if solely CT or other later imaging findings also included	Indirectness: • Unclear if all had head injury as no details provided Included in spinal assessment guideline Risk stratification: unclear		
Somppi 2018 ⁴³ N=574 (n=495 analysed for X-ray) Conducted in USA	Children and adolescents (≤19 years) presenting with possible neck injury	Cervical spine injury - ligamentous or osseous injury documented by attending radiologist in their report	X-ray	Follow-up/other imaging? Unclear definition. Mentions the following: to ensure complete identification of spinal cord injuries,	 Indirectness: 40% of whole population had head CT but unclear if the remaining participants had head 		

Study	Population	Target condition	Index test	Reference standard	Comments
Retrospective	Unclear if all or most had head injury – 40% of whole population reported to have had a head CT			medical records for all patients with a negative imaging study (CT or MRI) were reviewed for up to 1 month after the index ED visit to assess for cervical spine pain on ED or outpatient visits to the institution	 injury as part of the injury mechanism unclear if reference standard matches protocol as poorly defined Risk stratification: unclear
CT as index test					
Derderian 2019 ¹² N=221 Conducted in USA Retrospective	Children (median age 9 years) following trauma and undergoing cervical spine CT and MRI scan Unclear if all or most had head injury – 15.8% reported to have isolated head injury and 66.5% multiorgan injury (unclear if this include head injury)	Clinical instability – defined as those undergoing surgical intervention (spinal fusion or halo placement)	CT scan (any abnormality - stable or unstable injuries used to calculate diagnostic accuracy data)	Clinical instability (requiring intervention or not) – assume this was ascertained through follow-up of records Follow-up duration unclear	 Indirectness: clear that a proportion suffered head injury as part of the injury but unclear if this was the case for most people in the study Not formally described as a diagnostic accuracy study and no sensitivity etc. reported, but data available to calculate sensitivity and specificity for clinically unstable injuries. Risk stratification: unclear
Henry 2013-2 ²⁵ N=84 Conducted in USA	Children (≤18 years) with suspected cervical spine injury following trauma with CT and MRI performed within 48 h	Soft tissue injuries (compression fractures, soft tissue oedema, ligamentous injury, muscular injury and	CT – CT assessed for ability to detect soft tissue injuries	MRI –MRI used as reference standard for soft tissue injuries of cervical spine	 Indirectness: unclear if all or most experienced some form of head injury as part of the injury mechanism

Study	Population	Target condition	Index test	Reference standard	Comments
Retrospective	Unclear if all or most had head injury.	spinal cord injury) of cervical spine			 outcome limited to soft tissue injuries only Risk stratification: unclear
Rana 2009 ³⁹ N=254 Conducted in USA Retrospective	Paediatric patients (<18 years) following trauma and undergoing cervical spine imaging by plain radiography and CT Unclear if all or most had head injury as no details provided	Cervical spine injury – poorly defined	CT of cervical spine	Subsequent imaging Unclear if everyone followed up for same duration	 Indirectness: Unclear if all had head injury as no details provided Unclear if reference standard included a 2- week follow-up period Included in spinal assessment guideline Risk stratification: unclear
Somppi 2018 ⁴³ N=574 (n=130 analysed for CT) Conducted in USA Retrospective	Children and adolescents (≤19 years) presenting with possible neck injury Unclear if all or most had head injury – 40% of whole population reported to have had a head CT	Cervical spine injury - ligamentous or osseous injury documented by attending radiologist in their report	СТ	Follow-up/other imaging? Unclear definition. Mentions the following: to ensure complete identification of spinal cord injuries, medical records for all patients with a negative imaging study (CT or MRI) were reviewed for up to 1 month after the index ED visit to assess for cervical spine pain on ED or outpatient visits to the institution	 Indirectness: 40% of whole population had head CT but unclear if the remaining participants had head injury as part of the injury mechanism unclear if reference standard matches protocol as poorly defined Risk stratification: unclear

Study	Population	Target condition	Index test	Reference standard	Comments
MRI as index test					
Derderian 2019 ¹² N=221 Conducted in USA Retrospective	Children (median age 9 years) following trauma and undergoing cervical spine CT and MRI scan Unclear if all or most had head injury – 15.8% reported to have isolated head injury and 66.5% multiorgan injury (unclear if this include head injury)	Clinical instability – defined as those undergoing surgical intervention (spinal fusion or halo placement)	MRI scan (any abnormality - stable or unstable injuries used to calculate diagnostic accuracy data)	Clinical instability (requiring intervention or not) – assume this was ascertained through follow-up of records Follow-up duration unclear	 Indirectness: clear that a proportion suffered head injury as part of the injury but unclear if this was the case for most people in the study Not formally described as a diagnostic accuracy study and no sensitivity etc. reported, but data available to calculate sensitivity and specificity for clinically unstable injuries. Risk stratification: unclear
Henry 2013-1 ²⁶ N=73 Conducted in USA Retrospective	Children (≤18 years) with suspected cervical spine injury that could not be cleared using clinical criteria undergoing MRI-STIR within 48 h	Cervical spine injury with instability – requiring surgical stabilisation: either undergoing surgery or demonstrating signs of instability, pain or neurological compromise during follow-up	MRI with STIR (short T1 inversion recovery) sequence	Follow-up or flexion- extension radiographs: injury requiring surgical intervention or presenting with clinical (significant pain or neurological compromise) or radiographic evidence of instability upon follow- up. Flexion-extension radiographs used to identify false positive findings on MRI.	 Indirectness: unclear if all or most experienced some form of head injury as part of the injury mechanism Included in spinal assessment guideline Risk stratification: unclear, those that could not be cleared clinically

Study	Population	Target condition	Index test	Reference standard	Comments
				Mean follow-up: 10.0 (18.4 months), range 4 days to 7.6 years	
Henry 2013-2 ²⁵ N=84 Conducted in USA Retrospective	Children (≤18 years) with suspected cervical spine injury following trauma with CT and MRI performed within 48 h Unclear if all or most had head injury.	Osseous injuries (fractures, locked facets, subluxations and dislocations) of cervical spine	MRI – MRI assessed for ability to detect osseous injuries of cervical spine	CT– CT used as reference standard for osseous injuries of cervical spine	 Indirectness: unclear if all or most experienced some form of head injury as part of the injury mechanism outcome limited to fractures Risk stratification: unclear
Somppi 2018 ⁴³ N=574 (n=21 analysed for MRI) Conducted in USA Retrospective	Children and adolescents (≤19 years) presenting with possible neck injury Unclear if all or most had head injury – 40% of whole population reported to have had a head CT	Cervical spine injury - ligamentous or osseous injury documented by attending radiologist in their report	MRI	Follow-up/other imaging? Unclear definition. Mentions the following: to ensure complete identification of spinal cord injuries, medical records for all patients with a negative imaging study (CT or MRI) were reviewed for up to 1 month after the index ED visit to assess for cervical spine pain on ED or outpatient visits to the institution	 Indirectness: 40% of whole population had head CT but unclear if the remaining participants had head injury as part of the injury mechanism unclear if reference standard matches protocol as poorly defined Risk stratification: unclear

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Table 8: Summary of studies included in the evidence review – children – only including those that are o	btunded, unconscious and/or
requiring intensive care unit admission	

Study	Population	Target condition	Index test	Reference standard	Comments
X-ray as index test					
Brockmeyer 2012 ⁵ N=24 Conducted in USA Prospective	Children (<17 years) with severe traumatic injuries admitted to ICU undergoing assessment of cervical spine by X- ray, CT and MRI Unclear if all or most had head injury.	Early cervical spine instability – required surgical correction	X-ray	Clinical outcome/diagnosis of early instability – undergoing surgical correction Follow-up possibly >2 weeks as mentions plain radiographs at follow-up of 3-4 months post- injury	 Indirectness: All included were severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury Note: only one patient had confirmed early instability in the study Included in spinal assessment guideline Risk stratification: unclear, all comatose with severe traumatic injuries
CT as index test					
Al-Sarheed 2020 ² N=65	Children (<15 years) with suspected cervical spine injury and that were unconscious	Cervical spine injury mandating stabilisation – no further details provided	CT scan	Radiology/clinical examination, including MRI for some where this was performed.	 All included were unconcious representing more severely injured

Study	Population	Target condition	Index test	Reference standard	Comments
Conducted in Saudi Arabia Retrospective	Unclear if all or most had head injury – 23.3% with skull fracture and 17.4% with intra/extra-axial brain haemorrhage, smaller proportions with skull/face laceration, brain oedema or brain herniation				 subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury unclear if reference standard matches protocol as poorly defined Risk stratification: unclear, all were unconscious and intubated
Brockmeyer 2012 ⁵ N=24 Conducted in USA Prospective	Children (<17 years) with severe traumatic injuries admitted to ICU undergoing assessment of cervical spine by X- ray, CT and MRI Unclear if all or most had head injury.	Early cervical spine instability – required surgical correction	CT of cervical spine	Clinical outcome/diagnosis of early instability – undergoing surgical correction Follow-up possibly >2 weeks as mentions plain radiographs at follow-up of 3-4 months post- injury	 Indirectness: All included were severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury Note: only one patient had confirmed early instability in the study Included in spinal assessment guideline

Study	Population	Target condition	Index test	Reference standard	Comments
					Risk stratification: unclear, all comatose with severe traumatic injuries
Qualls 2015 ³⁸ N=63 Conducted in USA Retrospective	Children (median age 9.6 years) at a children's hospital admitted with severe traumatic brain injury and assessed for cervical spine injury with CT and MRI All had severe traumatic brain injury to be included	Unstable cervical spine injury: resulting in neurological deficit localised to cervical spinal cord, operative stabilisation, halo placement or cervical immobilisation of 3 months of greater	CT alone	CT followed by MRI (CT + MRI) combined Some also had plain radiography of cervical spine and unclear if this also used as part of reference standard for these patients	 Indirectness: All included were severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury Risk stratification: unclear, all with severe traumatic brain injury
MRI as index test					
Brockmeyer 2012 ⁵ N=24 Conducted in USA Prospective	Children (<17 years) with severe traumatic injuries admitted to ICU undergoing assessment of cervical spine by X- ray, CT and MRI Unclear if all or most had head injury.	Early cervical spine instability – required surgical correction	MRI of cervical spine	Clinical outcome/diagnosis of early instability – undergoing surgical correction Follow-up possibly >2 weeks as mentions plain radiographs at follow-up of 3-4 months post- injury	 All included were severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury Note: only one patient had confirmed early instability in the study

Study	Population	Target condition	Index test	Reference standard	Comments
					Included in spinal assessment guideline Risk stratification: unclear, all comatose with severe traumatic injuries
Qualls 2015 ³⁸ N=63 Conducted in USA Retrospective	Children (median age 9.6 years) at a children's hospital admitted with severe traumatic brain injury and assessed for cervical spine injury with CT and MRI All had severe traumatic brain injury to be included	Unstable cervical spine injury: resulting in neurological deficit localised to cervical spinal cord, operative stabilisation, halo placement or cervical immobilisation of 3 months of greater	MRI alone	CT followed by MRI (CT + MRI) combined Some also had plain radiography of cervical spine and unclear if this also used as part of reference standard for these patients	 Indirectness: All included were severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury Risk stratification: unclear, all with severe traumatic brain injury

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2 See Appendix D for full evidence tables.

1 **1.1.6 Summary of the diagnostic evidence**

The assessment of the evidence quality was conducted with emphasis on test sensitivity and specificity as this was identified by the committee as the primary measure in guiding decision-making. Clinical decision thresholds of sensitivity/specificity =0.9 and 0.60 above which a test would be recommended and 0.7 and 0.4 below which a test is of no clinical use were set. Of sensitivity and specificity, it was agreed that sensitivity is the most important measure as the consequences of missing injuries, particularly those that are found to be clinically significant, may be severe.

Results are separated into the four main population groups identified for adults and two main population groups identified for children (see
Diagnostic evidence section above for details), which are presented in separate GRADE tables. Within each GRADE table studies are further
separated based on whether or not most had head injury or suspected head injury, the reference standard and the outcome (for example any
cervical spine injury is separated from those studies reporting clinically significant injuries and those reporting a specific injury only such as
fractures are separated from those covering both osseous and ligamentous cervical spine injuries). Although some studies were similar in terms of
population, index test, reference standard and outcome, pooling was not performed given the amount of variation across studies included in the

12 review.

13 Those where the first column has been highlighted in green indicate studies where all or most were thought to have concomitant head injury. Note

this does not include those that were in obtunded, unconscious or severely injured populations where we have assumed head injury was present based on the nature of the injuries.

16

17 Adults – all having index test and not limited to those that were admitted

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19 X-ray as index test

20 Table 9: Clinical evidence summary: X-ray

Index Test/stu dy Those wit	Numbe r of studies th blunt tra	n auma with	Ref. standard all having h	Follow- up ead CT, C ⁻	Outcome definitio n T as referen	Sensitivity (95% CI) ce standard, cervic	Specificity (95% CI) al spine fracture as	e ontcome bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Gale 2005	1	400	CT of cervical spine	Unclear	Cervical spine fracture –	0.32 (0.13 to 0.57)	0.99 (0.98 to 1.00)	Sensitivity Very serious ^a	, Seriou s ^b	None	None	VERY LOW

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear					no further details			Specificit Very serious ^a	y Seriou s ^ь	None	None	VERY LOW
outcome	th trauma	and low ris	sk (one NEX	US criterio	on), unclear Clinically	if head injury, CT a 0.40 (0.12 to	s reference standa 0.97 (0.94 to	rd, clinical Sensitivit		cant cervio	cal spine in	ijury as
nloo 2021	1	220	cervical spine	Unclear	significan t cervical spine	0.74)	0.99)	Very serious	y Seriou s ^c	None	Serious ^d	VERY LOW
					injury (based on neurologi cal recomme ndation for subluxati on/disloc ation or acute fracture or both)			Specificit	y			
								Very serious ª	Seriou s ^c	None	None	VERY LOW
	-				-	ury, final diagnosis				ficant inju	ry as outco	ome
Bailitz 2009 – high risk	1	Unclear (n=15 positive on	Final diagnosis in medical	Unclear	Clinically significan t cervical spine	0.47 (0.21 to 0.73)	NR	Sensitivit Very serious ^a	y Very seriou s ^c	None	Serious ^d	VERY LOW

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE	
		referenc e standar d)	record at discharge		injury – requiring operative procedur e, halo applicatio n and/or rigid cervical collar			Specificity NA	/				
Bailitz	1	Unclear	Final	Unclear	Clinically	0.37 (0.16 to	NR	Sensitivity					
2009 – moderat e risk	positive in t cervica on medical spine referenc record at injury –				Very seriousª	Very seriou s ^c	None	None	VERY LOW				
		e standar d)	discharge		requiring operative procedur e, halo applicatio n and/or rigid cervical collar			Specificity NA	/				
Bailitz	1	Unclear	Final	Unclear	Clinically	0.25 (0.07 to	NR	Sensitivity	/				
2009 – Iow risk	2009 –	(n=16 diagnosis positive in on medical		significan t cervical spine	0.52)		Very seriousª	Very seriou s ^c	None	None	VERY LOW		
		referenc			injury –			Specificity	/				

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE	
		e standar d)	record at discharge		requiring operative procedur e, halo applicatio n and/or rigid cervical collar			NA					
Mathen 2007	1	667	Final	Unclear	Clinically significan	0.44 (0.25 to 0.65)	0.95 (0.93 to 0.97)	Sensitivity	/				
Risk			diagnosis based on all prospecti		t cervical spine injury – requiring surgery or long- term stabilisati on with a collar or halo	0.03)	0.97)	Very seriousª	Very seriou s ^c	None	None	VERY LOW	
stratifica tion:			vely			requiring			Specificity	/			
those not meeting NEXUS low risk criteria			collected clinical data and imaging results			a		Very serious ^a	Very seriou s ^c	None	None	VERY LOW	
Any follo	wing traun	na, uncleai	r if head inju	ıry, CT as	reference st	andard, cervical sp	ine fractures as ou	tcome					
Lee	1	604	Helical	Unclear	Cervical	0.33 (0.19 to	NR	Sensitivity	/				
2001 Risk		(gives results	atients CT scan spine gives fracture esults no furth	fracture – no further	0.51)		Very seriousª	Very seriou s ^e	None	None	VERY LOW		
stratifica		for total			details			Specificity	/				

tion: describe das patients high n- index of suspicio cervical nfor with cervical nfor with spine injury which rule based on) Takami tione tracture sine tracture sine tracture sine tracture injury tracture sine tracture injury tracture on NR A A A A A A A A A A A A A	Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
2014 spine same admissi on fracture - no further details 0.85) 0.85) Very serious Very serious of the series of the se	describe d as high index of suspicio n for cervical spine injury (unclear which rule based		not patients – includin g some with multiple fracture										
	2014 Risk stratifica tion: unclear, those with high-	1	179		same admissi	spine fracture – no further		NR	Very serious ^a Specificity	Very seriou s ^f	None	Serious ^d	
	Duane 2010	1	49	MRI of cervical spine	Unclear	Ligament ous cervical	0.00 (0.00 to 0.37)	0.98 (0.87 to 1.00)	Sensitivity Very seriousª	/ Very seriou s ^g	None	None	VERY LOW

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear					spine injury	Of 8 injuries missed, 5 were significant (2 with associated fractures requiring prolonged collar and 3 requiring operation)		Specificity Very serious ^a	/ Very seriou s ^g	None	None	VERY LOW
outcome		-	-			ead injury, reference		-		vical spine	e injuries a	S
Griffen 2003 Risk	1	1199	Unclear, possible all imaging/f ollow-up	Unclear	Cervical spine injury - poorly defined	0.65 (0.55 to 0.73)	NR	Sensitivity Very serious ^a	/ Very seriou s ^c	None	Serious ^d	VERY LOW
stratifica tion: unclear					donnod			Specificity NA	/			
Any with outcome	blunt inju	ry and cerv	vical spine a	issessmer	nt, unclear if	⁻ head injury, refere	nce standard uncle	ar/final dia	gnosis, c	ervical sp	ine fractur	es as
Nguyen	1	19	Diagnosi	Unclear	Cervical	0.93 (0.68 to	0.95 (0.74 to	Sensitivity	/			
2005 – high risk			s based on final reports		spine fractures – no	1.00)	1.00)	Very seriousª	Very seriou s ^h	None	Serious ^d	VERY LOW
			including all		further details			Specificity	/			
			imaging		details			Very serious ^a	Very seriou s ^h	None	None	VERY LOW
	1	78		Unclear				Sensitivity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Nguyen			Diagnosi		Cervical	Not estimable as	1.00 (0.95 to	NA				
2005 – Iow risk			s based on final		spine	there were no	1.00)	Specificity	/			
IOW TISK			including all imaging		fractures – no further details	reference standard positive sin this low-risk group		Very seriousª	Very seriou s ^h	None	None	VERY LOW

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Common issues contributing to risk of bias were: it was unclear whether or not a consecutive sample was enrolled, it was unclear if the index test and/or reference standard were interpreted without knowledge of the other, the time interval between index test and reference standard was unclear and it was unclear if the

5 reference standard consisted of the same components for all patients or there was likely to be a difference in components between patients

6 ^b Downgraded by 1 increment as outcome limited to fractures rather than any cervical spine injury

^c Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, and unclear if reference standard included
 a 2 week follow-up period

^d Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use

^e Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, outcome limited to fractures rather than any

cervical spine injury, and results interpreted at fracture level not patient level (patients could have more than one fracture and these included
 individually in analysis)

¹⁴ ^f Downgraded by 2 increments as head injury mentioned for a small proportion of participants but unclear if head injury was part of the injury 15 mechanism for all or most, and outcome focuses specifically on fractures rather than any cervical spine injury

^g Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, and outcome limited to ligamentous injuries
 rather than any cervical spine injury

- ^h Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, outcomes focuses only on fractures and
 unclear if reference standard included a 2 week follow-up period
- 3
- Č
- 4
- 5 **CT as index test**

6 Table 10: Clinical evidence summary: CT

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Those me	eting at le	ast one NI	EXUS criteri	on, unclea	r if head inj	ury, final diagnosis	as reference stand	ard, clinic	ally signi	ficant injur	y as outco	ome
Bailitz	1	Unclear	Final	Unclear	Clinically	1.00 (0.78 to	NR	Sensitivity	/			
2009 – high risk		(n=15 positive on	diagnosis in medical		significan t cervical spine	1.00)		Very seriousª	Very seriou s ^b	None	Serious⁰	VERY LOW
		referenc e	record at discharge		injury – requiring			Specificity	/			
		standar d)	alcontarge		operative procedur e, halo applicatio n and/or rigid cervical collar			NA				
Bailitz	1	Unclear	Final	Unclear	Clinically	1.00 (0.82 to	NR	Sensitivity	/			
2009 – moderat e risk		(n=19 positive on	diagnosis in medical		significan t cervical spine	1.00)		Very seriousª	Very seriou s ^b	None	Serious ^c	VERY LOW
		referenc			injury –			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
		e standar d)	record at discharge		requiring operative procedur e, halo applicatio n and/or rigid cervical collar			NA				
Bailitz	1	Unclear	Final	Unclear	Clinically	1.00 (0.79 to	NR	Sensitivity	/			
2009 – Iow risk		(n=16 positive on referenc	diagnosis based on all		significan t cervical spine	1.00)		Very seriousª	Very seriou s ^ь	None	Serious ^c	VERY LOW
		e	prospecti vely					Specificity	/			
		standar d)	collected clinical data and imaging results		spine injury – requiring operative procedur e, halo applicatio n and/or rigid cervical collar			NA				
Mathen	1	667	Final	Unclear	Clinically	1.00 (0.87 to	0.94 (0.92 to	Sensitivity	/			
2007 Risk stratifica			diagnosis based on all prospecti		significan t cervical spine injury –	1.00)	0.96)	Very serious ^a	Very seriou s ^b	None	Serious ^c	VERY LOW
suaunca			vely		requiring			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
tion: those not meeting NEXUS low risk criteria			collected clinical data and imaging results		surgery or long- term stabilisati on with a collar or halo			Very serious ^a	Very seriou s ^b	None	None	VERY LOW
	-		-		•••	•	erence standard, cli			ractures as	s outcome	
Inaba 2016	1	10,276	Final	Median	Clinically significan	0.98 (0.96 to 1.00)	0.91 (0.90 to 0.92)	Sensitivity				
Risk			diagnosis at discharge	length of stay was 2	t cervical spine	1.00)	0.92)	Very seriousª	Very seriou s ^d	None	None	VERY LOW
stratifica			, including	(IQR 1- 6) days	fracture – requiring			Specificity	/			
tion: patients failing low-risk NEXUS criteria			results of all imaging and operative findings	, .	surgical stabilisati on, Halo Orthotic placemen t or use of a Cervical- Thoracic Orthotic			Very serious ^a	Very seriou s ^d	None	None	VERY LOW
Any with outcome	-	y and cerv	vical spine a		unclear if he	ead injury, referenc	e standard unclear/	final diagn	osis, cer	vical spine	injuries a	S
Duane	1	9227	Later		1.00 (0.99 to	1.00 (1.00 to	Sensitivity	/				
2016 Risk			diagnosis of injury – poorly		ligamento us injury	1.00)	1.00)	Very seriousª	Very seriou s ^e	None	None	VERY LOW
stratifica			defined		Oľ			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
tion: unclear, describe d as patients with criteria for trauma team alert					cervical spine – no further details			Very serious ^a	Very seriou s ^e	None	None	VERY LOW
Griffen	1	1199	Unclear,	Unclear	Cervical	1.00 (0.97 to	NR	Sensitivity	/			
2003 Risk			possible all imaging/f		spine injury - poorly	1.00)		Very seriousª	Very seriou s ^b	None	None	VERY LOW
stratifica			ollow-up		defined			Specificity	/			
tion: unclear								NA				
Vanguri	1	5676	Unclear,	Unclear	Cervical	1.00 (0.99 to	1.00 (1.00 to	Sensitivity	/			
2014 Risk			possibly including other			1.00)	1.00)	Very seriousª	Very seriou s ^b	None	None	VERY LOW
stratifica			imaging		denned			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
tion: unclear, those meeting criteria for trauma team activatio n			(such as MRI and flexion- extension radiograp hs dependin g on patient)					Very serious ^a	Very seriou s ^b	None	None	VERY LOW
Any with outcome	-	ry and cerv	vical spine a	ssessmer	nt, unclear if	head injury, refere	nce standard uncle	ear/final dia	gnosis, c	ervical sp	ine fractur	es as
Nguyen	1	19	Diagnosi	Unclear	Cervical	1.00 (0.78 to	1.00 (0.82 to	Sensitivity	/			
2005 – high risk			s based on final reports		spine fractures – no	1.00)	1.00)	Very seriousª	Very seriou s ^f	None	Serious	VERY LOW
			including all		further details			Specificity	/			
			imaging		uotallo			Very seriousª	Very seriou s ^f	None	None	VERY LOW
Nguyen	1	78	Diagnosi	Unclear	Cervical	Not estimable as	1.00 (0.99 to	Sensitivity	/			
2005 – Iow risk			s based on final		spine fractures	there were no	1.00)	NA				
IOW TISK			reports		– no	reference standard positive		Specificity	/			
			including all imaging		further details	sin this low-risk group		Very seriousª	Very seriou s ^f	None	None	VERY LOW
	1	676		Unclear				Sensitivity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Ptak 2001			Final clinical diagnosis		Cervical spine fracture –	0.98 (0.91 to 1.00)	1.00 (0.99 to 1.00)	Very seriousª	Very seriou s ^f	None	None	VERY LOW
Risk			, in altrativa at		no further			Specificity	/			
stratifica tion: unclear			including operative and discharge notes (possibly incorpora ting CT results)		details		eference standard (Very serious ^a	Very seriou s ^f	None	None	VERY LOW

Any with suspected blunt cervical spine injury, >75% with head CT, CT + MRI as reference standard (only sensitivity possible), any cervical spine injury as outcome

		Unclear	Jnclear		•		Sensitivity	/									
cervical 0.89) no false positives possible when an	spine									spine	0.89)	possible when an	Very seriousª	None	None	None	LOW
injury – index test forms no further part of the													Specificity	Y			
details reference												reference	NA				
standard												standard					

Any with suspected blunt cervical spine injury, unclear if head injury, CT + MRI as reference standard (only sensitivity possible), unstable cervical spine injury as outcome

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Songur	1	88	MRI	Unclear	Unstable	0.78 (0.67 to	Not estimable as	Sensitivity	/			
2020			reported to be		cervical spine	0.86)	no false positives possible when an	Very seriousª	Seriou s ^g	None	Serious ^c	VERY LOW
Risk stratifica			reference standard		injury – based on		index test forms part of the	Specificity	/			
tion: unclear			in the paper, but data available to calculate using CT + MRI as reference standard		neurologi cal status of the patient, degree of spinal canal stenosis, and the degree of instability . Denis' 1983 delineatio n was used in the definition of unstable injury	inium: CT + MDI or	reference standard	NA				
Any with injury as		l blunt cer	vical spine i	njury, unc	lear if head	injury, CT + MRI as	reference standard	l (only sen	sitivity p	ossible), a	ny cervical	spine
Malhotra	1	1080	MRI	Unclear	Any	0.71 (0.67 to	Not estimable as	Sensitivity	/			
2018			reported to be		cervical spine	0.75)	no false positives possible when an	Very seriousª	Seriou s ^g	None	Serious	VERY LOW

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear			reference standard in the paper, but data available to calculate using CT + MRI as reference standard		injury (including osseous and ligamento us injuries)		index test forms part of the reference standard	Specificity NA	/			
Novick 2018	1	241	Referenc e standard	Unclear	Cervical spine injuries	0.87 (0.79 to 0.93)	Not estimable as no false positives possible when an	Sensitivity Very serious ^a	/ Seriou s ^h	None	Serious ^c	VERY LOW
Risk stratifica tion: unclear			not reported in the paper, but data available to calculate using CT + MRI as reference standard		(ligament ous or bony injuries)		index test forms part of the reference standard	Specificity NA	/			
Schoenf eld 2018	1	668	Referenc e standard not	Unclear	Cervical spine injury –	0.79 (0.73 to 0.84)	Not estimable as no false positives possible when an index test forms	Sensitivity Very serious ^a	/ Seriou s ^g	None	None	VERY LOW
			not					Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear			reported in the paper, but data available to calculate using CT + MRI as reference standard		poorly defined		part of the reference standard	NA				

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at 1

high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Common issues contributing to risk 2 3 of bias were: it was unclear whether or not a consecutive sample was enrolled or it was clear convenience sampling was performed, it was unclear

or unlikely that the index test and/or reference standard were interpreted without knowledge of the other, the time interval between index test and

4 reference standard was unclear or likely inappropriate (>48 h) and it was unclear if the reference standard consisted of the same components for 5

6 all patients or there was likely to be a difference in components between patients

7 ^b Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, and unclear if reference standard included a 2 week follow-up period 8

9 ^c Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for sensitivity to determine if an imaging test should be recommended or was of no clinical use 10

11 ^d Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, and reference standard indirectness as outcome only includes fractures and does not involve a period of 2 weeks follow-up as specified in the protocol 12

13 ^e Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, and unclear if reference standard matches protocol as poorly defined 14

^f Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, outcomes focuses only on fractures and 15

unclear if reference standard included a 2 week follow-up period 16

- 1
- ⁹ Downgraded by 1 increment as head injury not mentioned and unclear if all or most had head injury
- ³ ^h Downgraded by 1 increment as head injury status only clear for 17%, unclear if others had suspected head injury/head imaging
- 4
- 5

6 MRI as index test

7 Table 11: Clinical evidence summary: MRI

		al evident	ce Summai	y . WITN								
Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Any with injury as	•	l blunt cerv	vical spine i	njury, >75	% with head	CT, CT + MRI as re	ference standard (c	only sensit	ivity pos	sible), any	cervical s	pine
Friesen	1	108	MRI	Unclear	Any	0.71 (0.62 to	Not estimable as	Sensitivity	/			
2013			reported to be reference		cervical spine	0.78)	no false positives possible when an index test forms	Very serious ^a	None	None	Serious ^b	VERY LOW
Risk			standard		injury – no further		part of the	Specificity	/			
stratifica tion: 98% met at least one NEXUS criterion for imaging			in the paper, but data available to calculate using CT + MRI as reference standard		details		reference standard	NA				

Any with suspected blunt cervical spine injury, unclear if head injury, CT + MRI as reference standard (only sensitivity possible), unstable cervical spine injury as outcome

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Songur 2020 Risk stratifica tion: unclear	1	88	MRI reported to be reference standard in the paper, but data available to calculate using CT + MRI as reference standard	Unclear	Unstable cervical spine injury – based on neurologi cal status of the patient, degree of spinal canal stenosis, and the degree of instability . Denis' 1983 delineatio n was used in the definition of unstable	1.00 (0.95 to 1.00)	Not estimable as no false positives possible when an index test forms part of the reference standard	Sensitivity Very serious ^a Specificity NA	Seriou s ^c	None	None	VERY LOW
Any with injury as Malhotra 2018	outcome	i blunt cer 1080	vical spine i MRI reported	njury, unc Unclear	injury	injury, CT + MRI as 0.83 (0.79 to 0.86)	reference standarc Not estimable as no false positives	Sensitivity	/			I I spine VERY
_0.0			to be		spine		possible when an	Very serious ^a	Seriou s ^c	None	None	LOW

dy	r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear			reference standard in the paper, but data available to calculate using CT + MRI as reference standard		injury (including osseous and ligamento us injuries)		index test forms part of the reference standard	Specificity NA	,			
	241	Referenc e standard	Unclear	Cervical spine injuries	0.77 (0.68 to 0.85)	Not estimable as no false positives possible when an	Sensitivity Very serious ^a	, Seriou s ^d	None	Serious ^b	VERY LOW	
Risk stratifica tion: unclear			not reported in the paper, but data available to calculate using CT + MRI as reference standard		(ligament ous or bony injuries)		index test forms part of the reference standard	Specificity NA	,			
Schoenf eld 2018	1	668	Referenc e standard not	Unclear	Cervical spine injury –	1.00 (0.99 to 1.00)	Not estimable as no false positives possible when an index test forms	Sensitivity Very serious ^a Specificity	Seriou s ^c	None	None	VERY LOW

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear			reported in the paper, but data available to calculate using CT + MRI as reference standard		poorly defined		part of the reference standard	NA				

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at

2 high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Common issues contributing to risk

3 of bias were: it was unclear whether or not a consecutive sample was enrolled, it was unclear or unlikely that the index test and/or reference

standard were interpreted without knowledge of the other, and the time interval between index test and reference standard was unclear or likely
 inappropriate (>48 h)

^b Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use

- 8 °Downgraded by 1 increment as head injury not mentioned and unclear if all or most had head injury
- 9 ^d Downgraded by 1 increment as head injury status only clear for 17%, unclear if others had suspected head injury/head imaging

10

11 Adults – only including those admitted, not those subsequently discharged following index test

1 X-ray as index test

2 Table 12: Clinical evidence summary: X-ray

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
-	itted with I is outcome		/ and cervic	al spine as	ssessed, un	clear if head injury	, reference standard	d unclear/f	inal diagi	nosis, cerv	vical spine	
Cohn	1	60	Referenc	Unclear	Acute	0.57 (0.18 to	NR	Sensitivity	/			
1991 Risk			e standard unclear,		cervical spine injuries –	0.90)		Very seriousª	Very seriou s ^ь	None	Very serious⁰	VERY LOW
stratifica			possibly final		poorly defined			Specificity	/			
tion: unclear			diagnosis based on any further imaging performe d (including flexion/ex tension views, cervical CT scans or tomogra ms where indicated)					NA				
Any adm	itted follow	ving traum	a, unclear if	head inju	ry, CT as re	ference standard, o	ervical spine fractu	ires as out	come			
	1	1004		Unclear				Sensitivit	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Duane 2008			CT of cervical spine		Cervical spine fracture –	0.19 (0.11 to 0.29)	0.99 (0.98 to 1.00)	Very seriousª	Very seriou s ^d	None	None	VERY LOW
Risk					no further			Specificity	/			
stratifica tion: unclear					details			Very serious ^a	Very seriou s ^d	None	None	VERY LOW

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Issues contributing to risk of bias were (vary depending on the study): it was unclear whether or not a consecutive sample was enrolled, it was unclear whether the reference
 standard was interpreted without knowledge of the index test, the time interval between index test and reference standard was unclear and it was unclear if the reference standard consisted of the same components for all patients

- ^b Downgraded by 2 increments as head CT performed for 50% but unclear if remaining also had head injury as part of injury mechanism, and
 unclear if reference standard included a 2 week follow-up period
- ^c Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use
- ^d Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, and outcome limited to fractures rather than
 any cervical spine injury

1 CT as index test

2 Table 13: Clinical evidence summary: CT

Index Test/stu dy	Number of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
	tted follow			l spine as	sessed, unc	lear if head injury,	final diagnosis as	reference :	standard,	clinically	significant	
Resnick	1	830	Final	Unclear	Clinically	1.00 (0.85 to	1.00 (1.00 to	Sensitivity	/			
2014 Risk			diagnosis at discharge		significan t cervical spine	1.00)	1.00)	Very seriousª	Very serious	None	Serious	VERY LOW
stratifica			(all imaging		injury – required			Specificity	/			
tion: unclear			and operative findings)		either surgical interventi on for stabilisati on or halo placemen t, or mandator y use of a hard collar to protect an unstable ligamento us injury			Very serious ^a	Very serious	None	None	VERY LOW

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Issues contributing to risk of bias were: it was unclear if the reference standard includes a period of at least 2 weeks follow-up, it was unclear if the reference standard was
 interpreted without knowledge of the index test and it was likely that the reference standard was alightly different between petients.

- 4 interpreted without knowledge of the index test and it was likely that the reference standard was slightly different between patients
- 5
- ^b Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, and unclear if reference standard
 incorporates 2 week follow-up period specified in the protocol
- ^c Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use
- 10
- 11 Adults only including those that are obtunded, unconscious and/or requiring intensive care unit admission
- 12 CT as index test
- 13 Table 14: Clinical evidence summary: CT

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
	trauma pa iry as outo 1		n, neurolog i Final	ical sympt	oms or obtu Cervical	undation), unclear in	f head injury, final d	liagnosis a Sensitivity		ce standa	rd, any cei	rvical
2006 Risk stratifica			diagnosis based on MRI, CT and		spine injury – poorly defined	so CÌs no calculable)	so Cls no calculable)	Very serious ^a	Very seriou s ^b	None	Serious	VERY LOW

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
tion: deemed high risk for axial trauma, unclear which stratifica tion rule used			clinical decision- making of spinal consultan ts					Very serious ^a	Very seriou s ^b	None	Serious ^d	VERY LOW
High risk Widder	-					-	nce standard, cervi	-		ity as outc	ome	
2004	diagnosis suggest	Cervical spine	1.00 (0.81 to 1.00)	NK	Sensitivity Serious ^a	/ Seriou	None	Serious	VERY			
			at discharge	s follow- up post-	abnormal ity –			Centra	S ^e	None	Centra	LOW
Risk stratifica			consideri	dischar	poorly			Specificity	/			
tion: high-risk severely injured patients, unclear which stratifica tion rule used			ng any readmissi ons	ge as readmis sions mention ed	defined			NA				
High risk injury as		ma and ad	mission to i	intensive o	are unit, un	clear if head injury	, final diagnosis as	reference	standard	, unstable	cervical sp	oine
	1	58		Unclear				Sensitivity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Berne 1999			Final diagnosis based on		Unstable cervical spine	1.00 (0.63 to 1.00)	1.00 (0.93 to 1.00)	Very seriousª	Very seriou s ^f	None	Very serious ^c	VERY LOW
Risk			all		injury – in			Specificity	,			
stratifica tion: describe d as high-risk blunt trauma, unclear which stratifica tion rule used			imaging/s tudies		consultati on with neurosur gical- orthopae dic spine service based on published guideline s			Very serious ^a	Very seriou s ^f	None	None	VERY LOW
Unconcio	ous intubat		patients, u	nclear if he	ead injury, f	inal diagnosis as re	ference standard, ι	instable ce	rvical sp	ine injury	as outcom	е
Brohi	1	381	Final	Unclear,	Unstable	1.00 (0.88 to	0.99 (0.97 to	Sensitivity	,			
2005 Risk			diagnosis , including	through hospital stay	cervical spine injury –	1.00)	1.00)	Very serious ^a	Very seriou s ^g	None	Serious⁰	VERY LOW
stratifica					using White			Specificity	,			
tion: not reported , but all were unconsc ious, intubate d patients	ca all imaging performe d (MRI in some) sc and follow-up through hospital		and Punjabi and three- column model of Denis			Very serious ^a	Very seriou s ^g	None	None	VERY LOW		

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
			ollowing inju ries as outc	-	rvical spine	assessed, unclear	if head injury, final	diagnosis	as refere	nce stand	ard, clinica	ally
Raza	1	53	Final	Unclear	Clinically	1.00 (no raw data	NR	Sensitivity	/			
2013			diagnosis of injury at	duration , includes	significan t cervical	so CIs no calculable)		Very seriousª	Seriou s ^h	None	Very serious ^c	VERY LOW
Risk stratifica			at hospital	follow-	spine injury –			Specificity	/			
tion: high-risk severely injured patients, unclear which stratifica tion rule used			discharge , follow- up appointm ents or any readmissi ons	up post- dischar ge as readmis sions mention ed	poorly defined			NA				
			emorrhage as outcome	_	following n	on high-impact trau	ıma, CT + MRI as re	ference sta	andard (o	only sensit	ivity possi	ble),
Tan	1	83	MRI	Unclear	Unstable	1.00 (no raw data	Not estimable as	Sensitivity	/			
2014			reported to be reference		cervical spine injury –	so CIs no calculable)	no false positives possible when an index test forms	Very seriousª	Seriou s ^h	None	Serious	VERY LOW
			REFERENCE		nijury —			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all obtunde d patients admitted to ICU with intracran ial haemorr hage	d patients,	possibly a	standard in the paper, but data available to calculate using CT + MRI as reference standard		poorly defined	RI as reference stan	part of the reference standard	NA				
outcome		pecciony c		200000				.,	,, un y co	- nour opin	ie inganiee i	
Lau	1	63	MRI	Unclear	Cervical	0.872 (no raw	Not estimable as	Sensitivity				

Lau	1	63	MRI	Unclear	Cervical	0.872 (no raw	Not estimable as	Sensitivity	/			
2018			reported to be reference		spine injuries – poorly	data so CIs no calculable)	no false positives possible when an index test forms	Very serious ^a Specificity	Seriou s ^h	None	Very serious ^c	VERY LOW
								opeening				

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all unconsc ious requiring mechani cal ventilati on			standard in the paper, but data available to calculate using CT + MRI as reference standard		defined but appears to include bony and ligamento us injuries		part of the reference standard	NA				
	d patients ury as outo	•	lults), uncle	ar if head i	njury, CT +	MRI as reference s	tandard (only sensit	tivity poss	ible), clin	ically sigr	nificant cer	vical
Fisher 2013	1	277	Diagnosi s of	Unclear	Clinically significan	0.83 (no raw data so CIs no	Not estimable as no false positives	Sensitivit	y Soriou		Sariauas	

Fis		1	277	Diagnosi	Unclear	Clinically	0.83 (no raw data	Not estimable as	Sensitivity	/			
201	13			s of clinically significan		t cervical	so CIs no calculable)	no false positives possible when an	Very serious ^a	Seriou s ^h	None	Serious	VERY LOW
					spine		index test forms	Specificity	/				

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE	
Risk stratifica tion: unclear, all obtunde d patients			t cervical spine injury by any modality (CT or MRI)		injury – ligamento us injury in two adjacent spinal columns, subluxati ons, cord injury, nerve root injury, disc herniatio ns, and fractures except those specified in NEXUS		part of the reference standard	NA					
Unconcio	ous adults,	unclear if	head injury	, CT + MRI	as reference	e standard (only se	ensitivity possible),	any cervic	al spine	injury as o	outcome		
Parmar	1	27	7 MRI reported to be reference		Cervical spine injuries – poorly	0.74 (0.54 to 0.89)	Not estimable as no false positives possible when an index test forms	Sensitivity					
2018								Very serious ^a	Seriou s ⁱ	None	Serious	VERY LOW	
								Specificity	/				

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all unconsc ious requiring mechani cal ventilati on			standard in the paper, but data available to calculate using CT + MRI as reference standard		defined but appears to include bony and ligamento us injuries		part of the reference standard	NA				

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Common issues contributing to risk of bias were: it was unclear whether or not a consecutive sample was enrolled, it was unclear or unlikely that the index test and/or reference standard were interpreted without knowledge of the other, the time interval between index test and reference standard was unclear or likely inappropriate (>48 h), and it was unclear whether or likely that the components of the reference standard differed between patients

^b Downgraded by 2 increments asall included were high-risk representing a more severely injured subgroup which may be less applicable to
 general population of those attending ED with suspected cervical spine injury, and it is unclear if the reference standard included a 2-week follow up period and reference standard possibly places focus on MRI results

 $^{\circ}$ Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for sensitivity to determine if an imaging test should be recommended or was of no clinical use. Where confidence intervals could not be calculated due to lack of raw data reporting, studies were downgraded by 1 increment if the sample size was ≥70 and <350 and by 2 increments if the sample size was <70.

^d Where confidence intervals could not be calculated due to lack of raw data reporting, studies were downgraded by 1 increment if the sample size was \geq 70 and <350 and by 2 increments if the sample size was <70. ^e Downgraded by 1 increment as all were within more severely injured subgroup which may be less applicable to general population of those
 attending ED with suspected cervical spine injury

^f Downgraded by 2 increments as all were high-risk representing a more severely injured subgroup which may be less applicable to general
 population of those attending ED with suspected cervical spine injury, and unclear if reference standard included a 2 week follow-up period

5 ^g Downgraded by 2 increments as all were unconscious representing a more severely injured subgroup which may be less applicable to general 6 population of those attending ED with suspected cervical spine injury, and unclear if reference standard included a 2 week follow-up period

- ^h Downgraded by 1 increment as all were obtunded representing a more severely injured subgroup which may be less applicable to general
 population of those attending ED with suspected cervical spine injury
- ⁱ Downgraded by 1 increment as all were unconcious representing a more severely injured subgroup which may be less applicable to general
 population of those attending ED with suspected cervical spine injury
- 11

12 *MRI as index test*

13 Table 15: Clinical evidence summary: MRI

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
			emorrhage as outcome		following n	on high-impact trau	ıma, CT + MRI as re	ference sta	andard (o	only sensit	ivity possi	ble),
Tan	1	83	MRI	Unclear	Unstable	1.00 (no raw data	Not estimable as	Sensitivity	,			
2014			reported to be		cervical spine	so Cls no calculable)	no false positives possible when an	Very seriousª	Seriou s ^ь	None	Serious	VERY LOW
			reference		injury –		index test forms	Specificity				

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all obtunde d patients admitted to ICU with intracran ial haemorr hage			standard in the paper, but data available to calculate using CT + MRI as reference standard		poorly defined		part of the reference standard	NA				
Obtunded outcome	l patients,	possibly a	III had brain	assessmo	ent, CT + MF	RI as reference stan	idard (only sensitivi	ity possible	e), any ce	rvical spir	ne injuries	as
Lau 2018	1	63	MRI reported	Unclear	Cervical	1.00 (no raw data	Not estimable as	Sensitivity			. <i>(</i>	

Lau	1	63	MRI	Unclear	Cervical	1.00 (no raw data	Not estimable as	Sensitivity	/			
2018			reported to be		spine injuries –	so CIs no calculable)	no false positives possible when an	Very seriousª	Seriou s ^ь	None	Very serious ^c	VERY LOW
			reference		poorly		index test forms	Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all unconsc ious requiring mechani cal ventilati on			standard in the paper, but data available to calculate using CT + MRI as reference standard		defined but appears to include bony and ligamento us injuries		part of the reference standard	NA				
	d patients ury as outo		ults), unclea	ar if head i	njury, CT +	MRI as reference st	andard (only sensit	tivity poss	ible), clin	ically sign	ificant cer	vical
Fisher 2013	1	277	Diagnosi s of	Unclear	Clinically significan	0.89 (no raw data so CIs no	Not estimable as no false positives	Sensitivity Verv	/ Seriou	None	Serious	VERY

2013	1	211	s of clinically	Unclear	t cervical	so Cls no calculable)	no false positives possible when an	Very serious ^a	/ Seriou s⁵	None	Serious	VERY LOW
			significan		spine		index test forms	Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all obtunde d patients			t cervical spine injury by any modality (CT or MRI)		injury – ligamento us injury in two adjacent spinal columns, subluxati ons, cord injury, nerve root injury, disc herniatio ns, and fractures except those specified in NEXUS		part of the reference standard	NA				
Unconcio	ous adults,	unclear if	head injury	, CT + MRI	as reference	e standard (only se	ensitivity possible),	any cervic	al spine	injury as o	outcome	
Parmar	1	27	MRI	Unclear	Cervical	0.96 (0.81 to	Not estimable as	Sensitivity	/			
2018			reported to be reference		spine injuries – poorly	1.00)	no false positives possible when an index test forms	Very Seriou serious ^a s ^d		None	Serious⁰	VERY LOW
			Gerence		poorty			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all unconsc ious requiring mechani cal ventilati on			standard in the paper, but data available to calculate using CT + MRI as reference standard		defined but appears to include bony and ligamento us injuries		part of the reference standard					

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Common issues contributing to risk of bias were: it was unclear whether or not a consecutive sample was enrolled, it was unlikely that the index test and reference standard were interpreted without knowledge of the other, and the time interval between index test and reference standard was unclear or likely inappropriate (>48 h)

^b Downgraded by 1 increment as all were obtunded representing a more severely injured subgroup which may be less applicable to general
 population of those attending ED with suspected cervical spine injury

 $^{\circ}$ Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for sensitivity to determine if an imaging test should be recommended or was of no clinical use. Where confidence intervals could not be calculated due to lack of raw data reporting, studies were downgraded by 1 increment if the sample size was ≥70 and <350 and by 2 increments if the sample size was <70.

^d Downgraded by 1 increment as all were unconcious representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury

1

2 Adults – other very specific populations

3

4 X-ray as index test

5 Table 16: Clinical evidence summary: X-ray

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
	th diffuse i of cervical 1			Possibly	with low en	ergy trauma, unclea	ar if head injury, wh	ole spine (erence sta	ndard, acu	ite
Lantsma n 2020			spine CT scan	at least 1 month but	cervical spine fracture – those not	so CIs no calculable)	so CIs no calculable)	, Seriousª	Very seriou s ^b	None	Serious⁰	VERY LOW
					IIIUSE IIUI			Specificity				

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear Those wi				ion-extens		aphs for continued	cervical pain (fract	Seriousª	Very seriou s ^b	None	Serious	VERY LOW
Goodnig		379	All	Unclear	Ligament	ical spine injury as 1.00 (0.54 to	0.97 (0.95 to	Sensitivity	1			
ht 2008 (flexion- extensio			available evidence, including MRI in		ous cervical spine	1.00)	0.99)	Very seriousª	Very seriou s ^e	None	Very serious ^c	VERY LOW
n X- rays)			some		injury – poorly			Specificity	/			
Risk stratifica tion: unclear		some poorly	defined			Very serious ^a	Very seriou s ^e	None	None	VERY LOW		

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Issues contributing to risk of bias were (vary depending on the study): it was unclear whether or not a consecutive sample was enrolled, it was unclear whether the reference standard was interpreted without knowledge of the index test, the time interval between index test and reference standard was unclear, not all patients were analysed due to missing radiographs or poor quality radiographs, and it was unlikely that the reference standard consisted of the same components for all patients

^b Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, it was a very specific population of those
 with DISH, a condition making injuries more likely following lower impact trauma, and injury reported was specifically fracture not any type of injury

^o Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use. Where confidence intervals could not be calculated

due to lack of raw data reporting, studies were downgraded by 1 increment if the sample size was \geq 70 and <350 and by 2 increments if the sample

12 size was <70.

^d Where confidence intervals could not be calculated due to lack of raw data reporting, studies were downgraded by 1 increment if the sample size was \geq 70 and <350 and by 2 increments if the sample size was <70.

15

^e Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, population where those with confirmed
 fractures were excluded meaning may differ from population presenting without any prior imaging, outcome limited to ligamentous injuries and
 unclear if reference standard included a 2 week follow-up period

19

20 CT as index test

21 Table 17: Clinical evidence summary: CT

Index Test/stu dy Intoxicate outcome	Numbe r of studies ed adults w	n vith blunt t	Ref. standard rauma, uncl	Follow- up ear if head	Outcome definitio n d injury, fina	Sensitivity (95% CI) Il/discharge diagno	Specificity (95% CI) sis as reference sta	ndard, uns	ness ness ness	lucousi stency vical spin	e injury as	GRADE
	1	631		Unclear				Sensitivity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Bush 2016			Status at discharge /follow-		Unstable cervical spine	0.93 (0.66 to 1.00)	1.00 (0.99 to 1.00)	Very seriousª	Very seriou s ^ь	None	Very serious ^c	VERY LOW
Risk			up, including		injury –			Specificity	/			
stratifica tion: unclear, but all intoxicat ed adults			MRI findings, operative findings and clinical status at discharge		any unstable or potentiall y unstable injury that required surgical stabilisati on or prolonge d immobilis ation			Very serious ^a	Very seriou s ^b	None	None	VERY LOW
				d flexion-extension ra			cervical pain, uncle	ear if head	injury, al	l available	evidence a	as
Goodnig	1		Ligament	1.00 (0.54 to	0.97 (0.94 to	Sensitivity	/					
ht 2008 (flexion- extensio	2008 exion-		available evidence, including MRI in		ous cervical spine injury –	1.00)	0.98)	Very seriousª	Very seriou s ^d	None	Veru serious ^c	VERY LOW
		MRI in			nijury —			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
n X- rays)			some patients		poorly defined			Very seriousª	Very seriou s ^d	None	None	VERY LOW
Risk stratifica tion: unclear												

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Issues contributing to risk of bias were (vary depending on the study): it was unclear whether or not a consecutive sample was enrolled, reasons for exclusion were not reported, the follow-up period for assessing the reference standard was unclear, it was unclear whether the reference standard was interpreted without knowledge of the index test, the time interval between index test and reference standard was unclear, and it was unlikely that the reference standard consisted of the same components for all patients

^b Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, also limited to very specific population of
 those that were intoxicated and unclear time-point for reference standard and whether it matches protocol

^o Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use

^d Downgraded by 2 increments as head injury not mentioned and unclear if all or most had head injury, population where those with confirmed fractures were excluded meaning may differ from population presenting without any prior imaging, outcome limited to ligamentous injuries and

- 13 unclear if reference standard included a 2 week follow-up period
- 14

- 2 Children all having index test and not limited to those that were admitted
- 3

1

4 X-ray as index test

5 Table 18: Clinical evidence summary: X-ray

			oo oummu	J J								
Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Children	with possi	ble neck ir	njury, 40% h	ad head C	T, reference	e standard unclear/	ollow-up/other ima	ging, cervi	cal spine	injury as	outcome	
	1	495	Unclear,	Follow-	Cervical	0.83 (0.36 to	0.97 (0.96 to	Sensitivity	/			
2018 Risk			possibly all imaging	up of records for up to	spine injury (ligament	0.99)	0.99)	Very seriousª	Very seriou s ^ь	None	Very serious⁰	VERY LOW
stratifica	ion: unclear		and follow-up	1 month after	ous and osseous			Specificity	/			
tion: unclear				index ED visit	injuries)			Very seriousª	Very seriou s ^b	None	None	VERY LOW
Children	with cervio	cal spine ir	naging, und	lear if hea	d injury, ref	erence standard as	CT, cervical spine	injury as o	utcome			
Rana	1	54	CT of	Unclear	Cervical	0.615 (no raw	0.016 (no raw	Sensitivity	/			
2009	2009 Risk stratifica ion:	cervical spine		spine injury –	data so Cls no calculable)	data so CIs no calculable)	Very seriousª	Seriou s ^d	None	Very serious ^c	VERY LOW	
Risk				poorly defined			Specificity	/				
stratifica tion: unclear								Very seriousª	Seriou s ^d	None	Very serious ^e	VERY LOW

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at

high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Issues contributing to risk of bias
 were (vary depending on the study): it was unclear whether or not a consecutive sample was enrolled, it was unclear whether the index test and/or

1 reference standard was interpreted without knowledge of the other, the reference standard used for each index test was unclear, the time interval

between index test and reference standard was unclear, and it was unclear whether the reference standard consisted of the same components for
 all patients

^b Downgraded by 2 increments as unclear if all or the majority also sustained a head injury (40% said to have had head CT) and reference standard poorly defined so unclear if matches protocol

^c Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for sensitivity to determine if an imaging test should be recommended or was of no clinical use. Where confidence intervals could not be calculated due to lack of raw data reporting, studies were downgraded by 1 increment if the sample size was \geq 70 and <350 and by 2 increments if the sample size was <70.

- 10 ^d Downgraded by 1 increment as unclear if all or the majority also sustained a head injury
- ^e Where confidence intervals could not be calculated due to lack of raw data reporting, studies were downgraded by 1 increment if the sample size was \geq 70 and <350 and by 2 increments if the sample size was <70.
- 13

14 CT as index test

15 Table 19: Clinical evidence summary: CT

Table 13.		ai evident	ce Summa	<u>y. 01</u>								
Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Children	with possi	ble neck ir	njury, 40% h	ad head C	T, reference	e standard unclear/f	ollow-up/other ima	ging, cervi	cal spine	injury as o	outcome	
Somppi	1	130	Unclear,	Follow-	Cervical	1.00 (0.52 to	1.00 (0.96 to	Sensitivity	/			
2018 Risk			possibly all imaging	up of records for up to	spine injury (ligament	1.00)	1.00)	Very serious ^a	Very seriou s ^b	None	Very serious ^c	VERY LOW
stratifica			and follow-up	1 month after	ous and osseous			Specificity	/			
tion: unclear			- p	index ED visit	injuries)			Very seriousª	Very seriou s ^b	None	None	VERY LOW

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Chlidren outcome	with cervio	cal spine ir	naging, unc	lear if hea	d injury, ref	erence standard as	other imaging find	ings (uncle	ear), cerv	ical spine	injury as	
Rana	1	54	Clinical	Unclear	Cervical	1.00 (no raw data	0.976 (no raw	Sensitivity	/			
2009 Risk			outcome, including subseque		spine injury – poorly	so Cls no calculable)	data so CIs no calculable)	Very seriousª	Very seriou s ^d	None	Very serious ^c	VERY LOW
stratifica	atifica		nt imaging		defined			Specificity	/			
tion: unclear			where performe d					Very seriousª	Very seriou s ^d	None	Very serious ^e	VERY LOW
Children	following t	trauma, un	clear if head	d injury, re	ference sta	ndard as final diagr	nosis/unclear, unsta	able cervic	al spine i	njuries as	outcome	
Derderia	dren following trauma, unclear if h deria 1 221 Unclear		Unclear,	Unclear	Unstable	1.00 (0.89 to	0.85 (0.79 to	Sensitivity	/			
n 2019		confirme d clinical		cervical spine	1.00)	0.90)	Very seriousª	Seriou s ^f	None	Serious	VERY LOW	
			instability		injury –			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear			- records of those undergoi ng interventi on		surgical interventi on (spinal fusion or halo placemen t) indicated clinically unstable while radiologic ally unstable were those with disruption of two or more spinal columns (defined by Denis)			Very serious ^a	Seriou s ^f	None	None	VERY LOW
Children	following	trauma, un	clear if hea	d injury, re	ference sta	ndard as MRI, any s	oft tissue cervical	spine injur	y as outc	ome		
Henry	1	84	MRI of	Unclear	Soft	0.23 (0.05 to	1.00 (0.95 to	Sensitivity	/			
2013-2 Risk			cervical spine		0.54)	1.00)	Very seriousª	Very seriou s ^g	None	None	VERY LOW	
stratifica					spine –			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
tion: unclear					soft tissue oedema, ligamento us injury, muscular injury and spinal cord injury			Very serious ^a	Very seriou s ^g	None	None	VERY LOW

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at

2 high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Common issues contributing to risk

3 of bias were: it was unclear whether or not a consecutive sample was enrolled, it was unclear whether or unlikely that the index test and/or

4 reference standard was interpreted without knowledge of the other, the time interval between index test and reference standard was unclear, and it

5 was unclear whether the reference standard consisted of the same components for all patients

- ^b Downgraded by 2 increments as unclear if all or the majority also sustained a head injury (40% said to have had head CT) and reference
 standard poorly defined so unclear if matches protocol
- ^c Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for sensitivity to determine if an imaging test should be recommended or was of no clinical use. Where confidence intervals could not be calculated due to lack of raw data reporting, studies were downgraded by 1 increment if the sample size was \geq 70 and <350 and by 2 increments if the sample size was <70.
- ^d Downgraded by 2 increments as unclear if all or the majority also sustained a head injury, and unclear if follow-up of at least 2 weeks as part of the reference standard
- ^e Where confidence intervals could not be calculated due to lack of raw data reporting, studies were downgraded by 1 increment if the sample size was \geq 70 and <350 and by 2 increments if the sample size was <70.
- ¹⁶ ^f Downgraded by 1 increment as unclear if all or the majority also sustained a head injury

- ⁹ Downgraded by 2 increments as unclear if all or the majority also sustained a head injury and outcome limited to ligamentous injury not any
- 2 cervical spine injury
- 3
- 4 MRI as index test

5 Table 20: Clinical evidence summary: MRI

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Children	with possi	ible neck iı	njury, 40% h	ad head C	T, reference	e standard unclear/	follow-up/other ima	ging, cervi	cal spine	injury as	outcome	
Somppi	1	21	Unclear,	Follow-	Cervical	1.00 (0.51 to	1.00 (0.75 to	Sensitivity	/			
2018 Risk			possibly all imaging	up of records for up to	spine injury (ligament	1.00)	1.00)	Very seriousª	Very seriou s⁵	None	Very serious ^c	VERY LOW
stratifica	ratifica an on: fol	and follow-up	1 month after	ous and osseous			Specificity	/				
tion: unclear				index ED visit	injuries)			Very seriousª	Very seriou s ^b	None	None	VERY LOW
Children	following	trauma, un	clear if hea	d injury, re	ference sta	ndard as final diagı	nosis/unclear, unsta	able cervic	al spine i	njuries as	outcome	
Derderia	1	221	Unclear,	Unclear	Unstable	1.00 (0.89 to	0.45 (0.37 to	Sensitivity	/			
n 2019			confirme cer d clinical spin		cervical spine injury –	1.00)	0.52)	Very seriousª	Seriou s ^d	None	Serious⁰	VERY LOW
			instability		nijury —			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear			- records of those undergoi ng interventi on		surgical interventi on (spinal fusion or halo placemen t) indicated clinically unstable while radiologic ally unstable were those with disruption of two or more spinal columns (defined by Denis)			Very serious ^a	Seriou s ^d	None	Serious ^e	VERY LOW
Henry	1	73	Injury	Follow-	Cervical	1.00 (0.03 to	0.97 (0.90 to	Sensitivity	/			
2013-1			requiring surgical interventi	up mean	spine instability	1.00)	1.00)	Very seriousª	Seriou s ^d	iou None Very serious ^c	VERY LOW	
					_			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, those that could not be cleared clinically			on or presentin g with clinical or radiograp hic evidence of instability on follow- up	10.0 months	requiring surgical stabilisati on			Very serious ^a	Seriou s ^d	None	None	VERY LOW
Children	following	trauma, un	clear if head	d injury, re	eference sta	ndard as CT, any o	sseous cervical spi	ne injury a	s outcom	e		
Henry	1	84	CT of	Unclear	Osseous	1.00 (0.54 to	0.97 (0.91 to	Sensitivity	/			
2013-2 Risk			cervical spine		injury of cervical spine –	1.00)	1.00)	Very seriousª	Very seriou s ^f	None	Very serious ^c	VERY LOW
stratifica					fractures, locked			Specificity	/			
tion: unclear		facets, subluxati ons and dislocatio ns			Very seriousª	Very seriou s ^f	None	None	VERY LOW			

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at
 high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Common issues contributing to risk
 of bias were: it was unclear whether or not a consecutive sample was enrolled, it was unclear whether or unlikely that the index test and/or
 reference standard was interpreted without knowledge of the other, the time interval between index test and reference standard was unclear, and it
 was unclear whether the reference standard consisted of the same components/same follow-up for all patients

^b Downgraded by 2 increments as unclear if all or the majority also sustained a head injury (40% said to have had head CT) and reference
 standard poorly defined so unclear if matches protocol

1 2

- ^c Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use
- 5 ^d Downgraded by 1 increment as unclear if all or the majority also sustained a head injury
- ^e Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.7 and 0.4, respectively, which were the thresholds used for
 specificity to determine if an imaging test should be recommended or was of no clinical use
- ^f Downgraded by 2 increments as unclear if all or the majority also sustained a head injury and outcome limited to fractures not any cervical spine
 injury

10

- 11 Children only including those that are obtunded, unconscious and/or requiring intensive care unit admission
- 12
- 13 X-ray as index test

14 Table 21: Clinical evidence summary: X-ray

Index Test/stu dy	Number of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci	GRADE
						clear if head injury urgery as outcome		ro as clinio	cal outcor	ne/diagnos	sis at time	OT
Brockm	1	24	Clinical	Possibly	Early	1.00 (0.03 to	0.96 (0.78 to	Sensitivity	/			
eyer 2012			outcome/ diagnosis	>2 weeks	cervical spine	1.00)	1.00)	Very seriousª	Seriou s ^ь	None	Very serious ^c	VERY LOW
			of early	as	instability			Specificity	/			

Test/stu	Number of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all comatos e with severe traumati c injuries			instability – undergoi ng surgical correctio n	mention plain radiogra phs at follow- up of 3- 4 months post- injury	– surgical correctio n			Very serious ^a	Seriou s ^b	None	None	VERY LOW

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Issues contributing to risk of bias were: it was unclear if index tests and reference standard were interpreted without knowledge of the other, time interval between index tests and reference standard consisted of the same components for all patients

^b Downgraded by 1 increment as all were within a severely injured subgroup which may be less applicable to general population of those attending
 ED with suspected cervical spine injury

^c Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use

- 9
- 10 CT as index test
- 11 Table 22: Clinical evidence summary: CT

DRAFT FOR CONSULTATION Imaging of the cervical spine

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
		•	assessment v as outcome		med severe	traumatic brain i	njury, reference sta		+ MRI po		her imaging	g,
Qualls	1	63	Final	Unclear	Unstable	1.00 (0.48 to	0.84 (0.73 to	Sensitivit	y			
2015			diagnosis based on		cervical spine	1.00)	0.93)	Very seriousª	Seriou s⁵	None	Very serious ^c	VERY LOW
Risk			all imaging		injury – injuries			Specificit	y			
stratifica tion: unclear, all with severe traumati c brain injury	with sover	e iniuries	reports (CT, MRI and possibly other imaging)	s uncloar	resulting in neurologi cal deficit localised to cervical spine cord, oeprpativ e stabilisati on, halo placemen t or cervical immobilis ation of 3 months or greater	ry roforonco star	ndard as final diagn	Very serious ^a	Seriou s ^b	None	None	VERY LOW
			ion as outco			,			· · · ,	•••••	1. 2	
	1	65		Unclear				Sensitivit	y			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% CI)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Al- Sarheed 2020			Radiolog y/clinical examinati	Cervical spine injury	0.85 (0.68 to 0.95)	1.00 (0.89 to 1.00)	Very seriousª	Very seriou s ^d	None	Very serious ^c	VERY LOW	
			on, including		requiring stabilisati			Specificity	/			
Risk stratifica tion: unclear, all were unconsc ious and intubate d			MRI for some where performe d		on			Very serious ^a	Very seriou s ^d	None	None	VERY LOW
Brockm	1	24	Clinical	Possibly	Early 1.00 (0.03 to	1.00 (0.85 to	Sensitivity	/				
eyer 2012			outcome/ diagnosis	>2 weeks	cervical spine	1.00)	1.00)	Very seriousª	Seriou s ^e	None	Very serious⁰	VERY LOW
Dist			of early instability	as mention	instability – surgical			Specificity				
Risk stratifica tion: unclear, all comatos e with severe traumati c injuries			undergoi ng surgical correctio n	plain radiogra phs at follow- up of 3- 4 months post- injury	correctio n			Very serious ^a	Seriou s ^e	None	None	VERY LOW

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at
 high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Common issues contributing to risk
 of bias were: it was unclear whether or not a consecutive sample was enrolled, it was unclear whether or unlikely that the index test and/or
 reference standard was interpreted without knowledge of the other, the time interval between index test and reference standard was unclear or not

- 5 appropriate (>48 h), and it was unclear whether or clear that the reference standard did not consist of the same components for all patients
- 6
- ^b Downgraded by 1 increment as all were within a severely injured subgroup which may be less applicable to general population of those attending
 ED with suspected cervical spine injury
- ^o Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use
- ^d Downgraded by 2 increments as all were unconcious representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury, and unclear if the reference standard matches protocol as definition provided is limited to 'radiology/clinical examination'
- ^e Downgraded by 1 increment all were within a severely injured subgroup which may be less applicable to general population of those attending
 ED with suspected cervical spine injury
- 16
- 17 *MRI as index test*

18 Table 23: Clinical evidence summary: MRI

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% Cl)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE	
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Children with cervical spine assessment and confirmed severe traumatic brain injury, reference standard as CT + MRI possibly other imaging, unstable cervical spine injury as outcome

Qualls	1	63	Final	Unclear	Unstable	al 0.99) 0.90)	0.91 (0.69 to	Sensitivity				
2015			diagnosis based on		cervical spine	0.99)	· ·	Very seriousª	Seriou s ^ь	None	Very serious ^c	VERY LOW
			all		injury –			Specificity	/			

Index Test/stu dy	Numbe r of studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all with severe traumati c brain injury			imaging reports (CT, MRI and possibly other imaging)		injuries resulting in neurologi cal deficit localised to cervical spine cord, oeprpativ e stabilisati on, halo placemen t or cervical immobilis ation of 3 months or greater			Very serious ^a	Seriou s ^b	None	Serious ^d	VERY LOW
						nclear if head injury urgery as outcome	r, reference standar	d as clinic	al outcon	ne/diagnos	sis at time	of
Brockm	1	24	Clinical	Possibly	Early	1.00 (0.03 to	0.74 (0.52 to	Sensitivity	/			
eyer 2012			outcome/ diagnosis	>2 weeks	cervical spine	1.00)	0.90)	Very seriousª	Seriou s ^e	None	Very serious⁰	VERY LOW

Specificity

. instability

of early

as

Index Number Test/stu r of dy studies	n	Ref. standard	Follow- up	Outcome definitio n	Sensitivity (95% Cl)	Specificity (95% CI)	Risk of bias	Indirect ness	Inconsi stency	Impreci sion	GRADE
Risk stratifica tion: unclear, all comatos e with severe traumati c injuries		instability – undergoi ng surgical correctio n	mention plain radiogra phs at follow- up of 3- 4 months post- injury	– surgical correctio n			Very serious ^a	Seriou s ^e	None	Serious ^d	VERY LOW

^a Risk of bias was assessed using the QUADAS-2 checklist. The evidence was downgraded by 1 increment if the majority of studies were rated at high risk of bias, and downgraded by 2 increments if the majority of studies were rated at very high risk of bias. Issues contributing to risk of bias were (varied depending on study): it was unclear whether or not a consecutive sample was enrolled, it was unclear whether or unlikely that the index test and/or reference standard was interpreted without knowledge of the other, the time interval between index test and reference standard was unclear whether or clear that the reference standard did not consist of the same components for all patients

^b Downgraded by 1 increment as all were within a severely injured subgroup which may be less applicable to general population of those attending
 ED with suspected cervical spine injury

^o Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.9 and 0.7, respectively, which were the thresholds used for
 sensitivity to determine if an imaging test should be recommended or was of no clinical use

^d Downgraded by 1 or 2 increments if the confidence intervals crossed one or both of 0.7 and 0.4, respectively, which were the thresholds used for specificity to determine if an imaging test should be recommended or was of no clinical use

^e Downgraded by 1 increment as all were within a severely injured subgroup which may be less applicable to general population of those attending
 ED with suspected cervical spine injury

1 **1.1.7 Economic evidence**

2 1.1.7.1 Included studies

- No published health economic studies were included. However, two models were identified
 from previous NICE guidelines:
- NICE Head injury guideline (CG176) 2014 <u>https://www.nice.org.uk/guidance/cg176/evidence/appendices-pdf-191719838</u> (Appendix
 M)
- 8 NICE Spinal injury guideline (NG41) 2016 -
- 9 <u>https://www.nice.org.uk/guidance/ng41/evidence/appendices-jp-pdf-2358425775</u>
 10 (Appendix L)
- These economic evaluations are described in section 1.1.8 Summary of included economicevidence.

13 1.1.7.2 Excluded studies

- 14 No relevant health economic studies were excluded due to assessment of limited
- 15 applicability or methodological limitations.
- 16 See also the health economic study selection flow chart in Appendix F.

1 **1.1.8 Summary of included economic evidence**

A description of two relevant guideline models can be found in Table 24 with an assessment of their applicability and quality. Table 25 shows a
 description of the CG176 model strategies. It indicates the implausible assumption in the base case where far more people have X-ray than CT in
 the CT strategy, Strategy 5. The sensitivity analysis in the lower panel seem more plausible and will be the focus of this review.

5 Table 24: Comparison of previous guideline model characteristics

	NICE Head injury guideline (CG176) 2014	NICE spinal injury guideline (NG41) 2016
Comparators	7 strategies Canadian C-Spine rule vs NEXUS c-spine rule vs image all vs no CT vs X-ray (then MRI if positive or indeterminate)	18 strategies Canadian C-Spine rule vs NEXUS c-spine rule vs image all CT vs X-ray vs MRI Further imaging after a positive scan
Population	Adults with suspected cervical spine injury_and head injury	Adults with suspected (cervical) spinal column injury (bony or ligamentous) and <u>no other injuries</u>
Perspective	NHS & personal social services	NHS & personal social services
Study design	Decision tree	Decision tree
Main outcome	False negatives averted	Quality-adjusted life-years (QALYs)
Applicability assessment	Partially applicable Due to absence of quality-adjusted life-years (or any measure of health outcome.	Partially applicable Due to population not being exclusively people with head injury.
Time horizon	Hospital episode	Lifetime
Treatment effects	95% of missed spinal injuries deteriorate with a cost of \pounds 7,214	0.5% of missed column injuries convert to a cord injury
Cost components	 Imaging costs (x-ray, CT and MRI) Observation (depending on test results) Treatment Litigation cost (in a secondary analysis) 	 Imaging costs (x-ray, CT and MRI) Treatment of column injury (depending on whether true positive, false positive or false negative) Spinal cord injury Litigation cost Cancer treatment cost from radiation

	NICE Head injury guideline (CG176) 2014	NICE spinal injury guideline (NG41) 2016
Limitations	 Most probabilities in model based on expert opinion Indeterminate results Accuracy of tests after an indeterminate test or 2nd-line test Specificity of prediction rule differed a lot for CT and X-ray in the base case analysis. How they were applied in the model was not clearly described. Furthermore, the strategy labelled "CT according to Canadian C-Spine rule" actually had fewer CT scans in the base case analysis than the strategy labelled "X-ray according to Canadian C-Spine rule for CT". This is due to the specificity of CT being misapplied in the model and due to the assumption in both strategies that 50% of false negatives get the other imaging modality. However, there was a sensitivity analysis with far more plausible assumptions for the Canadian C-spine rule strategies – see Table 25. 	 Does not explicitly model the pathway for indeterminate results Assumes accuracy of 2nd-line test is independent of 1st test result Prevalence and evidence for treatment effects based on expert opinion
Quality assessment	Potentially serious limitations	Potentially serious limitations

1 Table 25: Specification of strategies in head injury guideline model (CG176)

2 a) Base case analysis

Probability of having a given initial image	Initial of	clinical decision		Initi	al clinical decisio	n
strategy	(for thos	(for those with injury)				
7	No imaging	CT first	X ray first	No imaging	CT first	X ray first
Strategy 1: No imaging	100%			100%		
Strategy 2: CT all		100%			100%	
Strategy 3: x ray all			100%			100%
Strategy 4: Canadian C spine for Xray	29%	29%	43%	0%	0%	100%
Strategy 5: Canadian C Spine for CT	49.7%	0.6%	49.7%	0%	100%	0%
Strategy 6: NEXUS for Xray	32%	32%	37%	4.65%	4.65%	90.70%
Strategy 7: NEXUS for CT	38%	24%	38%	5%	90%	5%

3 4

b) Sensitivity analysis: 'Committee estimates for initial imaging decisions'

Probability of having a given initial		clinical decision se vithout injury	Initial clinical decision			
image strategy	(for the	lot	those with inju	iry)		
7	No imaging	CT first	X ray first	No imaging	CT first	X ray first
Strategy 1: No imaging	100%			100%		
Strategy 2: CT all		100%			100%	
Strategy 3: x ray all			100%			100%
Strategy 4: Canadian C spine for Xray	54%	3%	43%	0%	0%	100%
Strategy 5: Canadian C Spine for CT	54%	46%	0%	0%	100%	0%
Strategy 6: NEXUS for Xray	60%	3%	37%	4.65%	4.65%	90.70%
Strategy 7: NEXUS for CT	22%	40%	38%	5%	90%	5%

5 6 7

8

Table 25 shows for the CG176 model, the proportion of people having each test for those who do have an underlying spine injury and those that do not. This is shown for every comparator in the base case analysis (top panel) and for a key sensitivity analysis (bottom panel) – For details see

H.1.3.3 and H.1.4 of Appendix H below. The prevalence of injury was only 0.5% therefore the left-hand side of the table covers most patients. In 1 the base case analysis (top panel) CT strategy 5, 49.7% of these patients had an X-ray and only 0.6% had a CT scan. Paradoxically in X-ray 2 strategy 4, 29% had a CT, which was more than in the CT strategy. The current committee concluded that this was illogical and therefore only 3 considered the results of the model based on the sensitivity analysis. 4

Table 26 compares the estimates of accuracy used in each model with those found in the current guideline review. 5

6 Table 26: Diagnostic accuracy of imaging used in previous guideline models

			2021 guideline review – see 1.1.6.
	Head injury guideline model (CG176)	Spinal injury guideline model (NG41)	(depending on reference standard)
X-ray Sensitivity	56.8%	70%	32%-65%
X-ray Specificity	99.7%	84%	95%-99%
CT sensitivity	83.0%	98%	93%-100%
CT specificity	99.9%	100%	91%-100%

7

- Table 27 and Table 28 show the results of each model. Both models found the use of CT with the Canadian C-Spine CT rule to be the most cost-8 9 effective strategy.
- 10 For the 2014 head injury guideline model, we can say little about the sensitivity of results, because the guideline's sensitivity analyses were based on variations from its flawed base case analysis. For the spinal injury guideline model, there was a lot of uncertainty around model parameters but 11 12 it was concluded that the results were robust to plausible changes
- 13 • in the accuracy estimates,
- 14 • in the discount rate,
- when litigation costs were included, 15
- 16 • when the QALY loss associated with false negatives was increased,
- 17 when the time horizon was extended.
- 18 • when the risk and consequences of radiation exposure were included.
- The spinal injury guideline concluded that at the assumed prevalence rates and accuracy data, CT scans in combination with a decision rule are 19
- most likely to be cost effective. Therefore, CT scanning only those with a positive X-ray at the assumed prevalence and accuracy rates results in 20 21
- many missed injuries.

1 Table 27: Sensitivity analysis^a results from NICE head injury model (CG176): 'Committee estimates for initial imaging decisions'

Strategy	Mean cost	False negatives	Cost per false negative avoided (vs no imaging)	Mean cost including litigation cost ^b	Rank ^c
No imaging	£1 ^d	0.500%		£1,001	6
CT all	£329	0.140%	£90,974	£612	2
X-ray all	£558	0.280%	£253,076	£1,116	7
Canadian C-spine rule for Xray	£335	0.280%	£151,890	£893	5
Canadian C-Spine rule for CT	£295	0.140%	£81,478	£578	1
NEXUS C-Spine rule for Xray	£311	0.280%	£140,780	£877	4
NEXUS C-Spine rule for CT	£301	0.170%	£90,866	£633	3

(a) The base case analysis results are fatally flawed. A plausible sensitivity analysis is reported here instead.

(b) Litigation cost was assumed to be £200,000 for each false negative test result.

(c) Rank of mean cost including litigation cost – 1=lowest cost, 7=highest cost.

(d) In the no imaging arm, almost all patients were discharged and there were no treatment costs. There were observation costs for a very small proportion of patients.

1 Table 28: Base case results from NICE spinal injury model (NG41)

		/		
Strategy	Mean cost	Mean Quality-adjusted life- years	Net Health Benefit (£20K per QALY)*	Rank
1. X-ray	£158	20.85252	20.8446	14
2. CT	£121	20.85275	20.8467	7
3. MRI	£191	20.85270	20.8431	18
4. X-ray + CT	£127	20.85251	20.8461	12
5. CT + MRI	£129	20.85268	20.8462	11
6. MRI + CT	£187	20.85268	20.8433	17
7. Canadian C-spine rule + X-ray	£111	20.85252	20.8470	5
8. Canadian C-spine rule + CT	£81	20.85275	20.8487	1
9. Canadian C-spine rule + MRI	£122	20.85270	20.8466	9
10. NEXUS C-spine rule + X-ray	£146	20.85252	20.8452	13
11. NEXUS C-spine rule + CT	£111	20.85274	20.8472	4
12. NEXUS C-spine rule + MRI	£173	20.85269	20.8440	16
13. Canadian C-spine rule + X-ray + CT	£95	20.85251	20.8478	3
14. Canadian C-spine rule + CT + MRI	£89	20.85267	20.8482	2
15. Canadian C-spine rule + MRI + CT	£121	20.85267	20.8466	8
16. NEXUS C-spine rule + X-ray + CT	£119	20.85251	20.8466	10
17. NEXUS C-spine rule + CT + MRI	£119	20.85267	20.8467	6
18. NEXUS C-spine rule + MRI + CT	£170	20.85267	20.8442	15

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3 **1.1.9 Economic model**

4 Original modelling was not conducted for this guideline. The model from CG176 was summarised in 1.1.8 above and the full report can be found in 5 Appendix H.

1 **1.1.10 Unit costs**

2 Relevant unit costs are provided below to aid consideration of cost effectiveness.

Code	Description	Unit cost
RD01A	Magnetic Resonance Imaging Scan of One Area, without Contrast, 19 years and over	£146.75
RD01B	Magnetic Resonance Imaging Scan of One Area, without Contrast, between 6 and 18 years	£215.63
RD01C	Magnetic Resonance Imaging Scan of One Area, without Contrast, 5 years and under	£140.83
RD20A	Computerised Tomography Scan of One Area, without Contrast, 19 years and over	£88.06
RD20B	Computerised Tomography Scan of One Area, without Contrast, between 6 and 18 years	£159.25
RD20C	Computerised Tomography Scan of One Area, without Contrast, 5 years and under	£104.27
PF	Plain Film (including x-ray)	£28.62

3 Direct access costs from NHS Reference costs: 2019-2020 version 2³⁴

4 1.1.11 Evidence statements

5 Economic

One original comparative cost analysis conducted for the 2014 NICE Head Injury guideline, found that using the Canadian C-Spine rule for CT was the least costly strategy when compared to No imaging; CT all, Xray all; Canadian C-spine rule for Xray; NEXUS C-Spine rule for Xray and NEXUS C-Spine rule for CT for initial imaging for adults with suspected cervical spine injury and head injury. This analysis was assessed as partially applicable with potentially serious limitations.

One original cost-utility analysis conducted for the 2016 NICE Spinal Injury guideline,
 found that Canadian C-Spine rule and CT was the cost-effective strategy compared to 17
 other strategies for initial imaging for adults with suspected cervical spinal column injury
 and no other injuries. This analysis was assessed as partially applicable with potentially
 serious limitations.

17 **1.1.12** The committee's discussion and interpretation of the evidence

18 **1.1.12.1. The outcomes that matter most**

19 Diagnostic accuracy

20 Diagnostic accuracy for any significant cervical spine injury (including fracture/bony injury, soft tissue/ligament damage, spinal cord injuries and vascular injuries) was relevant for the 21 diagnostic accuracy component of this review. Sensitivity was considered the most important 22 measure because the initial imaging method should pick up as many true positives as 23 possible to avoid missing those with significant cervical spine injuries and subsequent 24 25 negative consequences for the person with head injury and cervical spine injury, such as disability. It was noted that a high sensitivity contributes to management, as it provides 26 reassurance that the test is good at ruling out injury and allowing early discharge or 27 mobilisation. It was also noted that for imaging it is unlikely that many false positives will 28 occur, so specificity values are generally higher than for other diagnostic tests or scoring 29 systems, such as the clinical decision rules. 30

31 Diagnostic test and treat

For the diagnostic test and treat component of the review, all outcomes were considered
equally important for decision-making and were primary outcomes, including all-cause
mortality at 3 months, quality of life at ≥3 months, objectively reported scores of disability
(such as the Glasgow Outcome Score or extended Glasgow Outcome Score) at ≥3 months,
length of hospital stay, unscheduled readmission (28 days or longer) and neurological
deterioration.

No studies comparing clinical outcomes between two different imaging strategies wereidentified.

9 **1.1.12.2 The quality of the evidence**

10 Possible population indirectness was present for most studies included in the review, as the proportion of people with confirmed head injury was not reported. The population was 11 described only as those with suspected cervical spine injury undergoing imaging of the 12 cervical spine. It was noted that many of those with cervical spine injury are likely to have 13 14 experienced head injury based on the nature of the injury, for example those with whiplash 15 are likely to have suffered a head injury as well. Therefore, the population of included studies 16 varies, with few where it was clear all had confirmed or suspected head injury. Some studies included a population that was limited to those that were unconscious or obtunded, often 17 requiring intensive care unit admission. Although head injury was not specifically mentioned 18 in many studies, it was assumed that these groups did have at least suspected head injury 19 20 given the severity of their injuries; these studies were not downgraded based on head injury 21 not being mentioned.

Studies limiting to those more severely injured, only including people that were unconscious or obtunded, were however downgraded for indirectness. This is because the severe nature of their injuries makes them a very specific subgroup of the population that attend the emergency department with suspected cervical spine injury. Results would be less applicable than those of people who are discharged from the emergency department without admission to hospital or intensive care.

28 Reference standards used across studies differed. Many studies used 'final diagnosis' at 29 discharge or including any readmissions as the reference standard, which was not always 30 well-defined. Where studies had used this as the reference standard, if they had not included 31 at least a 2-week period as part of the follow-up or this was unclear, this was taken into account when assessing indirectness. For studies that had used other reference standards 32 33 listed in the protocol, for example CT or MRI where X-ray was used as the index test or CT 34 and MRI combined for any index test, this was accepted and there was no reason to 35 downgrade the reference standard for indirectness, regardless of whether or not there was a follow-up of 2 weeks. Similarly, a further factor considered in the risk of bias assessment for 36 37 studies using final diagnosis as a reference standard was the fact that not all people included 38 in the study had the same reference standard; for example, some may have had MRI while others did not. 39

Some studies did not use an external reference standard, but data was available to calculate
the sensitivity of both CT and MRI when using CT and MRI combined as the reference
standard; as this means that any person with a positive result on CT or MRI is considered to
be reference standard-positive, false positives are not possible such that only sensitivity, and
not specificity, can be calculated.

In terms of outcome definitions, some studies did report data for 'significant' cervical spine injuries, as specified in the protocol. However, some studies only reported any severity of cervical spine injury, not limiting to significant ones. Data from these studies was still included, but for studies reporting data for any severity and significant injuries, only data for the significant injuries was analysed as it is more relevant to the review protocol. A further way in which outcome definitions varied across studies was the types of injuries (for example, bony or ligamentous/soft tissue) that were included. Some studies included any
type of injury in the outcome/target condition whereas others focused the study on specific
types of injuries, for example only fractures or only ligamentous injuries.

Given the differences between studies described above, pooling of results was not thought to
be appropriate. Studies that were broadly similar in terms of index test, population, reference
standard and target condition were grouped under the same headings but not formally
pooled.

8 Most of the included evidence was very low quality based on the assessment of risk of bias, 9 indirectness and a measure of imprecision for sensitivity and specificity. The exception was 10 studies where head injury was clearly confirmed or suspected in studies, in which case the 11 quality was low rather than very low. There were very few prospective studies, meaning 12 many of the same issues were present in terms of risk of bias assessment.

- Some of the most common reasons that studies were downgraded for risk of bias included:
- 15
- o a consecutive sample not being enrolled or this being unclear
- it being unclear if the index test and/or reference standard were interpreted
 without knowledge of the other
- 18 o the interval between index test and reference standard being unclear
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- o not all patients within a study having the same reference standard
- Indirectness was often present, with studies downgraded for one of the following reasons:
 - Head injury was not mentioned in the paper and included anyone with suspected cervical spine injury
 - They were very specific populations that may not be representative of the general population this review would apply to, including studies limiting specifically to those that were obtunded or unconscious with or without admission to an intensive care unit
- Studies using 'final diagnosis' as a reference standard where it was unclear if
 a 2-week follow-up was incorporated into this standard, given that not all
 people received specific types of imaging such as MRI
- Imprecision was assessed separately for sensitivity and specificity. Thresholds of
 ≥90% and ≥60% for sensitivity and specificity, respectively, were used as values
 above which a test would be recommended and values of 0.7 and 0.4 below which a
 test is of no clinical use were set for sensitivity and specificity, respectively.

35 The limitations of the evidence were taken into account when considering any possible changes to existing recommendations. For children, these limitations contributed to the 36 37 decision not to make any major changes to recommendations. Factors to consider for children include radiation exposure and risk of cancer. For adults, despite the limitations the 38 committee agreed that the evidence supported the removal of X-ray as a primary imaging 39 modality in people with head injury and suspected cervical spine injury. The committee 40 agreed that current practice was already moving away from X-ray in adults and therefore 41 although the evidence did have limitations it supported this change. Although it was unclear if 42 43 many of the studies represented a head injury population, it was agreed that evidence from any people with suspected cervical spine injury is still relevant to the subgroup that also have 44 45 head injury as many with suspected cervical spine injury are likely to have at least some type of head injury depending on the mechanism of injury. 46

1 1.1.12.3 Benefits and harms

2 <u>Adults</u>

Despite the limitations of the included evidence discussed in the previous section, including
very few studies where it was clear the population had a head injury and methodology issues
contributing to concerns about risk of bias, the committee were able to draw some
conclusions from the evidence and use this to support decisions that were made.

7 The sensitivity values for X-ray in adult populations were consistently very low, with fourteen of sixeen analyses demonstrating values less than the 90% threshold specified in the 8 9 protocol as a test that should be recommended and all fourteen of these studies having values also below 70%, with many below 50%. This included studies that reported any 10 severity of injury and also those reporting clinically significant injuries. Where imprecision 11 12 was present, this was because confidence intervals crossed the lower threshold of 70%, 13 meaning even when considering confidence intervals, the sensitivity values for these studies could not be consistent with a value >90%. Results from one study reporting results 14 15 separately for low-, moderate- and high-risk groups demonstrated that sensitivity was worse as the risk decreased; however, it was a very small study. For the two analyses where values 16 17 >90% were reported, this included one very small analysis of 19 participants that were at 18 high-risk and another was assessing the sensitivity of flexion-extension X-rays in those that had already been confirmed as CT-negative. This may represent secondary imaging rather 19 20 than primary imaging and be less relevant to the review protocol. In both cases imprecision was also present, meaning the confidence intervals indicated uncertainty as to whether or 21 22 not the true sensitivity value was >90%. Of the included studies, one of them was clearly in a 23 suspected head injury population (as all had head CT) and the results for this study was one 24 of those with a low sensitivity value for X-ray, which in this case was testing the ability to detect cervical spine fractures of any severity (32%; n=400; very low quality). Across all 25 26 results for X-ray, specificity values were very good, with values >90% where they could be 27 calculated.

28 In contrast, the committee noted that the sensitivity values for CT (for any severity of injury 29 and clinically significant injuries) across studies were higher compared to X-ray. Across 30 analyses using a reference standard other than CT+MRI combined (for example 'final 31 diagnosis'), sensitivity values were all >90%, with fourteen of the eighteen analyses having 32 values of 100%. This included one specific subgroup of intoxicated adults following blunt trauma (where it was unclear if head injury was present) where sensitivity was reported to be 33 34 93%, with imprecision present. The same study as above for X-ray also demonstrated that sensitivity values were 100% in low-, moderate- and high-risk groups for CT, though it was a 35 36 very small study. Imprecision for sensitivity was present for some of the smaller studies, but 37 there were some much larger studies (for example, two studies with ~10,000 people 38 analysed) that also confirmed the high sensitivity values for CT of the cervical spine with no 39 imprecision present. This improvement in sensitivity compared to X-ray did not come at the 40 expense of specificity, as where reported these values were >90% in all but one analysis, 41 with the other being 88%. It was noted that none of the studies discussed here for CT were 42 clearly in a population with head injury or suspected head injury.

43 Studies where CT+MRI was used as a combined reference standard provided information 44 about injuries that may be picked up on MRI but not on CT. This included three studies 45 where it was clear head imaging had been performed in most or all people included and was 46 therefore considered to be more relevant to the head injury population. Some reported any 47 severity of injury while others reported unstable cervical spine injuries. Results 48 demonstrated sensitivity values that were lower than those discussed in the previous paragraph, with all but one being lower than 90% (ranging from 71% to 100%). Using a 49 50 combined reference standard of CT+MRI makes interpretation more difficult, as the lack of external reference standard makes it difficult to determine if a case missed on CT is a false 51 52 negative on CT or a false positive on MRI. This is less of a problem where only unstable

injuries have been reported as this often requires an intervention. The committee noted that
 MRI may pick up ligamentous injuries that are not identified on CT, which may explain the
 lower sensitivity of CT when including results of MRI in the reference standard.

4 The only studies available for MRI as an index test in adults were those using combined 5 CT+MRI as the reference standard, including three studies where it was clear head imaging had been performed in most or all people included and was therefore considered to be more 6 7 relevant to the head injury population. The results were similar to those mentioned in the 8 previous paragraph for CT, but there were five analyses reporting sensitivity values >90% 9 rather than one (values ranged from 71% to 100%). Some reported any severity of injury while others reported unstable cervical spine injuries. These results suggest that some 10 11 studies show that MRI picks up all injuries that were identified on CT, while others suggest it 12 misses some identified on CT. The lack of results for MRI using an external reference standard, for example 'final diagnosis' as for CT and X-ray index tests discussed above. 13 means the evidence identified for MRI as the initial imaging method in those with suspected 14 15 cervical spine injury is more limited.

16 Based on the information discussed in the previous paragraphs, clinical experience and knowledge of current practice, the committee agreed that X-ray should not be used as an 17 initial imaging strategy for the cervical spine in adults with head injury due to it's poor 18 19 sensitivity compared to CT. This was further supported by the committee as they noted that it 20 is being used less frequently in current practice for adults and they highlighted that it can be time-consuming and distracting, with multiple views often required which takes up time (up to 21 3-4 hours), possibly delays the diagnosis process. It was also noted that it can be technically 22 23 difficult in some people, for example those with large shoulders, and inadequate X-rays then 24 mean a CT is done anyway. The committee agreed that they considered the quality of CTs to 25 be more reliable than X-ray. The evidence from the very small study showing worsening sensitivity values for X-ray as risk group reduced (low-, moderate- and high-risk) was cited, 26 27 as it suggested that even though a group is lower risk for cervical spine injury, this does not mean the sensitivity of X-ray is adequate to pick up injuries. This meant that the 28 recommendation for X-ray in the group that have neck pain or tenderness but no high-risk 29 indications for a CT cervical spine was edited so that CT is also performed in this group of 30 31 people and the recommendation for CT in those at high-risk was retained. Other 32 recommendations were edited to remove mention of X-ray, given it is now not included as an 33 initial imaging strategy for adults with suspected cervical spine injury. This was also 34 supported by cost-effectiveness evidence, as discussed in a later section.

Based on evidence from one study showing the sensitivity of X-ray to be poor for detecting
fractures in those with diffuse idiopathic skeletal hyperostosis (DISH) with low-energy
trauma, the committee agreed that people with a condition predisposing them to a higher risk
of injury to the cervical spine (for example, ankylosing spondylitis) should be included as an
additional factor in the group with neck pain or tenderness but not high-risk indications for a
CT cervical spine. This was also extrapolated and included in the respective
recommendation for children.

The committee highlighted the role that MRI has in imaging of the cervical spine, but agreed that there is limited evidence and it is rarely used as the first imaging strategy for adults with suspected cervical spine injury, meaning it cannot be recommended as an initial imaging method. Recommendations about the use of MRI as an additional form of imaging in certain circumstances were retained from the previous version of the guideline.

47 <u>Children</u>

Although limitations associated with the evidence for children were similar to those identified for adults, the committee agreed that additional factors complicated the recommendations for children, including the fact that there are concerns about radiation exposure and the risk of

51 cancer.

1 Similar to the results for adults, some of the results for X-ray as an index test in children also 2 demonstrated sensitivity values <90%. However, there were fewer studies in children and the 3 sensitivity values for X-ray were in general higher compared to the adult population (values 4 of 62%, 83% and 100% across the three studies, compared to adults where sensitivity values 5 were often <50%); however, two of the studies were also small, with <100 people included, 6 and the other was larger with 495 people analysed. There were also no studies where it was 7 clear head injury had been confirmed or was suspected. The two studies with lower 8 sensitivity values were those reporting any severity of cervical spine injury, while the single 9 study reporting unstable injuries reported 100% sensitivity. However, for all three studies 10 there was imprecision was sensitivity, meaning there was uncertainty about the true sensitivity of X-ray in children. Of the three studies, two reported specificity values >90% 11 12 while the other reported a very low specificity of 1.6%.

13 Results for CT across seven included studies, with one specific to head injury as only those with confirmed severe traumatic brain injury were included, demonstrated sensitivity values 14 that were similar to those for adults; five of the seven analyses reported values of 100%, with 15 the other two reporting values of 85% and 23%. The study reporting the very low value of 16 17 23% was assessing the ability of CT to pick up ligamentous injuries specifically, with MRI as the reference standard; as it is established that CT is less able to pick up ligamentous 18 injuries this result was expected, and it was unclear how many of these injuries were 19 20 clinically significant. As for the X-ray results, imprecision was present for these sensitivity values meaning there was uncertainty about the true sensitivity of CT in children. For 21 22 specificity, where reported the values were high, with all being >80% and many being >90%.

23 Results for MRI across six included studies, with one specific to head injury as only those 24 with confirmed severe traumatic brain injury were included, demonstrated sensitivity values 25 that were similar to those discussed for CT in the previous paragraph; five of the six analyses reported values of 100%, with the other (the study with confirmed head injury) reporting a 26 27 value of 80%. This included some studies reporting any severity of injury and others reporting unstable injuries. One study specifically reporting fractures also reported a value of 28 29 100%, even though MRI is usually better at picking up ligamentous injuries. As for the X-ray and CT results, imprecision was present for these sensitivity values meaning there was 30 31 uncertainty about the true sensitivity of MRI in children. For specificity, where reported the 32 values were high, with all but one being >70% and the other being lower at 45%.

33 Based on the information discussed in the previous paragraphs, clinical experience and 34 knowledge of current practice, the committee agreed that there was insufficient evidence to 35 change any of the existing recommendations for children. Although some evidence 36 suggested sensitivity values <90% for X-ray in children, fewer studies reported this compared to adults and the values were higher than those observed for adults. CT performed well in 37 terms of sensitivity for children, as with for adults, but concerns about radiation exposure and 38 39 cancer risk is a factor that needs to be considered in children. Radiation risk of CT was described as the biggest drawback for using CT in children for assessing the cervical spine 40 41 as the risk of radiation-induced tumours (for example, thyroid tumours) may be higher for 42 children as their organs are rapidly developing. Although the committee noted that the 43 evidence for this is based on forecasting tools and extrapolation of risks rather than actual 44 data, it is a risk that should be considered in decision-making and balancing risks and 45 benefits. MRI also demonstrated good sensitivity values across most studies, however the committee noted that limitations of MRI in children include the need for immobilisation and/or 46 47 sedation of children, as it is a longer process than CT and X-ray and requires children to 48 remain still. Additionally, there was no cost-effectiveness evidence available for children for 49 any of the index tests.

50 As the evidence was limited and changing to CT or MRI rather than X-ray as initial imaging in 51 children would lead to a large change in practice for children with head injury and suspected 52 cervical spine injury, the committee did not make any changes to the recommendations for 53 cervical spine imaging in children; CT was retained only for those at higher risk and X-ray 1 recommended for those with no high-risk indications for CT but neck pain or tenderness. The 2 committee further noted that in their experience, clinically significant cervical spine injury in 2 children is much recent than for adults, which further supported the decision net to support the

children is much rarer than for adults, which further supported the decision not to expand the
 use of CT in children any further as the risks of radiation were considered to outweigh any

5 possible benefit in terms of picking up cervical spine injuries.

In terms of MRI, the committee noted that existing recommendations about MRI as an
additional form of imaging in certain circumstances were not age-specific and also applied to
children.

9 As for adults, people with a condition predisposing them to a higher risk of injury to the 10 cervical spine (for example, axial spondylarthritis was also included as an additional factor in the group with neck pain or tenderness but no high-risk indications for a CT cervical spine. 11 12 There was no evidence for this in children but the committee agreed that collagen vascular disease or osteogenesis imperfecta may be important factors in children, as ankylosing 13 14 spondylitis is extremely uncommon. Although in the study the results showed poor sensitivity of X-ray, the recommendation for children was to X-ray these children, as this is the method 15 16 of imaging agreed for those considered to have no high-risk factors for CT in children.

17 **1.1.12.4 Cost effectiveness and resource use**

18 CT and MRI scanning are generally more expensive than plain X-ray but sometimes it can be 19 difficult and time-consuming to get the right views for X-ray of the neck.

The clinical evidence review showed that CT is significantly more sensitive at diagnosing spinal injuries although it involves a substantial dose of radiation. Since untreated spinal injury could lead to damage to the spinal cord, with serious implications for the patient's quality of life and long-term costs for the NHS, the use of more costly imaging could be cost effective, depending on the prevalence of spine injury in the head injury population.

25 Adults

26 No published economic evaluations were found but two previous NICE guideline models27 were identified.

28 The model from the previous head injury CG176 compared strategies that encompassed 29 both the Canadian C-spine and NEXUS rules with plan X-ray or CT scan. The optimal strategy was labelled "CT according to Canadian C-Spine rule" but on close inspection there 30 31 were fewer CT scans in this strategy than in the strategy labelled "X-ray according to 32 Canadian C-Spine rule for CT". This is due to the specificity of CT being misapplied in the 33 model and due to the assumption in both strategies that 50% of false negatives get the other imaging modality. For this reason, the base case analysis of the model is fatally flawed. 34 35 However, there was a sensitivity analysis 'GDG (guideline development group) estimates for initial imaging decisions', where the proportions having each test are far more plausible and 36 37 reflect better the diagnostic accuracy evidence. The result of this sensitivity analysis (and the base case analysis) was that CT according to the Canadian C-spine rule was dominant - it 38 39 had the lowest cost and the fewest false negatives. This model had some other limitations, 40 for example the costs were for the initial hospital episode, not the long-term cost, although 41 litigation costs were included in a sensitivity analysis.

The model from the NICE spinal injury guideline (NG41) also compared strategies that encompassed both the Canadian C-spine and NEXUS rules with plan X-ray, CT scan or MRI. Although the model population specified was suspected c-spine injury <u>and no other</u> injury, the diagnostic accuracy evidence base overlaps a great deal. The strategy that used CT according to the Canadian C-Spine rule was the most cost-effective yielding both the lowest long-term NHS cost and the most QALYs. This model had some limitations, for example the risk of spinal injury deterioration was based on expert opinion. Also, the

- sensitivity of x-ray seems to be higher than the estimates found in the current guideline
 review, whereas the specificity seems to be a bit lower.
- Both guideline models support the use of CT scanning over plain x-ray overall. However,
 they did not explicitly model the previous guideline recommendations, which are CT for high-

5 risk patients and X-ray for moderate risk. This was recommended by the previous guideline

6 committee because they were concerned about the resource implications of referring

7 everyone for CT, the exposure to radiation and the relatively poor specificity of the C-spine

- 8 prediction rules. Modelling of that strategy does not seem to be possible because of the lack
- 9 of data on the prevalence of c-spine injury in the moderate risk population.
- 10 Although there is some uncertainty around the cost effectiveness of extending CT scanning
- 11 to adults who are assessed as moderate risk of cervical spine injury, the committee decided
- 12 to recommend CT because of its much greater sensitivity and because it has become current 13 practice for adults in recent years.

14 <u>Children</u>

15 No economic evaluations were found.

- 16 Given concerns about radiation exposure to the thyroid in children and insufficient evidence
- about the diagnostic accuracy of MRI, the committee decided not to change the
- 18 recommendations for children and therefore plain x-ray was retained for children identified as
- 19 at moderate risk. The cost effectiveness of this strategy is uncertain but since the committee
- 20 were not considering moving away from current practice there is not a cost impact.

21 **1.1.12.5 Other factors the committee took into account**

Some additional changes to recommendations were made that were not based on the
 content of the evidence review, which included removing some older recommendations that
 were no longer seen as necessary given they are now all carried out routinely anyway.
 Recommendations on when to carry out full cervical spine immobilisation were removed as
 there is a guideline on spinal injury which should be referred to.

Other edits included editing the wording for clarity. This included making it clear that MRI recommended if there are neurological signs and symptoms referable to the cervical spine would be subsequent imaging, for example following CT or X-ray, and would not be as an initial form of imaging.

Another edit was made to one of the factors listed as a high-risk indicator for CT in adults and children; the example given for when 'a definitive diagnosis of cervical spine injury is needed urgently' was edited from 'before surgery' to explain further, with manipulation of the cervical spine being requiring during surgery or anaesthesia being added.

35

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- 24

1 Appendices

2 Appendix A – Review protocols

3 Review protocol for CT, MRI and X-ray of the cervical spine in people with head injury – diagnostic accuracy

ID	Field	Content
0.	PROSPERO registration number	CRD42021283523
1.	Review title	What is the diagnostic accuracy of CT, MRI and X-ray of the cervical spine for initial imaging in people with head injury?
2.	Review question	What is the diagnostic accuracy of CT , MRI and X-ray of the cervical spine for initial imaging in people with head injury?
3.	Objective	To determine which of the currently available imaging techniques is best to diagnose cervical spine injury.
4.	Searches	The following databases (from inception) will be searched:
		Cochrane Central Register of Controlled Trials (CENTRAL)
		 Cochrane Database of Systematic Reviews (CDSR)
• Embas		• Embase
• MEDL		MEDLINE
		Searches will be restricted by:
		English language studies
		Human studies
		Letter and comments excluded

		Other searches:
		[Inclusion lists of systematic reviews]
		The searches may be re-run 6 weeks before the final committee meeting and further studies retrieved for inclusion if relevant.
		The full search strategies will be published in the final review.
		Medline search strategy to be quality assured using the PRESS evidence-based checklist (see methods chapter for full details).
5.	Condition or domain being studied	Cervical spine injury in patients who have experienced a head injury.
		Cervical spine injury in patients who have experienced a nead injury.
6.	Population	
0.	Population	Inclusion: Infants, children and adult with head injury and suspected cervical spine injury
		Strata:
		Adults at:
		o high risk
		o moderate risk
		o low risk
		Children + infants at:
		o high risk
		o moderate risk

		○ low risk
		 Adults (aged ≥16 years)
		 Children and infants (aged 0 to <16 years).
		Mixed population studies will be included but downgraded for indirectness. Cut-off of 60% will be used for all age groups
		Exclusion: Adults, and children (including infants under 1 year) with superficial injuries to the eye or face without suspected or confirmed head or brain injury.
		C spine injury risk stratification based on:
		Canadian C Spine Rule
		NEXUS
		Stratification rules should be kept separate as they have different features.
		Both for adults and children. Both rules validated in adults and extrapolated for children. These are not specific for head injury .
7.	Test	Computed tomography scan (CT) of cervical spine
		Magnetic resonance imaging (MRI) of cervical spine
		X-ray of cervical spine
8.	Reference standard	For diagnostic accuracy:
		Reference standard for CT:
		CT and MR imaging of cervical spine
		Or

		2 weeks follow-up after CT including autopsy findings
		 <u>Reference standard for MR imaging:</u> CT and MR imaging of cervical spine
		 Or 2 weeks follow-up after MR imaging including autopsy findings
		 <u>Reference standard for X-ray:</u> CT or MR imaging of cervical spine Or CT and MRI imaging of cervical spine or
		 2 weeks follow-up after X-ray including autopsy findings For X-ray only include children and people below 65 years.
		People >65 years are considered as high risk and will be offered CT cervical spine within 8 hours of injury (CG 176).
		Vascular injuries will be picked up by MR imaging
9.	Types of study to be included	Diagnostic accuracy: Diagnostic cross-sectional studies, cohort studies (prospective and retrospective)

		Systematic reviews and meta-analyses of the above
10.	Other exclusion criteria	Non-English language studies.
		Conference abstracts will be excluded as it is expected there will be sufficient full text published studies available.
11.	Context	Head-injured patient may also have sustained concomitant injury to the cervical spine. Some head injured patients who require a CT head scan will also need cervical spine imaging. The purpose of this review is to inform the optimal diagnostic pathways for these patients using the best evidence available.
12.	Primary outcomes (critical outcomes)	All outcomes are considered equally important for decision making and therefore have all been rated as critical:
		Diagnostic accuracy outcomes
		Diagnostic accuracy CT, MRI and X-ray of the cervical spine for
		any significant cervical spine injury
		(fracture/bony injury, soft tissue/ligament damage, spinal cord injuries, vascular injuries)
		No objective definition for significant cervical spine injury. Note definitions as reported in the papers.
		Diagnostic test accuracy to be reported by test sensitivity/specificity
		For measurement of imprecision, clinical decision thresholds for sensitivity and specificity are set at 90% and 60%.
		Sensitivity is considered to be more important than specificity. Sensitivity is more important as that will change management. Often, the decision is whether

		someone can be discharged from ED. A test with high sensitivity that is negative is very reassuring in ruling out an injury and allowing early discharge or mobilisation. It's unlikely that imaging will produce false positives (i.e. low specificity). COMET database was searched for relevant core outcome sets and none were identified.
13.	Data extraction (selection and coding)	All references identified by the searches and from other sources will be uploaded into EPPI reviewer and de-duplicated.
		10% of the abstracts will be reviewed by two reviewers, with any disagreements resolved by discussion or, if necessary, a third independent reviewer.
		This review will make use of the priority screening functionality within the EPPI- reviewer software.
		The full text of potentially eligible studies will be retrieved and will be assessed in line with the criteria outlined above.
		A standardised form will be used to extract data from studies (see <u>Developing</u> <u>NICE guidelines: the manual</u> section 6.4).
		10% of all evidence reviews are quality assured by a senior research fellow. This includes checking:
		papers were included /excluded appropriately
		a sample of the data extractions
		correct methods are used to synthesise data
		a sample of the risk of bias assessments
		Disagreements between the review authors over the risk of bias in particular studies will be resolved by discussion, with involvement of a third review author where necessary.

14.	Risk of bias (quality) assessment	For diagnostic reviews Diagnostic test accuracy studies: QUADAS-2
 synthesized in a quantitative data analysis. Endnote will be used for bibliography, citations, simanagement. 		 Aggregate data on diagnostic accuracy of investigations will be collected and synthesized in a quantitative data analysis. Endnote will be used for bibliography, citations, sifting and reference management. WinBUGS will be used for meta-analysis of diagnostic accuracy studies if included
16.	Analysis of sub-groups	Subgroups that will be investigated if heterogeneity is present: Types of injuries: • Bone injuries • Spinal cord injuries • other soft tissue injuries • vascular injuries
17.	Type and method of review	□ Intervention ⊠ Diagnostic □ Prognostic

		Qua	Qualitative		
			lemiologic		
		□ Serv	vice Delivery		
		□ Oth	er (please specify)		
18.	Language	English			
19.	Country	England			
20.	Anticipated or actual start date	[For the purposes of PRO review can be defined as	[For the purposes of PROSPERO, the date of commencement for the systematic review can be defined as any point after completion of a protocol but before formal screening of the identified studies against the eligibility criteria begins.		
		A protocol can be deemed complete after sign-off by the NICE team with responsibility for quality assurance.]			
21.	Anticipated completion date	be edited at any time. Al	[Give the date by which the guideline is expected to be published. This field may be edited at any time. All edits will appear in the record audit trail. A brief explanation of the reason for changes should be given in the Revision Notes facility 1		
22.	Stage of review at time of this submission	Review stage	Started	Completed	
		Preliminary searches			
	Piloting of the study select process		ction		
		Formal screening of sea against eligibility criteria	rch results		
		Data extraction			

r				
		Risk of bias (quality) assessment		
		Data analysis		
23.	Named contact	5a. Named contact		
		National Guideline Centre		
		5b Named contact e-mail		
		[Guideline email]@nice.org.uk		
		[Developer to check with Guideline	Coordinator for ema	il address]
		5e Organisational affiliation of the re	eview	
		National Institute for Health and Car Alliance / National Guideline Centre Public Health Guideline Development template text here and one of the cer recognise this as a NICE protocol]	/ NICE Guideline U nt Team] [Note it is	pdates Team / NICE essential to use the
24.	Review team members	[Give the title, first name, last name member of the review team. Affiliation review team members belong.]		
		From the National Guideline Centre	:	
		[Guideline lead]		
		[Senior systematic reviewer]		
		Systematic reviewer		
		[Health economist]		

		[Information specialist]
		[Others]
25.	Funding sources/sponsor	This systematic review is being completed by the National Guideline Centre which receives funding from NICE.
26.	Conflicts of interest	All guideline committee members and anyone who has direct input into NICE guidelines (including the evidence review team and expert witnesses) must declare any potential conflicts of interest in line with NICE's code of practice for declaring and dealing with conflicts of interest. Any relevant interests, or changes to interests, will also be declared publicly at the start of each guideline committee meeting. Before each meeting, any potential conflicts of interest will be considered by the guideline committee Chair and a senior member of the development team. Any decisions to exclude a person from all or part of a meeting will be documented. Any changes to a member's declaration of interests will be recorded in the minutes of the meeting. Declarations of interests will be published with the final guideline.
27.	Collaborators	Development of this systematic review will be overseen by an advisory committee who will use the review to inform the development of evidence-based recommendations in line with section 3 of <u>Developing NICE guidelines: the manual</u> . Members of the guideline committee are available on the NICE website: [NICE guideline webpage].
28.	Other registration details	[Give the name of any organisation where the systematic review title or protocol is registered (such as with The Campbell Collaboration, or The Joanna Briggs Institute) together with any unique identification number assigned. If extracted data will be stored and made available through a repository such as the Systematic Review Data Repository (SRDR), details and a link should be included here. If none, leave blank.]
29.	Reference/URL for published protocol	[Give the citation and link for the published protocol, if there is one.]
30.	Dissemination plans	NICE may use a range of different methods to raise awareness of the guideline. These include standard approaches such as:
		 notifying registered stakeholders of publication

		 publicising the guideline through NICE's newsletter and alerts issuing a press release or briefing as appropriate, posting news articles on the NICE website, using social media channels, and publicising the guideline within NICE. 		
		[Add in any addit	ional agree dissemination plans.]	
31.	Keywords	[Give words or pl	nrases that best describe the review.]	
32.	Details of existing review of same topic by same authors	[Give details of earlier versions of the systematic review if an update of an existing review is being registered, including full bibliographic reference if possible. NOTE: most NICE reviews will not constitute an update in PROSPERO language. To be an update it needs to be the same review question/search/methodology. If anything has changed it is a new review]		
33.	Current review status	\boxtimes	Ongoing	
			Completed but not published	
			Completed and published	
			Completed, published and being updated	
			Discontinued	
34.	Additional information	[Provide any other information the review team feel is relevant to the registration of the review.]		
35.	Details of final publication	www.nice.org.uk		

2 Review protocol for CT, MRI and X-ray of the cervical spine in people with head injury – diagnostic accuracy

ID Field		Content	
0.	PROSPERO registration number	CRD42021283526	

1.	Review title	What is the clinical and cost effectiveness of CT, MRI and X-ray of the cervical spine for initial imaging in people with head injury?
2.	Review question	What is the clinical and cost effectiveness of CT, MRI and X-ray of the cervical spine for initial imaging in people with head injury?
3.	Objective	To determine which of the currently available imaging techniques is best to diagnose cervical spine injury.
4.	Searches	The following databases (from inception) will be searched:
		Cochrane Central Register of Controlled Trials (CENTRAL)
		Cochrane Database of Systematic Reviews (CDSR)
		• Embase
		MEDLINE
		Searches will be restricted by:
		English language studies
		Human studies
		Letter and comments excluded
		Other searches:
		[Inclusion lists of systematic reviews]
		The searches may be re-run 6 weeks before the final committee meeting and further studies retrieved for inclusion if relevant.
		The full search strategies will be published in the final review.

Condition or domain being studied Population	Cervical spine injury in patients who have experienced a head injury. Inclusion: Infants, children and adult with head injury and suspected cervical spine
Population	Inclusion: Infante, childron and adult with head injuny and suspected conviced spine
	injury
	Strata:
	Adults at:
	 o high risk
	o moderate risk
	○ low risk
	Children + infants at:
	o high risk
	o moderate risk
	 o low risk
	• Adults (aged ≥16 years)
	• Children and infants (aged 0 to <16 years).
	Mixed population studies will be included and downgraded for indirectness. Cut- off of 60% will be used for all age groups.
	Exclusion: Adults, and children (including infants under 1 year) with superficial injuries to the eye or face without suspected or confirmed head or brain injury.

		C spine injury risk stratification based on: Canadian C Spine Rule NEXUS Stratification rules should be kept separate as they have different features
		Both for adults and children. Both rules validated in adults and extrapolated for children. These are not specific for head injury
7.	Intervention	 Computed tomography scan (CT) of cervical spine Magnetic resonance imaging (MRI) of cervical spine X-ray of cervical spine
8.	comparator	MRI of cervical spine, X-ray of cervical spine and CT of cervical spine compared to each other For X-ray only include children and people below 65 years.
		People >65 years are considered as high risk and will be offered CT cervical spine within 8 hours of injury (CG 176). Vascular injuries will be picked up by MR imaging
9.	Types of study to be included	
		 Diagnostic test and treat: Randomised controlled trials (RCTs), systematic reviews of RCTs. If no RCT evidence is available, non-randomised studies will be considered if they adjust for key confounders, starting with prospective cohort studies.

		Key confounders • Age • Gender • GCS or pupillary response at presentation
10.	Other exclusion criteria	Non-English language studies. Conference abstracts will be excluded as it is expected there will be sufficient full text published studies available.
11.	Context	Head-injured patient may also have sustained concomitant injury to the cervical spine. Some head injured patients who require a CT head scan will also need cervical spine imaging. The purpose of this review is to inform the optimal diagnostic pathways for these patients using the best evidence available.
12.	Primary outcomes (critical outcomes)	 All outcomes are considered equally important for decision making and therefore have all been rated as critical: Mortality at 3 months
		 Mortality at 3 months Quality of life - 3 months or more
		 Objectively applied score of disability e.g. Glasgow Outcome Score (GOS) or extended GOS - at 3 months or more
		Length of hospital stay
		Unscheduled re-admission (28 days or longer)
		Neurological deterioration

<u>г</u>		
		Neurological deterioration could be because of either no imaging or no appropriate imaging.
		Spinal injuries are determined by different scales– e.g. <u>American Spinal Injury</u> <u>Association</u> (ASIA), functional independence measure (FIM). Different scales are used. Report as in the studies
		Vascular insult would be picked up in outcome neurological deterioration
		COMET database was searched for relevant core outcome sets and none were identified.
13.	Data extraction (selection and coding)	All references identified by the searches and from other sources will be uploaded into EPPI reviewer and de-duplicated.
		10% of the abstracts will be reviewed by two reviewers, with any disagreements resolved by discussion or, if necessary, a third independent reviewer.
		This review will make use of the priority screening functionality within the EPPI- reviewer software.
		The full text of potentially eligible studies will be retrieved and will be assessed in line with the criteria outlined above.
		A standardised form will be used to extract data from studies (see <u>Developing</u> <u>NICE guidelines: the manual section 6.4</u>).
		10% of all evidence reviews are quality assured by a senior research fellow. This includes checking:
		 papers were included /excluded appropriately
		• a sample of the data extractions
		 correct methods are used to synthesise data
		 a sample of the risk of bias assessments

		Disagreements between the review authors over the risk of bias in particular studies will be resolved by discussion, with involvement of a third review author where necessary.
14.	Risk of bias (quality) assessment	 For diagnostic Test and treat: Risk of bias will be assessed using the appropriate checklist as described in Developing NICE guidelines: the manual. For Intervention reviews Systematic reviews: Risk of Bias in Systematic Reviews (ROBIS) Randomised Controlled Trial: Cochrane RoB (2.0) Non randomised study, including cohort studies: Cochrane ROBINS-I
15.	Strategy for data synthesis	 For diagnostic test and treat: Pairwise meta-analyses will be performed using Cochrane Review Manager (RevMan5). Fixed-effects (Mantel-Haenszel) techniques will be used to calculate risk ratios for the binary outcomes where possible. Continuous outcomes will be analysed using an inverse variance method for pooling weighted mean differences. Heterogeneity between the studies in effect measures will be assessed using the l² statistic and visually inspected. An l² value greater than 50% will be considered indicative of substantial heterogeneity. Sensitivity analyses will be conducted based on pre-specified subgroups using stratified meta-analysis to

16.	Analysis of sub-groups	heterogeneity, the GRADEpro outcome, taking int results. The 4 main imprecision) will be when there are mo The risk of outcome using an a Development and B GRADE working gr Where meta-analys individually per out Subgroups that will Types of injuries: Bone injuri Spinal core	I be investigated if heterogeneity is present: es d injuries issue injuries
17.	Type and method of review		Intervention
			Diagnostic
			Prognostic
			Qualitative
			Epidemiologic
			Service Delivery

		□ Other (please	specify)		
18.	Language	English	English		
19.	Country	England			
20.	Anticipated or actual start date	pated or actual start date [For the purposes of PROSPERO, the da review can be defined as any point after of formal screening of the identified studies		completion of a protocol but before	
		A protocol can be deemed complete responsibility for quality assurance		y the NICE team with	
21.	Anticipated completion date	[Give the date by which the guideline is expected to be published. This field may be edited at any time. All edits will appear in the record audit trail. A brief explanation of the reason for changes should be given in the Revision Notes facility.]			
22.	Stage of review at time of this submission	Review stage	Started	Completed	
		Preliminary searches			
		Piloting of the study selection process			
		Formal screening of search result against eligibility criteria	3 □		
		Data extraction			
		Risk of bias (quality) assessment			
		Data analysis			
23.	Named contact	5a. Named contact			

		National Guideline Centre
		5b Named contact e-mail
		[Guideline email]@nice.org.uk
		[Developer to check with Guideline Coordinator for email address]
		5e Organisational affiliation of the review
		National Institute for Health and Care Excellence (NICE) and [National Guideline Alliance / National Guideline Centre / NICE Guideline Updates Team / NICE Public Health Guideline Development Team] [Note it is essential to use the template text here and one of the centre options to enable PROSPERO to recognise this as a NICE protocol]
24.	Review team members	[Give the title, first name, last name and the organisational affiliations of each member of the review team. Affiliation refers to groups or organisations to which review team members belong.]
		From the National Guideline Centre:
		[Guideline lead]
		[Senior systematic reviewer]
		Systematic reviewer
		[Health economist]
		[Information specialist]
		[Others]
25.	Funding sources/sponsor	This systematic review is being completed by the National Guideline Centre which receives funding from NICE.

26.	Conflicts of interest	All guideline committee members and anyone who has direct input into NICE guidelines (including the evidence review team and expert witnesses) must declare any potential conflicts of interest in line with NICE's code of practice for declaring and dealing with conflicts of interest. Any relevant interests, or changes to interests, will also be declared publicly at the start of each guideline committee meeting. Before each meeting, any potential conflicts of interest will be considered by the guideline committee Chair and a senior member of the development team. Any decisions to exclude a person from all or part of a meeting will be documented. Any changes to a member's declaration of interests will be recorded in the minutes of the meeting. Declarations of interests will be published with the final guideline.
27.	Collaborators	Development of this systematic review will be overseen by an advisory committee who will use the review to inform the development of evidence-based recommendations in line with section 3 of <u>Developing NICE guidelines: the manual</u> . Members of the guideline committee are available on the NICE website: [NICE guideline webpage].
28.	Other registration details	[Give the name of any organisation where the systematic review title or protocol is registered (such as with The Campbell Collaboration, or The Joanna Briggs Institute) together with any unique identification number assigned. If extracted data will be stored and made available through a repository such as the Systematic Review Data Repository (SRDR), details and a link should be included here. If none, leave blank.]
29.	Reference/URL for published protocol	[Give the citation and link for the published protocol, if there is one.]
30.	Dissemination plans	NICE may use a range of different methods to raise awareness of the guideline. These include standard approaches such as:
		 notifying registered stakeholders of publication
		 publicising the guideline through NICE's newsletter and alerts
		 issuing a press release or briefing as appropriate, posting news articles on the NICE website, using social media channels, and publicising the guideline within NICE.

		[Add in any additi	[Add in any additional agree dissemination plans.]		
31.	Keywords	[Give words or ph	[Give words or phrases that best describe the review.]		
32.	Details of existing review of same topic by same authors	[Give details of earlier versions of the systematic review if an update of an existing review is being registered, including full bibliographic reference if possible. NOTE: most NICE reviews will not constitute an update in PROSPERO language. To be an update it needs to be the same review question/search/methodology. If anything has changed it is a new review]			
33.	Current review status	\boxtimes	Ongoing		
			Completed but not published		
			Completed and published		
			Completed, published and being updated		
			Discontinued		
34.	Additional information	[Provide any other information the review team feel is relevant to the registration of the review.]			
35.	Details of final publication	www.nice.org.uk			

2 Health economic review protocol

Review question	All questions – health economic evidence
Objective	To identify health economic studies relevant to any of the review questions.
Search criteria	 Populations, interventions and comparators must be as specified in the clinical review protocol above. Studies must be of a relevant health economic study design (cost–utility analysis, cost-effectiveness analysis, cost–benefit analysis, cost–consequences analysis, comparative cost analysis). Studies must not be a letter, editorial or commentary, or a review of health economic evaluations. (Recent reviews will be
	ordered although not reviewed. The bibliographies will be checked for relevant studies, which will then be ordered.)

- Unpublished reports will not be considered unless submitted as part of a call for evidence.
- Studies must be in English.

Search A health economic study search will be undertaken using population-specific terms and a health economic study filter – see appendix B below. The search covered all years

Review Studies not meeting any of the search criteria above will be excluded. Studies published before 2006, abstract-only studies and studies from non-OECD countries or the USA will also be excluded.

Studies published in 2006 or later that were included in the previous guidelines will be reassessed for inclusion and may be included or selectively excluded based on their relevance to the questions covered in this update and whether more applicable evidence is also identified.

Each remaining study will be assessed for applicability and methodological limitations using the NICE economic evaluation checklist which can be found in appendix H of Developing NICE guidelines: the manual (2014).³²

Inclusion and exclusion criteria

- If a study is rated as both 'Directly applicable' and with 'Minor limitations' then it will be included in the guideline. A health economic evidence table will be completed and it will be included in the health economic evidence profile.
- If a study is rated as either 'Not applicable' or with 'Very serious limitations' then it will usually be excluded from the guideline. If it is excluded then a health economic evidence table will not be completed and it will not be included in the health economic evidence profile.
- If a study is rated as 'Partially applicable', with 'Potentially serious limitations' or both then there is discretion over whether it should be included.

Where there is discretion

The health economist will make a decision based on the relative applicability and quality of the available evidence for that question, in discussion with the guideline committee if required. The ultimate aim is to include health economic studies that are helpful for decision-making in the context of the guideline and the current NHS setting. If several studies are considered of sufficiently high applicability and methodological quality that they could all be included, then the health economist, in discussion with the committee if required, may decide to include only the most applicable studies and to selectively exclude the remaining studies. All studies excluded on the basis of applicability or methodological limitations will be listed with explanation in the excluded health economic studies appendix below.

The health economist will be guided by the following hierarchies.

Setting:

- UK NHS (most applicable).
- OECD countries with predominantly public health insurance systems (for example, France, Germany, Sweden).
- OECD countries with predominantly private health insurance systems (for example, Switzerland).
- Studies set in non-OECD countries or in the USA will be excluded before being assessed for applicability and methodological limitations.

Health economic study type:

- Cost-utility analysis (most applicable).
- Other type of full economic evaluation (cost-benefit analysis, cost-effectiveness analysis, cost-consequences analysis).
- Comparative cost analysis.
- Non-comparative cost analyses including cost-of-illness studies will be excluded before being assessed for applicability and methodological limitations.

Year of analysis:

- The more recent the study, the more applicable it will be.
- Studies published in 2006 or later (including any such studies included in the previous guidelines) but that depend on unit costs and resource data entirely or predominantly from before 2006 will be rated as 'Not applicable'.
- Studies published before 2006 (including any such studies included in the previous guidelines) will be excluded before being assessed for applicability and methodological limitations.

Quality and relevance of effectiveness data used in the health economic analysis:

• The more closely the clinical effectiveness data used in the health economic analysis match with the outcomes of the studies included in the clinical review the more useful the analysis will be for decision-making in the guideline.

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Appendix B – Literature search strategies

- 2 This literature search strategy was used for the following questions:
 - What is the diagnostic accuracy of CT, MRI and X-ray of the cervical spine for initial imaging in people with head injury?
 - What is the clinical and cost effectiveness of CT, MRI and X-ray of the cervical spine for initial imaging in people with head injury?
- The literature searches for this review are detailed below and complied with the methodology
 outlined in Developing NICE guidelines: the manual.³²
- 9 For more information, please see the Methodology review published as part of the
- 10 accompanying documents for this guideline.

B₁1 Clinical search literature search strategy

- 12 Searches were constructed using a PICO framework where population (P) terms were
- 13 combined with Intervention (I) and in some cases Comparison (C) terms. Outcomes (O) are
- 14 rarely used in search strategies as these concepts may not be indexed or described in the
- 15 title or abstract and are therefore difficult to retrieve.

16 **Table 29: Database parameters, filters and limits applied**

Database	Dates searched	Search filter used			
Medline (OVID)	1946 – 22 June 2022	Exclusions (animal studies, letters, comments, editorials, case studies/reports) English language			
Embase (OVID)	1974 – 22 June 2022	Exclusions (animal studies, letters, comments, editorials, case studies/reports, conference abstracts) English language			
The Cochrane Library (Wiley)	Cochrane Reviews to 2022 Issue 6 of 12 CENTRAL to 2022 Issue 6 of 12				

17 Medline (Ovid) search terms

exp Spinal Injuries/
Spinal Cord Injuries/
exp Neck Injuries/
whiplash.ti,ab.
((neck or spine or spinal) adj3 (trauma or injur* or fracture*)).ti,ab.
or/1-5
cervical.ti,ab.
6 and 7
(cervical adj3 (trauma* or injur* or fracture*)).ti,ab.
8 or 9
letter/

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12.	editorial/
13.	news/
14.	exp historical article/
15.	Anecdotes as Topic/
16.	comment/
17.	case report/
18.	(letter or comment*).ti.
19.	or/11-18
20.	randomized controlled trial/ or random*.ti,ab.
21.	19 not 20
22.	animals/ not humans/
23.	exp Animals, Laboratory/
24.	exp Animal Experimentation/
25.	exp Models, Animal/
26.	exp Rodentia/
27.	(rat or rats or mouse or mice or rodent*).ti.
28.	or/21-27
29.	10 not 28
30.	limit 29 to English language
31.	tomography/
32.	exp magnetic resonance imaging/
33.	exp tomography, emission-computed/
34.	exp tomography, x-ray/
35.	Radiography/
36.	Neuroradiography/
37.	(compute* adj2 tomograph*).ti,ab.
38.	(CT or CAT or PET or SPECT).ti,ab.
39.	((MR or magnetic resonance or NMR) adj2 (imag* or tomograph* or angiograph*)).ti,ab.
40.	MRI.ti,ab.
41.	(radiograph* or xray* or x-ray* or x ray*).ti,ab.
42.	or/31-41
43.	30 and 42

18 Embase (Ovid) search terms

1.	spine injury/
2.	cervical spine injury/
3.	spinal cord injury/
4.	cervical spinal cord injury/
5.	neck injury/
6.	whiplash injury/
7.	whiplash.ti,ab.
8.	((neck or spine or spinal) adj3 (trauma or injur*)).ti,ab.
9.	or/1-8
10.	letter.pt. or letter/
11.	note.pt.

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12.	editorial.pt.
13.	(conference abstract or conference paper).pt.
14.	case report/ or case study/
15.	(letter or comment*).ti.
16.	or/10-15
17.	randomized controlled trial/ or random*.ti,ab.
18.	16 not 17
19.	animal/ not human/
20.	nonhuman/
21.	exp Animal Experiment/
22.	exp Experimental Animal/
23.	animal model/
24.	exp Rodent/
25.	(rat or rats or mouse or mice or rodent*).ti.
26.	or/18-25
27.	9 not 26
28.	limit 27 to English language
29.	tomography/
30.	brain tomography/
31.	exp computer assisted tomography/
32.	exp emission tomography/
33.	exp nuclear magnetic resonance imaging/
34.	radiography/
35.	(compute* adj2 tomograph*).ti,ab.
36.	(CT or CAT or PET or SPECT).ti,ab.
37.	((MR or magnetic resonance or NMR) adj2 (imag* or tomograph* or angiograph*)).ti,ab.
38.	MRI.ti,ab.
39.	(radiograph* or xray* or x-ray* or x ray*).ti,ab.
40.	neuroradiology/ or brain radiography/
41.	or/29-40
42.	28 and 41

19 Cochrane Library (Wiley) search terms

#1.	MeSH descriptor: [Spinal Injuries] explode all trees
#2.	MeSH descriptor: [Spinal Cord Injuries] this term only
#3.	MeSH descriptor: [Neck Injuries] explode all trees
#4.	whiplash:ti,ab
#5.	((neck or spine or spinal) near/3 (trauma or injur* or fracture*)):ti,ab
#6.	#1 or #2 or #3 or #4 or #5
#7.	cervical:ti,ab
#8.	#6 and #7
#9.	(cervical near/3 (trauma* or injur* or fracture*)):ti,ab
#10.	#8 or #9
#11.	MeSH descriptor: [Tomography] this term only
#12.	MeSH descriptor: [Magnetic Resonance Imaging] explode all trees

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#13.	MeSH descriptor: [Tomography, Emission-Computed] explode all trees	
#14.	MeSH descriptor: [Tomography, X-Ray] explode all trees	
#15.	MeSH descriptor: [Radiography] this term only	
#16.	MeSH descriptor: [Neuroradiography] this term only	
#17.	(compute* near/2 tomograph*):ti,ab	
#18.	(CT or CAT or PET or SPECT):ti,ab	
#19.	((MR or magnetic resonance or NMR) near/2 (imag* or tomograph* or angiograph*)):ti,ab	
#20.	MRI:ti,ab	
#21.	(radiograph* or xray* or x-ray* or x ray*):ti,ab	
#22.	#11 or #12 or #13 or #14 or #15 or #16 or #17 or #18 or #19 or #20 or #21	
#23.	#10 and #22	

B22 Health Economics literature search strategy

21 Health economic evidence was identified by conducting searches using terms for a broad

Head Injury population. The following databases were searched: NHS Economic Evaluation

23 Database (NHS EED - this ceased to be updated after 31st March 2015), Health Technology

Assessment database (HTA - this ceased to be updated from 31st March 2018) and The

25 International Network of Agencies for Health Technology Assessment (INAHTA). Searches

for recent evidence were run on Medline and Embase from 2014 onwards for health

economics, and all years for quality-of-life studies.

28 Table 30: Database parameters, filters and limits applied

Database	Dates searched	Search filters and limits applied
Medline (OVID)	Health Economics 1 January 2014 – 22 June 2022	Health economics studies Quality of life studies Exclusions (animal studies,
	Quality of Life 1946 – 22 June 2022	letters, comments, editorials, case studies/reports) English language
		5 5 5
Embase (OVID)	Health Economics 1 January 2014 – 22 June 2022	Health economics studies Quality of life studies
	Quality of Life 1974 – 22 June 2022	Exclusions (animal studies, letters, comments, editorials, case studies/reports, conference abstracts)
		English language
NHS Economic Evaluation Database (NHS EED) (Centre for Research and Dissemination - CRD)	Inception –31 st March 2015	
Health Technology Assessment Database (HTA)	Inception – 31 st March 2018	

Database	Dates searched	Search filters and limits applied
(Centre for Research and Dissemination – CRD)		
The International Network of Agencies for Health Technology Assessment (INAHTA)	Inception – 22 June 2022	English language

29 Medline (Ovid) search terms

1.	craniocerebral trauma/ or exp brain injuries/ or coma, post-head injury/ or exp head injuries, closed/ or head injuries, penetrating/ or exp intracranial hemorrhage, traumatic/ or exp skull fractures/
2.	((skull or cranial) adj3 fracture*).ti,ab.
3.	((head or brain or craniocerebral or intracranial or cranial or skull) adj3 (injur* or trauma*)).ti,ab.
4.	(trauma* and ((subdural or intracranial or brain) adj2 (h?ematoma* or h?emorrhage* or bleed*))).ti,ab.
5.	or/1-4
6.	letter/
7.	editorial/
8.	news/
9.	exp historical article/
10.	Anecdotes as Topic/
11.	comment/
12.	case report/
13.	(letter or comment*).ti.
14.	or/6-13
15.	randomized controlled trial/ or random*.ti,ab.
16.	14 not 15
17.	animals/ not humans/
18.	exp Animals, Laboratory/
19.	exp Animal Experimentation/
20.	exp Models, Animal/
21.	exp Rodentia/
22.	(rat or rats or mouse or mice or rodent*).ti.
23.	or/16-22
24.	5 not 23
25.	limit 24 to English language
26.	economics/
27.	value of life/
28.	exp "costs and cost analysis"/
29.	exp Economics, Hospital/
30.	exp Economics, medical/
31.	Economics, nursing/

32.	economics, pharmaceutical/
33.	exp "Fees and Charges"/
34.	exp budgets/
35.	budget*.ti,ab.
36.	cost*.ti.
37.	(economic* or pharmaco?economic*).ti.
38.	(price* or pricing*).ti,ab.
39.	(cost* adj2 (effectiv* or utilit* or benefit* or minimi* or unit* or estimat* or variable*)).ab.
40.	(financ* or fee or fees).ti,ab.
41.	(value adj2 (money or monetary)).ti,ab.
42.	or/26-41
43.	quality-adjusted life years/
44.	sickness impact profile/
45.	(quality adj2 (wellbeing or well being)).ti,ab.
46.	sickness impact profile.ti,ab.
47.	disability adjusted life.ti,ab.
48.	(qal* or qtime* or qwb* or daly*).ti,ab.
49.	(euroqol* or eq5d* or eq 5*).ti,ab.
50.	(qol* or hql* or hqol* or h qol* or hrqol* or hr qol*).ti,ab.
51.	(health utility* or utility score* or disutilit* or utility value*).ti,ab.
52.	(hui or hui1 or hui2 or hui3).ti,ab.
53.	(health* year* equivalent* or hye or hyes).ti,ab.
54.	discrete choice*.ti,ab.
55.	rosser.ti,ab.
56.	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.
57.	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.
58.	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.
59.	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.
60.	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab.
61.	(sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.
62.	or/32-61
63.	25 and (42 or 62)

30 Embase (Ovid) search terms

1.	head injury/
2.	exp brain injury/
3.	skull injury/ or exp skull fracture/
4.	((head or brain or craniocerebral or intracranial or cranial or skull) adj3 (injur* or trauma*)).ti,ab.
5.	((skull or cranial) adj3 fracture*).ti,ab.
6.	(trauma* and ((subdural or intracranial or brain) adj2 (h?ematoma* or h?emorrhage* or bleed*))).ti,ab.

7.	or/1-6
8.	letter.pt. or letter/
9.	note.pt.
10.	editorial.pt.
11.	(conference abstract or conference paper).pt.
12.	case report/ or case study/
13.	(letter or comment*).ti.
14.	or/8-13
15.	randomized controlled trial/ or random*.ti,ab.
16.	14 not 15
17.	animal/ not human/
18.	nonhuman/
19.	exp Animal Experiment/
20.	exp Experimental Animal/
21.	animal model/
22.	exp Rodent/
23.	(rat or rats or mouse or mice or rodent*).ti.
24.	or/16-23
25.	7 not 24
26.	limit 25 to English language
27.	health economics/
28.	exp economic evaluation/
29.	exp health care cost/
30.	exp fee/
31.	budget/
32.	funding/
33.	budget*.ti,ab.
34.	cost*.ti.
35.	(economic* or pharmaco?economic*).ti.
36.	(price* or pricing*).ti,ab.
37.	(cost* adj2 (effectiv* or utilit* or benefit* or minimi* or unit* or estimat* or variable*)).ab.
38.	(financ* or fee or fees).ti,ab.
39.	(value adj2 (money or monetary)).ti,ab.
40.	or/27-39
41.	quality-adjusted life years/
42.	"quality of life index"/
43.	short form 12/ or short form 20/ or short form 36/ or short form 8/
44.	sickness impact profile/
45.	(quality adj2 (wellbeing or well being)).ti,ab.
46.	sickness impact profile.ti,ab.
47.	disability adjusted life.ti,ab.
48.	(qal* or qtime* or qwb* or daly*).ti,ab.

49.	(euroqol* or eq5d* or eq 5*).ti,ab.
50.	(qol* or hql* or hqol* or h qol* or hrqol* or hr qol*).ti,ab.
51.	(health utility* or utility score* or disutilit* or utility value*).ti,ab.
52.	(hui or hui1 or hui2 or hui3).ti,ab.
53.	(health* year* equivalent* or hye or hyes).ti,ab.
54.	discrete choice*.ti,ab.
55.	rosser.ti,ab.
56.	(willingness to pay or time tradeoff or time trade off or tto or standard gamble*).ti,ab.
57.	(sf36* or sf 36* or short form 36* or shortform 36* or shortform36*).ti,ab.
58.	(sf20 or sf 20 or short form 20 or shortform 20 or shortform20).ti,ab.
59.	(sf12* or sf 12* or short form 12* or shortform 12* or shortform12*).ti,ab.
60.	(sf8* or sf 8* or short form 8* or shortform 8* or shortform8*).ti,ab.
61.	(sf6* or sf 6* or short form 6* or shortform 6* or shortform6*).ti,ab.
62.	or/41-61
63.	26 and (40 or 62)

31 NHS EED and HTA (CRD) search terms

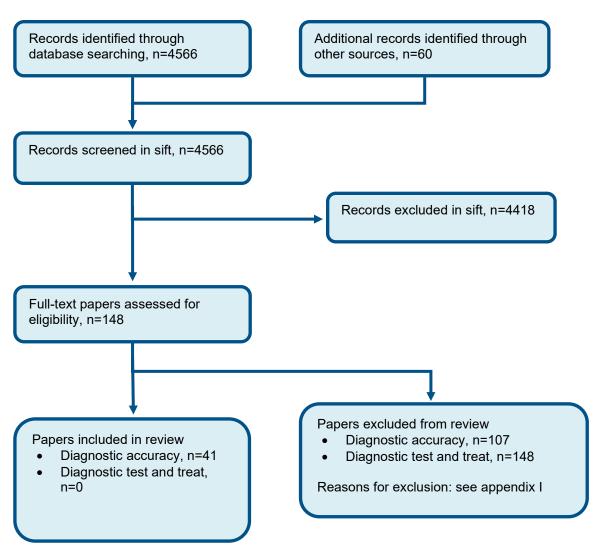
#1.	MeSH DESCRIPTOR Brain Injuries EXPLODE ALL TREES
#2.	MeSH DESCRIPTOR Craniocerebral Trauma
#3.	MeSH DESCRIPTOR Coma, Post-Head Injury
#4.	MeSH DESCRIPTOR Head Injuries, Closed EXPLODE ALL TREES
#5.	MeSH DESCRIPTOR Head Injuries, Penetrating
#6.	MeSH DESCRIPTOR Intracranial Hemorrhage, Traumatic EXPLODE ALL TREES
#7.	MeSH DESCRIPTOR Skull Fractures EXPLODE ALL TREES
#8.	(((skull or cranial) adj3 fracture*))
# 9.	(((head or brain or craniocerebral or intracranial or cranial or skull) adj3 (injur* or trauma*)))
#10.	((trauma* and ((subdural or intracranial or brain) adj2 (h?ematoma* or h?emorrhage* or bleed*))))
#11.	#1 OR #2 OR #3 OR #4 OR #5 OR #6 OR #7 OR #8 OR #9 OR #10

32 INAHTA search terms

1.	((((trauma* and ((subdural or intracranial or brain) and (haematoma* or hematoma* or haemorrhage* or hemorrhage* or bleed*))))[Title]) AND (((trauma* and ((subdural or intracranial or brain) and (haematoma* or hematoma* or haemorrhage* or
	hemorrhage* or bleed*))))[Title])) OR ((((skull or cranial) and fracture*))[Title] OR
	(((skull or cranial) and fracture*))[abs]) OR ((((head or brain or craniocerebral or
	intracranial or cranial or skull) and (injur* or trauma*)))[Title] OR (((head or brain or
	craniocerebral or intracranial or cranial or skull) and (injur* or trauma*)))[abs]) OR
	("Skull Fractures"[mhe]) OR ("Intracranial Hemorrhage, Traumatic"[mhe]) OR ("Head
	Injuries, Penetrating"[mh]) OR ("Head Injuries, Closed"[mhe]) OR ("Coma, Post-Head
	Injury"[mh]) OR ("Brain Injuries"[mhe]) OR ("Craniocerebral Trauma"[mh])

34 Appendix C – Diagnostic evidence study selection

Figure 1: Flow chart of clinical study selection for the review of CT, MRI and X-ray of the cervical spine in people with head injury



1

2 Appendix D Diagnostic evidence

3 Note that some evidence tables may contain outcomes that were not eventually included in the analysis, for example for studies that reported both

4 any severity of cervical spine injury and clinically significant/unstable injuries separately, the latter was used for analysis and the results for any

5 injury severity not used as this was closest to the target condition in the protocol.

D.a Adults

Reference	Adams 2006 ¹
Study type	Retrospective chart review
Study methodology	Data source: chart review of CT and MRI reports and changes in clinical management as part of Morristown Memorial Hospital Trauma Quality Improvement Initiative
	Recruitment: records of those within a 12 month period (January 1 st 2004 to December 31 st 2004) matching inclusion criteria were included
Number of patients	n = 97
	(n=99 undergoing MRI cervical spine identified and n=97 charts were complete and available for review)
Patient characteristics	Age, mean (SD): 40 (21) years
	Gender (male to female ratio): 71.1% male and 28.9% female
	Ethnicity: not reported

Reference	Adams 2006 ¹
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Injury severity score, mean (SD): 15 (11) in whole population and 24 (9) in obtunded group
	Mechanism of injury:
	 Motor vehicle crash, 45% Falls, 44% Pedestrian struck, 6% Assaults, 4%
	Setting: secondary care – hospital/trauma centre
	Country: USA
	Inclusion criteria: underwent MRI cervical spine trauma protocol; deemed high-risk for axial trauma due to pain, neurologic symptoms or obtundation after significant blunt trauma; and complete chart data available for review.
	Exclusion criteria: not reported

Reference	Adams 2006 ¹
	Adults with high suspicion of axial trauma undergoing MRI of cervical spine
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided (assumed head injury based on severity of injuries – all at high suspicion of axial trauma)
Index test(s) and	Index test
reference standard	CT of cervical spine
	Performed using GE light speed 4-slice CT scanner with collimation of 5 mm, pitch of 1.6. and reconstructions at 1 mm of image spacing from base of skull to first thoracic vertebrae.
	Reference standard
	Final diagnosis based on MRI and CT and clinical decision-making of spinal consultants, no follow-up mentioned.
	MRI of cervical spine performed without contrast. Sagittal T1- and T2-weighted images acquired from posterior fossa through 5 th thoracic vertebrae. 3 mm thin section contiguous axial and sagittal T1- and T2-weighted images obtained from 2 nd cervical vertebrae through 1 st thoracic vertebrae. When available, comparison was made with the CT of cervical spine by attending radiologist.
	All imaging studies evaluated based on radiology department protocols. After initial review by radiology residents, attending radiologist then read studies. No attempt at blinding radiology staff to results of CT or MRI was made and final printed radiology report accepted as official reading of the study. Presence or absence of acute spinal injury based on official MRI and CT scan reports and clinical decision-making of spinal consultants. Areas of discrepancy between CT and MRI were subjected to formal interrogation by dedicated thin cut CT imaging on the level in question.
	Time between measurement of index test and reference standard: unclear time interval between index test and subsequent tests/final confirmed diagnosis

Reference	Adams 2006 ¹
Outcome	Cervical spine injury – poorly defined
2×2 table	Raw data not reported to allow 2x2 tables to be calculated
Statistical measures	Cervical spine injury – CT as index test: reported in paper Sensitivity: 94.0% Specificity: 88.0% PPV: NR NPV: NR
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, index test and reference standard likely interpreted with knowledge of the other, unclear time interval between index test and reference standard and unclear if reference standard components differed between patients
	general population of those attending ED with suspected cervical spine injury, and reference standard possibly places focus on MRI results with it also unclear if follow-up included 2 weeks
Comments	None

Reference	Bailitz 2009 ³
Study type	Prospective observational study
Study methodology	Data source: conducted at single hospital trauma unit
	Recruitment: consecutive patients presenting to Cook County Hospital Trauma Unit meeting inclusion criteria between December 15 th 2004 and November 15 th 2006.
Number of patients	n = 1505
	(n=1583 had cervical spine trauma and n=78 patients were excluded as they did not have both CT and radiography, leaving n=1505 patients)
Patient characteristics	Age, mean (SD): 37 (SD not reported) years
	Gender (male to female ratio): 72% male and 28% female
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury:
	 Motor vehicle collisions, 40% Assault, 25% Fall, 20%

Reference	Bailitz 2009 ³
	Pedestrian struck by car, 9%
	Setting: secondary care – hospital trauma unit
	Country: USA
	Inclusion criteria: meeting at least one of NEXUS criteria and therefore requiring cervical spine imaging for vertebral bone blunt cervical trauma injury (criteria were: midline pain or tenderness, neurologic findings, altered mental status, intoxication and distracting injury); and had both CT and X-ray performed.
	Exclusion criteria: <16 years
	People meeting at least one NEXUS criterion and suspicion of cervical spine injury – CT and X-ray performed
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	X-ray of cervical spine – three-view
	CT of cervical spine

Reference	Bailitz 2009 ³				
	Final readings for index tests were performed by two attending radiologists provided with same basic clinical information while blinded to the results of other imaging study and earlier preliminary readings. No further details provided for index tests				
	Reference standa	ard			
	Final diagnosis in	medical record at disc	harge		
		easurement of index tes /final confirmed diagnos		rd: unclear time in	terval between index tests and
Outcome	Clinically significa	ant cervical spine injury	 reported separately f 	or different risk str	ata (NEXUS) as in protocol
	Clinically significant injuries were defined as those requiring one or more of following interventions recommended by neurosurgical consultation: operative procedure, halo application and/or rigid cervical collar.				
2×2 table	Clinically significant cervical spine injury – X-ray as index test (high risk)				
		Reference standard +	Reference standard	Total	Raw data reported insufficiently meaning
	Index test +	7	NR	NR	specificity could not be calculated and is not reported
	Index test -	8	NR	NR	
	Total	15	NR	NR	
	Clinically signifi	cant cervical spine in	jury – X-ray as index 1	est (moderate ris	k)

Reference	Bailitz 2009 ³				
		Reference standard +	Reference standard	Total	Raw data reported insufficiently meaning
	Index test +	7	NR	NR	specificity could not be calculated and is not reported
	Index test -	12	NR	NR	
	Total	19	NR	NR	
	Clinically signif	icant cervical spine in	jury – X-ray as index t	est (low risk)	
		Reference standard +	Reference standard -	Total	Raw data reported insufficiently meaning
	Index test +	4	NR	NR	specificity could not be calculated and is not reported
	Index test -	12	NR	NR	
	Total	16	NR	NR	
	Clinically signif	icant cervical spine in	jury – CT as index tes	t (high risk)	
		Reference standard +	Reference standard -	Total	Raw data reported insufficiently meaning
	Index test +	15	NR	NR	specificity could not be calculated and is not reported
	Index test -	0	NR	NR	
	Total	15	NR	NR	

Reference	Bailitz 2009 ³				
	Clinically significant cervical spine injury – CT as index test (moderate risk)				
		Reference standard +	Reference standard –	Total	Raw data reported insufficiently meaning
	Index test +	19	NR	NR	specificity could not be calculated and is not reported
	Index test -	0	NR	NR	
	Total	19	NR	NR	
	Clinically significant cervical spine injury – CT as index test (low risk)				
		Reference standard +	Reference standard -	Total	Raw data reported insufficiently meaning specificity could not be calculated and is not reported
	Index test +	16	NR	NR	
	Index test -	0	NR	NR	
	Total	16	R	NR	
Statistical measures	Clinically signifi	cant cervical spine in	jury – X-ray as index t	e st (high risk) : re	ported in paper
	Sensitivity: 46.0%	, 0			
	Specificity: NR				
	PPV: NR				

Reference	Bailitz 2009 ³
	NPV: NR
	Clinically significant cervical spine injury – X-ray as index test (moderate risk): reported in paper
	Sensitivity: 37.0%
	Specificity: NR
	PPV: NR
	NPV: NR
	Clinically significant cervical spine injury – X-ray as index test (low risk): reported in paper
	Sensitivity: 25.0%
	Specificity: NR
	PPV: NR
	NPV: NR
	Clinically significant cervical spine injury – CT as index test (same for high, moderate and low risk as all injuries detected): reported in paper
	Sensitivity: 100.0%
	Specificity: NR
	PPV: NR

Reference	Bailitz 2009 ³
	NPV: NR
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if reference standard interpreted without knowledge of index test, unclear interval between index test and reference standard and unclear if reference standard contained the same components for all patients – applies to both index tests
	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, and unclear if reference standard included a 2 week follow-up period
Comments	None
Reference	Berne 1999⁴
Study type	Prospective study
Study methodology	Data source: study performed at level 1 academic urban trauma centre

Study methodology	Data source: study performed at level 1 academic urban trauma centre
	Recruitment: performed over an 8 month period (November 1996 to June 1997).
Number of patients	n = 58
Number of patients	11 - 56
	(n=67 met inclusion criteria but n=9 were excluded as they did not get both plain radiography/X-ray and CT of the
	cervical spine, leaving n=58 analysed)
Patient characteristics	Age, mean (range): 43.1 (17-87) years
	Gender (male to female ratio): not reported
	Centrel (male to tentale). Not reported

Reference	Berne 1999⁴
	Ethnicity: not reported Head injury: 53% had associated head injury (intracranial bleed), unclear if remaining suffered some form of head injury as part of the injury mechanism Mechanism of injury: • Motor vehicle crash, 36.2%
	 Fall from a height, 24.1% Auto vs. pedestrian, 27.6% Auto vs. bicycle, 3.4% Motorcycle crash, 3.4% Assault, 3.4% Hanging, 1.7%
	Injury severity score, mean (range): 24.1 (4-66)
	Intubation: • In the field, 8.6% • In ED, 74.1%

Reference	Berne 1999⁴
	Associated injuries: Head (intracranial bleed), 53.4% Thoracic, 29.3% Abdominal, 15.5% Pelvic fracture, 5.2% Spinal injury, 8.6% Upper extremity, 6.9% Lower extremity, 10.3%
	Setting: secondary care – trauma centre
	Country: USA
	Inclusion criteria: high-risk blunt trauma; inability to evaluate patient's cervical spine clinically due to head injury, shock, alcohol or illicit drug use or pharmacological sedation and/or paralysis; need for CT scan of another body areas besides the cervical spine; and need for intensive care unit admission.
	Exclusion criteria: haemodynamic instability preventing transportation to CT suite; pregnancy; age <17 years; and/or identification of a surgical emergency while scanning another area (for example, mass lesion on head CT scan).
	High risk blunt trauma with suspected cervical spine injury

Reference	Berne 1999⁴
Target condition(s)	Suspected cervical spine injury – 53% had associated head injury (intracranial bleed), unclear if remaining suffered some form of head injury as part of the injury mechanism (assumed head injury based on severity of injuries – all admitted to ICU/at high risk)
Index test(s) and	Index test
reference standard	X-ray of cervical spine (data not included further in review as not relevant as initial imaging in severely injured)
	CT of cervical spine
	Complete cervical helical CT scan including all seven cervical vertebrae and first thoracic vertebrae performed when patient sent to CT suite to evaluate other body areas. Data on associated injuries, time from admission to CT scan, type and effects of adverse events occurring in CT scanner and CT readings by attending radiologist blinded to cases collected.
	Reference standard
	Final diagnosis based on all imaging/studies, including plain radiography even when CT used as index test, unclear duration of follow-up. Initially underwent plain radiography (X-ray) followed by complete cervical helical CT scan. May include clinical examination findings for some.
	Radiological study considered positive if diagnostic or suspicious for cervical spine injury. True positive defined if attending radiologist considered study to be diagnostic for injury or when confirmed by complementary imaging (CT or plain radiography) for suspicious films. If complementary CT or plain radiography did not confirm suspicious studies an additional radiological study (MRI or flexion-extension films) or subsequent clinical examination where appropriate (recovery of normal sensorium) was used to correlate initial radiological findings. False positives were those where initial films were suspicious but not diagnosed by complementary radiological studies or clinical examination. A study was negative where no cervical spine injury was identified. True

Reference	Berne 1999⁴					
	negatives were those where all radiological studies performed failed to reveal an abnormality suspicious for injury. False negative was when a complementary study in same patient revealed a previously unrecognised abnormality that was diagnostic for cervical spine injury.					
	Time between measurement of index test and reference standard: unclear time interval between index tests and subsequent tests/final confirmed diagnosis					
Outcome	Any cervical spine	e injury (stable and uns	table)			
	and Unstable cervical spine injury – classified as unstable in consultation with combined neurosurgical-orthopaedic					
	spine service and based on published guidelines					
2×2 table	Any cervical spi	Any cervical spine injury – X-ray as index test				
		Reference standard +	Reference standard -	Total		
	Index test +	12	0	12		
	Index test -	8	38	46		
	Total	20	38	58		
	Any cervical spine injury – CT as index test					

Reference	Berne 1999⁴				
		Reference standard +	Reference standard -	Total	
	Index test +	18	0	18	
	Index test -	2	38	40	
	Total	20	38	58	
	Unstable cervica	al spine injury – X-ray	as index test		
		Reference standard +	Reference standard -	Total	Raw data not provided so calculated using excel sheet from sensitivity/specificity etc. reported in the paper
	Index test +	5	0	5	
	Index test -	3	50	53	
	Total	8	50	58	
	Unstable cervica	al spine injury – CT as	index test		
		Reference standard +	Reference standard	Total	Raw data not provided so calculated using excel sheet
	Index test +	8	0	8	from sensitivity/specificity etc. reported in the paper
	Index test -	0	50	50	
	Total	8	50	58	

Reference	Berne 1999⁴
Statistical measures	Any cervical spine injury – X-ray as index test: reported in paper
	Sensitivity: 60.0%
	Specificity: 100.0%
	PPV: 100.0%
	NPV: 82.6%
	Any cervical spine injury – CT as index test: reported in paper
	Sensitivity: 90.0%
	Specificity: 100.0%
	PPV: 100.0%
	NPV: 95.0%
	Unstable cervical spine injury – X-ray as index test: reported in paper
	Sensitivity: 62.5%
	Specificity: 100.0%
	PPV: 100.0%
	NPV: 92.7%

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Unstable cervical spin Sensitivity: 100.0%	e injury – CT as index test: reported in paper
	e injury – CT as index test: reported in paper
Sensitivity: 100.0%	
Specificity: 100.0%	
PPV: 100.0%	
NPV: 100.0%	
Source of funding Not reported	
without knowledge of in	is – unclear if consecutive sample enrolled, unclear if reference standard interpreted dex test, unclear time interval between index test and reference standard and likely that ponents differed between patients – applies to both index tests
subgroup which may be	us – all included were at high-risk/admitted to ICU representing a more severely injured less applicable to general population of those attending ED with suspected cervical r if reference standard included a 2 week follow-up period
Comments None	

Reference	Brohi 2005 ⁶
Study type	Retrospective
Study methodology	Data source: people undergoing new spinal assessment protocol from single hospital
	Recruitment: new protocol introduced in February 2002 and patients included in the study between then and January 2004.
Number of patients	n = 421 analysed for X-ray and n=381 analysed for CT
	(n=442 considered relevant to review population; for X-ray, those without both CT and X-ray of cervical spine were excluded, leaving n=421; for CT, those without both CT and clinical outcome/MRI were excluded, leaving n=381)
Patient characteristics	Age, median (IQR): 34 (25-50) years
	Gender (male to female ratio): 2.6:1
	Ethnicity: not reported
	Head injury: unclear if all or most had head injury as no details provided
	Setting: secondary care – intubated trauma patients in hospital

Reference	Brohi 2005 ⁶
	Country: UK
	Inclusion criteria: unconscious, intubated trauma patients
	Exclusion criteria: not reported
	Unconscious, intubated patients with suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided (assumed head injury based on severity of injuries – all unconscious an intubated)
Index test(s) and	Index test
reference standard	X-ray of cervical spine (data not included further in review as not relevant as initial imaging in severely injured)
	CT of cervical spine
	Single-slice helical CT performed from base of skull to first thoracic vertebra. Performed at 2 mm thickness and 1.5 mm pitch with sagittal and coronal reformations.
	Reference standard
	Final diagnosis, including all imaging performed (MRI in some) and follow-up through hospital stay to identify missed injuries

Reference	Brohi 2005 ⁶				
	MRI was performed if CT scan or lateral radiograph suggested ligamentous injury or instability, if there were neurological signs of spinal cord injury before intubation or if there were contradictory findings between plain film and CT imaging. Consultant trauma radiologist (board-certified or equivalent) reported the images. If all imaging was normal, spine was cleared and spinal precautions removed. Patients assessed once they regained consciousness and followed through their hospital stay for any evidence of missed spinal injury.				
	Time between measurement of index test and reference standard: unclear duration between index tests and other imaging/final diagnosis being confirmed				
Outcome	Any cervical spir	ne injury			
	and Unstable cervical spine injury – defined using White and Punjabi system and three-column model of Denis				
2×2 table	Any cervical spine injury – X-ray as index test				
		Reference standard +	Reference standard -	Total	Results for 'all laterals' reported including all films
	Index test +	44	21	65	rather than just those deemed 'adequate'.
	Index test -	17	339	356	
	Total	61	360	421	
	Any cervical sp	oine injury – CT as inde	ex test		

Reference	Brohi 2005 ⁶				
		Reference standard +	Reference standard -	Total	
	Index test +	51	4	55	
	Index test -	1	325	326	
	Total	52	329	381	
	Unstable cervic	al spine injury – X-ray	as index test		
		Reference standard +	Reference standard -	Total	Results only available for the subgroup that had 'adequate'
	Index test +	24	14	38	films, which was n=200.
	Index test -	8	154	162	
	Total	31	168	200	
	Unstable cervic	al spine injury – CT as	index test		
		Reference standard +	Reference standard	Total	
	Index test +	29	4	33	
	Index test -	0	348	348	
	Total	29	352	381	

Reference	Brohi 2005 ⁶
Statistical measures	Any cervical spine injury – X-ray as index test: reported in paper (apart from PPV which was calculated using excel)
	Sensitivity: 72.1%
	Specificity: 94.2%
	PPV: 68.0%
	NPV: 95.2%
	Any cervical spine injury – CT as index test: reported in paper (apart from PPV which was calculated using excel)
	Sensitivity: 98.1%
	Specificity: 98.8%
	PPV: 93.0%
	NPV: 99.7%
	Unstable cervical spine injury – X-ray as index test: reported in paper (apart from PPV which was calculated using excel)
	Sensitivity: 75.0%
	Specificity: 91.7%
	PPV: 63.0%

Reference	Brohi 2005 ⁶
	NPV: 95.1%
	Unstable cervical spine injury – CT as index test: reported in paper (apart from PPV which was calculated using excel)
	Sensitivity: 100.0%
	Specificity: 99.0%
	PPV: 88.0%
	NPV: 100.0%
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if index test and reference standard interpreted without knowledge of the other, unclear time interval between index test and reference standard and likely that reference standard components differed between patients – applies to both index tests
	Indirectness: very serious – all included were at unconscious and intubated representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury, and unclear if reference standard included a 2 week follow-up period
Comments	None

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Reference	Bush 2016 ⁷
Study type	Prospective study
Study methodology	Data source: Conducted at an American College of Surgeons-verified Level 1 trauma centre.
	Recruitment: conducted between March 2014 and March 2015.
Number of patients	n = 1668 (only provides useable results for the n=632 that were intoxicated by alcohol and/or drugs)
	(n=1696 underwent cervical spine CT with n=28 of these subsequently excluded from the overall population for unclear reasons; useable results were only provided for n=632 in intoxicated subgroup)
Patient characteristics	Age, mean (SD): 45 (17), 39 (14) and 43 (17) in subgroups intoxicated by alcohol only, drugs only or alcohol and drugs
	Gender (male to female ratio): 71.7% male and 28.3% female
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury – reported for whole population as breakdown not given specifically for the intoxicated group
	Motor vehicle crash, 28.2%
	Ground-level fall, 20.2%

Reference	Bush 2016 ⁷
	Other mechanism, 15.8%
	• Fall from height, 14.5%
	Motorcycle crash, 6.9%
	Automobile vs. pedestrian, 5.5%
	• Assault, 5.5%
	Bicycle vs. automobile, 3.4%
	GCS score:
	• GCS 15, 52.8%
	• GCS 9-14, 33.7%
	• GCS <9, 13.4%
	Injury Severity Score, mean (range): 8 (8), 12 (10) and 9 (10) in subgroups intoxicated by alcohol only, drugs only or alcohol and drugs
	,
	Additional imaging, 85 (5%) – reported for whole population as breakdown not given specifically for the
	intoxicated group
	Setting: secondary care – trauma centre

Reference	Bush 2016 ⁷
	Country: USA
	Inclusion criteria: adults (≥18 years) with blunt trauma; and underwent evaluation of cervical spine with cervical CT scan – only provides results for the group of participants that were intoxicated (defined as a blood alcohol level greater than 80 mg/dL and/or apositive urine drug screen that was not attributable to field or emergency department medication administration).
	Exclusion criteria: patients presenting in a delayed manner after the index trauma, transfer patients without available CT images from the referring facility who did not undergo repeated CT scans at our facility, and patients with known recent cervical spinefractures or surgery
	Adults with suspected cervical spine injury undergoing CT (results only provided for those that were intoxicated)
Target condition(s)	Suspected cervical spine injury in those that were intoxicated – unclear if most/all had head injury
Index test(s) and	Index test
reference standard	CT of cervical spine
	CT scans performed using one of two emergency department imagers. CT protocol included continuous image acquisition from the skull base through the T1 vertebral body using 2-mm slice thickness. Axial images as well as coronal and sagittal reconstructions wereimmediately reviewed and interpreted one of eight certified radiologists, two of whom were neuroradiologists.

Reference	Bush 2016 ⁷				
	Reference standard				
	Cervical spine injury diagnosis at discharge/follow-up – includes composite end-point, which included MRI findings, operative findings and clinical status at discharge. Likely to differ between patients as not all would have had MRI and same imaging during stay. Also mentions identification of missed clinically significant injuries from outpatient notes following discharge. Unclear how long this followup was for and whether the same in all patients.				
	All patients followed up through completion of hospital stay and re-evaluated at time of discharge, which include recording of anyinterval diagnosis of cervical spine pathology as well as other details.				
	Time between measurement of index test and reference standard: unclear, varies between patients as components of additional imaging/follow-up prior to discharge and following discharge likely differ				
Outcome	Cervical spine injury – any bony, ligamentous or spinal cord injury.				
	Unstable cervical spine injury – any injury that required or benefitted from spine immobilisation or alternatively was at risk of any adverse effect because of the removal of spine precautions. Any injury defined as unstable or potentially unstable injury that required surgical stabilisation or prolonged immobilisation.				y injury defined as unstable or
2×2 table	All cervical spine injuries – CT vs. reference standard				
		Reference standard +	Reference standard -	Total	Note: results only given for the subgroup that were
	Index test +	56	1	57	intoxicated.
	Index test -	5	570	575	

Reference	Bush 2016 ⁷				
	Total	61	571	632	Raw data calculated from diagnostic accuracy measures provided in paper.
	Unstable cervical spine injuries – CT vs. reference standard				
		Reference standard +	Reference standard	Total	Note: results only given for the subgroup that were
	Index test +	13	0	13	intoxicated.
	Index test -	1	617	618	Raw data calculated from diagnostic accuracy
	Total	14	617	631	measures provided in paper – does not quite total 632 which is the number said to be analysed.
Statistical measures	All cervical spine injuries - CT vs. references standard (intoxicated patients) - reported in the paper				
	Sensitivity: 92.9%				
	Specificity: 99.8% PPV: 98.5%				
	NPV: 99.2%				
	Unstable cervical spine injuries – CT vs. reference standard (intoxicated patients) – reported in the paper Sensitivity: 91.6%			ients) – reported in the paper	
	Specificity: 100%				

Reference	Bush 2016 ⁷
	PPV: 100%
	NPV: 99.8%
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled and reasons for exclusion not reported, some concerns about whether the reference standard was interpreted without knowledge of the index test, unclear follow-up time period for assessing reference standard outcome and references standard components likely differed between participants
	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, also limited to very specific population of those that were intoxicated and unclear time-point for reference standard and whether it matches protocol
Comments	2x2 data not reported so calculated from the diagnostic accuracy measures reported (sensitivity etc.)

Reference	Cohn 1991 ⁹
Study type	Prospective study
Study methodology	Data source: data from a single university medical centre
	Recruitment: between July 1989 and August 1989, consecutive patients admitted with blunt traumatic injury were evaluated for cervical spine injury
Number of patients	n = 60

Reference	Cohn 1991 ⁹
	(n=60 said to be evaluated and n=60 analysed)
Patient characteristics	Age, mean (SD): not reported
	Gender (male to female ratio): not reported
	Ethnicity: not reported
	Head injury: 50% had head CT as part of diagnostic tests, unclear if remaining patients had some form of head injury as part of the injury mechanism
	Altered sensorium:
	 GCS <15, 48.3% Coma (GCS <8), 15.0% Intoxicated, 35.0%
	Abnormal neck exam, 11.7%
	Cord injury, 1.7%

Reference	Cohn 1991 ⁹
	Shock (BP <80), 3.3%
	Setting: secondary care – admitted to centre with blunt traumatic injury
	Country: USA
	Inclusion criteria: blunt traumatic injury evaluated for presence of cervical spine injury
	Exclusion criteria: not reported
	People with blunt traumatic injury admitted to centre, with suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – 50% had head CT as part of diagnostic tests, unclear if remaining patients had some form of head injury as part of the injury mechanism
Index test(s) and	Index test
reference standard	
	X-ray of cervical spine
	Performed in the trauma room after sensorium and neck assessment/examination. Initially evaluated by radiology resident and trauma chief resident for presence of pathology. Completion cervical spine studies included more sophisticated studies (wide supra) when needed to exclude injury. All patients managed in cervical collars until cleared of spine injury.

Reference	Cohn 1991 ⁹				
	Reference standard Reference standard unclear, possibly a final diagnosis based on any further imaging performed (including flexion/extension views, cervical CT scans or tomograms where indicated)				
	Time between measurement of index test and reference standard: unclear time interval between index tests and subsequent tests/final confirmed diagnosis			terval between index tests and	
Outcome	Acute cervical spine injuries – poorly defined				
2×2 table	Acute cervical spine injury – X-ray as index test				
		Reference standard +	Reference standard	Total	Data reported insufficiently meaning specificity could not
	Index test +	4	NR	NR	be calculated and it was not reported in the paper.
	Index test -	3	NR	NR	
	Total	7	53	60	
Statistical measures	Acute cervical spine injury – X-ray as index test: calculated using excel sheet				
	Sensitivity: 57.0%				
	Specificity: NR				
	PPV: NR				

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Reference	Cohn 1991 ⁹
	NPV: NR
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if reference standard interpreted without knowledge of index test, unclear interval between index test and reference standard and unclear if reference standard contained the same components for all patients
	Indirectness: very serious – head CT performed for 50% but unclear if remaining also had head injury as part of injury mechanism, and unclear if reference standard included a 2 week follow-up period
Comments	None
Reference	Dan Lantsman 2020 ¹⁰
Study type	Retrospective cohort study
Study methodology	Data source: review of records of those admitted to emergency room. Picture computerised archive system (PACS) used to extract whole spine CT examinations.
	Recruitment: review of medical records and whole spine CT scans performed between 2017 and 2018
Number of patients	n = 147 (n=129 analysed, as n=9 were excluded due to poor X-ray quality and n=9 were excluded due to missing radiographs)
Patient characteristics	Characteristics are provided for the n=147 included in the study initially, not the n=129 analysed as part of the diagnostic accuracy assessment

Reference	Dan Lantsman 2020 ¹⁰
	Age, median (IQR): 83 (77-88) years
	Gender (male to female ratio): 43.5% male and 56.5% female
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported
	Setting: secondary care – emergency room following low-energy trauma
	Country: Israel
	Inclusion criteria: radiographic diagnosis of diffuse idiopathic skeletal hyperostosis (DISH); following low-energy trauma (traumatic or accidental injury with a maximal Injury Severity Score of 9/75 not requiring invasive procedures); and underwent a whole spine CT with radiographs for at least the thoracic and lumbar spine.

Reference	Dan Lantsman 2020 ¹⁰
	Exclusion criteria: spinal CT examination did not confirm DISH; those with known malignant spinal involvement or ankylosing spondylitis; those with missing spinal radiographs (except for if they were missing only cervical spine radiographs); and with poor radiography quality
	Those with suspected spine injury with DISH following low-energy trauma (separate results for cervical spine injuries)
Target condition(s)	Suspected spine injury in those with DISH following low-energy trauma – spine in general but provides results separately for cervical spine region – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	X-ray (radiographs) – performed in anterior-posterior and lateral projections.
	Reference standard
	Whole spine CT scan – performed in axial plane on 64-Slice machine. Images reconstructed in bone and soft tissue algorithms and reformatted in sagittal and coronal planes.
	Spinal radiographs and whole spine CT were evaluated separately for presence of spinal fractures by a single reader (third year radiology resident) with at least 1-month interval between readings. Reader was blinded to patient clinical data (apart from age and gender) and the radiographic report. Fractures and locations recorded for radiographs and CT scans and classified as acute or chronic fractures. Acute fractures were those not present in previous studies and consisting of a radiographically depicted cortical disruption or impaction of the trabeculae and paravertebral soft tissue infiltration. Chronic fractures were those detected and unchanged from previous radiological studies and consisting of any degree of remodelling and smoothing of cortical edges with no anterior vertebral body buckling. Second reading by a senior, experienced musculoskeletal radiology was performed on 10% of radiographs and CT scans.

Reference	Dan Lantsman 2020 ¹⁰
	Time between measurement of index test and reference standard: describes at least 1 month interval between readings – unclear if this is between multiple readings of the same imaging (e.g. X-ray and a subsequent X-ray), or between X-ray and reference standard (CT).
Outcome	Acute fracture - those not present in previous studies and consisting of a radiographically depicted cortical disruption or impaction of the trabeculae and paravertebral soft tissue infiltration.
2×2 table	2x2 tables could not be reported as raw data not reported for cervical spine injuries specifically. Attempted to calculate raw data from diagnostic accuracy measures reported in the paper but not possible given 0 values for sensitivity and PPV.
Statistical measures	X-ray vs. reference standard (whole spine CT) – reported in paper
	Sensitivity: 0%
	Specificity: 100%
	PPV: 0%
	NPV: 96.9%
Source of funding	Reported to be no funding for the study.
Limitations	Risk of bias: serious – unclear if consecutive sample enrolled, not all were analysed as some were missing radiographs or had poor quality radiographs and unclear duration between index test and reference standard
	Indirectness: very serious - head injury not mentioned and unclear if all or most had head injury. Very specific population of those with DISH, a condition making injuries more likely following lower impact trauma. Also, injury reported was specifically fracture not any type of injury.

Reference	Dan Lantsman 2020 ¹⁰
Comments	2x2 tables could not be reported as raw data not reported for cervical spine injuries specifically. Attempted to calculate raw data from diagnostic accuracy measures reported in the paper but not possible given 0 values for sensitivity and PPV.

Reference	Duane 2008 ¹⁴
Study type	Prospective
Study methodology	Data source: evaluation of those admitted to a single level 1 trauma centre
	Recruitment: prospective evaluation between February 2004 and September 2006 of all blunt trauma team alert patients >16 years admitted to single level 1 trauma centre
Number of patients	n = 1004
	(N=4608 patients >16 years with blunt trauma identified, with n=1004 subsequently included and analysed as they had both lateral cervical spine films and CT of the cervical spine)
Patient characteristics	Note: n=84 in fracture group and n=920 in no fracture group
	Age, mean (SD): 41.3 (19.0) years in fracture group and 41.3 (21.0) years in no fracture group
	Gender (male to female ratio): 61.9% male in fracture group and 64.3% male in no fracture group
	Ethnicity: 62.0% white in fracture group and 59.0% white in no fracture group
	Lie edicium une detaile nemented, un de cuif ell en mont suffered concernitent inium to the based
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head

Reference	Duane 2008 ¹⁴
	Mechanism of injury: 72.6% in fracture group and 77.4% in no fracture group had injury due to motor vehicle collision.
	GCS at initial trauma, mean (SD): 12.4 (4.6) vs. 14.0 (3.0) in fracture and no fracture groups
	Injury Severity Score, mean (SD): 20.8 (14.2) vs. 11.6 (10.6) in fracture and no fracture groups
	Setting: secondary care – trauma centre
	Country: USA
	Inclusion criteria: blunt trauma team alert patients; >16 years; admitted to single level 1 trauma centre; and underwent both lateral cervical spine film and cervical spine CT
	Exclusion criteria: not reported
	People admitted to trauma centre with blunt injury and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided

Reference	Duane 2008 ¹⁴	Duane 2008 ¹⁴			
Index test(s) and	Index test	Index test			
reference standard	X-ray of cervical	spine			
	Reference standa	ard			
	CT of cervical sp	ine – gold standard for t	fractures		
	CT of cervical spine used as gold standard for diagnosis of cervical spine fracture. At time of study, standard approach for all trauma activation patients was thorough clinical examination followed by cervical spine film. All patients, regardless of level of consciousness or intoxication then had CT of cervical spine. Further radiographic studies then performed based on results of initial CT scan. Time between measurement of index test and reference standard: unclear time interval between index test and reference standard			ved by cervical spine film. All	
				erval between index test and	
Outcome	Cervical spine fra	octure			
2×2 table	Cervical spine f	racture – X-ray as inde	ex test		
		Reference standard +	Reference standard	Total	
	Index test +	16	7	23	
	Index test -	68	913	981	
	Total	84	920	1004	

Reference	Duane 2008 ¹⁴
Statistical measures	Cervical spine fracture – X-ray as index test: reported in paper
	Sensitivity: 19.0%
	Specificity: 99.2%
	PPV: 69.6%
	NPV: 93.1%
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if reference standard interpreted without knowledge of index test and unclear time interval between index test
	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, and outcome limited to fractures rather than any cervical spine injury
Comments	None
Reference	Duane 2010 ¹³
Study type	Retrospective
Study methodology	Data source: retrospective review of established trauma registry and chart review

Recruitment: retrospectively matching inclusion criteria between January 2000 and December 2008 from single level 1 trauma centre

Number of patients

n = 49

Reference	Duane 2010 ¹³
	(n=271 matching inclusion criteria that underwent flexion-extension films, with n=49 eventually included as they also had MRI of the cervical spine)
Patient characteristics	Age, mean (SD): 37.9 (17.7) years
	Gender (male to female ratio): 69.4% males
	Ethnicity: 49.0% white
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury:
	 Motor vehicle collision, 69.0% Falls, 16.0% Other, 15.0%
	GCS, mean (SD): 13.8 (3.5)
	Injury Severity Score, mean (SD): 15.6 (10.2)

Reference	Duane 2010 ¹³
	Duration of stay, mean (SD): 8.0 (11.2) days
	Initial lactate: 2.2 (1.7) mmol/L
	Setting: secondary care – data from trauma registry
	Country: USA
	Inclusion criteria: adults (≥18 years) sustaining blunt trauma; and had flexion-extension plain films and subsequent MRI evaluation of cervical spine.
	Exclusion criteria: not reported
	Adults with blunt trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	X-ray of cervical spine
	Reference standard

Reference	Duane 2010 ¹³				
	MRI – gold standard for ligamentous injuries				
	Flexion-extension considered complete if it visualised from the first cervical spine through to the bottom of the first thoracic spine and >30° excursion on both flexion and extension. All performed actively without fluoroscopy. Flexion-extension compared with MRI as the gold standard for diagnosis of ligamentous injury. MRI performed using Siemens Avanto 1.5T with scans performed without contrast. Multiple sequences included: T1 turbo spin echo (TSE) sagittal, T2 TSE sagittal, T2 short tau inversion recovery sagittal, T2* multiple echo data image combination or gradient echo axial, T2 TSE axial and fast low-angle shot two-dimensional sagittal gradient echo.				
Outcome	Ligamentous injury of the cervical spine				
2×2 table	Ligamentous cervical spine injury – X-ray as index test				
		Reference standard +	Reference standard	Total	Of the 8 ligamentous injuries missed by X-ray, five were
	Index test +	0	1	1	significant (n=2 associated fractures requiring prolonged
	Index test -	8	40	48	collar and n=3 operative intervention)
	Total	8	41	49	
Statistical measures	Ligamentous cervical spine injury – X-ray as index test: reported in paper				
	Sensitivity: 0.0%				

Reference	Duane 2010 ¹³
	Specificity: 98.0%
	PPV: 0.0%
	NPV: 83.0%
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if reference standard interpreted without knowledge of index test and unclear time interval between index test Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, and outcome limited to ligamentous injuries rather than any cervical spine injury
Comments	None
Reference	Duane 2016 ¹⁶
Study type	Retrospective study

Reference	Duane 2016 ¹⁶
Study type	Retrospective study
Study methodology	Data source: retrospective review of trauma registry
	Recruitment: those presenting as blunt trauma team alert patients between January 2008 and May 2014 at American College of Surgeons verified level I trauma centre (Virginia Commonwealth University Medical Center)
Number of patients	n = 9227
	(patient flow not well described)

Reference	Duane 2016 ¹⁶
Patient characteristics	Age, mean (SD): 39.4 (17.5) years
	Gender (male to female ratio): 64.4% male and 35.6% female
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury:
	Motor vehicle collision, 59%
	• Falls, 12%
	Motorcycle collision, 10%
	Pedestrian struck, 10%
	Other/unknown, 9%
	GCS, mean (SD; median): 14.3 (2.4; 15)
	Setting: secondary care – trauma centre

Reference	Duane 2016 ¹⁶
	Country: USA
	Inclusion criteria: adults (≥18 years) with blunt trauma; and underwent screening CT to diagnose or rule-out cervical spine injury.
	Exclusion criteria: none reported
	Adults following trauma undergoing assessment of cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	CT scan
	Using 64-multidetector CT between 2008 and 2011 and a 128-multidetector CT between 2011 and 2014. 2-mm thick axial cuts performed at 2-mm increments with multiplanar reformatted images.
	Reference standard
	Later found to have cervical spine injury – poorly described. Likely involves any other findings during follow-up but duration of follow-up available unclear. CT considered the standard for presence or absence of fracture and

Reference	Duane 2016 ¹⁶
	magnetic resonance imaging for ligament injury. MRI performed in some as indicated but not all patients. For MIR, 3-mm sagittal and axial cuts with 0.3 mm standard of error.
	Haemodynamically stable patients were evaluated by non-contrast CT of cervical spine. Those that were unstable had their cervical collar maintained and spine examination and CT deferred until patient had stabilised. Post-CT management was as follows: normal CT result and conscious had cervical spine re-evaluated for midline tenderness. If no tenderness then collar was removed. If tenderness present then collar was maintained for 2 weeks and examination repeated. If significant tenderness persisted then an MRI was obtained. If normal, collar was removed and physical therapy started. If there was an abnormal result, management was based on specialist consultation. Patients with a normal CT who could not participate in their own evaluation (GCS <15) underwent MRI and further management as above. For those with an abnormal CT result, liberal use of MRI and spine consultations was used.
	For all imaging, only final attending radiologist reads of scans were considered. Negative CT or MRI results were defined as high quality images without motion artifacts with no fracture and/or ligament injury identified. Positive CT or MRI defined as one where a fracture or ligament injury was identified or could not be excluded. Specific findings suggestive of ligamentous injury included abnormal vertebral alignment, increased space between ligamentous columns or other contiguous structures and prevertebral haematoma or oedema.
	Time between measurement of index test and reference standard: unclear, likely that reference standard components and timeinterval differs between patients.
Outcome	Fracture and/or ligamentous injury
	Ligamentous injury - specific findings suggestive of ligamentous injury included abnormal vertebral alignment, increased space between ligamentous columns or other contiguous structures and prevertebral haematoma or oedema.

Reference Duane 2016¹⁶

2×2 table

CT vs. reference standard (later diagnosis of injury) – fracture and/or ligamentous injury

	Reference standard +	Reference standard	Total	Raw data calculated from diagnostic accuracy
Index test +	561	6	567	measures provided in paper.
Index test -	0	8660	8660	Notes much an of two
Total	561	8666	9227	Note: number of true positives does not match that reported in paper (n=553) but insufficient data provided to calculate diagnostic accuracy results from raw data.

CT vs. reference standard (later diagnosis of injury) – ligamentous injury (with or without an associated fracture)

	Reference standard +	Reference standard -	Total	Raw data calculated from diagnostic accuracy
Index test +	29	9	38	measures provided in paper.
Index test -	28	9160	9188	
Total	57	9169	9226	Note: total number calculated from diagnostic accuracy measures reported does not quite match 9227 reported in the paper.

Reference	Duane 2016 ¹⁶	
		Note: all ligamentous injuries were also associated with a fracture so no individuals with a ligamentous injury were missed by CT, as they had a fracture that was picked up by CT.
Statistical measures	CT vs. reference standard (later diagnosis of injury) – fra paper	cture and/or ligamentous injury – reported in
	Sensitivity: 100.0%	
	Specificity: 99.93%	
	PPV: 98.93%	
	NPV: 100%	
	CT vs. reference standard (later diagnosis of injury) – lig fracture) – reported in paper	amentous injury (with or without an associated
	Sensitivity: 50.88%	
	Specificity: 99.90%	
	PPV: 76.31%	
	NPV: 99.69%	
	Note: all of these ligamentous injuries were also associated w	with a fracture, which were all picked up by CT
Source of funding	Not reported	

Reference	Duane 2016 ¹⁶
Limitations	Risk of bias: very serious –unclear if consecutive sample enrolled, reference standard poorly described and likely to have been interpreted with knowledge of index test, unclear time interval between index test and reference standard and likely that reference standard different between patients
	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, and unclear if reference standard matches protocol as poorly defined
Comments	2x2 data not reported so calculated from the diagnostic accuracy measures reported (sensitivity etc.)

Reference	Fisher 2013 ¹⁷
Study type	Retrospective study
Study methodology	Data source: subjects identified using trauma registry at University Medical Center, a level I academic trauma centre in Lubbock, Texas.
	Recruitment: included those matching inclusion criteria and admitted between 1 st January 2005 and 30 th March 2012.
Number of patients	n = 277(n=1354 blunt trauma patients admitted and n=277 subsequently included as they had both a CT and MRI performed of cervical spine)
Patient characteristics	Age, mean (range): 35.2 (0-93) years
	 Children <18 years, 14%
	Gender (male to female ratio): 70% males and 30% females

Reference	Fisher 2013 ¹⁷
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury:
	Motor vehicle collisions, 70%
	• Falls, 9%
	Assault, 8%
	Pedestrian/bike accidents, 3%
	• Other, 10%
	GCS score, mean (range): 6 (3-14)
	Injury Severity Score, mean (range): 22 (0-75)
	Duration of stay, mean (range): 15.3 (1-66) days

Reference	Fisher 2013 ¹⁷
	Setting: secondary care – trauma centre
	Country: USA
	Inclusion criteria: blunt trauma patients with GCS <15; and underwent both a CT scan and MRI of the cervical spine.
	Exclusion criteria: none reported
	People with blunt trauma and suspected cervical spine injury (majority adults at least 18 years old, 86%)
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided (assumed to have head injury based on severity of injuries – all obtunded)
Index test(s) and	Index test
reference standard	CT – performed with 64-slice or 16-slice scanner.
	OR
	MRI – performed using General Electric HDX scanner.

Reference	Fisher 2013 ¹⁷
	Reference standard
	Diagnosis of clinically significant cervical spine injury by any modality – this was used in the study as the denominator as they noted a lack of an external gold standard. The use of this as the denominator does not allow specificity to be calculated.
	Time between measurement of index test and reference standard: MRI was obtained an average of 3.3 days (range 0-39) days after admission. Unclear whether this gives an indication of time between CT and MRI as unclear if CT performed immediately on admission for all.
Outcome	Clinically significant cervical spine injury
	CT and MRI scans considered clinically significant if detecting one of the following: ligamentous injury in two adjacent spinal columns, subluxations, cord injury, nerve root injury, disc herniations, and fractures except the following types as specified by NEXUS: spinous process fracture without involvement of the lamina, transverse process fracture without involvement of the facet joint, osteophytefracture not including corner or teardrop fracture, isolated avulsion without associated ligamentous injury, simple wedge-compression fracture without loss of greater than or equal to 25% of vertebral body height, endplate fracture, type 1 odontoid fracture, and injury to the trabecular bone.
2×2 table	Raw data provided but difficult to understand – attempted calculations of true positives and false positives do not match sensitivity values reported in the paper for CT and MRI and therefore values reported in paper used and 2x2 tables not completed.
Statistical measures	CT vs. reference standard (CT + MRI) for clinically significant cervical spine injuries
	Sensitivity: 83%
	Specificity: NR
	PPV: NR

Reference	Fisher 2013 ¹⁷
	NPV: NR
	MRI vs. reference standard (CT + MRI) for clinically significant cervical spine injuries
	Sensitivity: 89%
	Specificity: NR
	PPV: NR
	NPV: NR
	Specificity could not be calculated as the combine CT + MRI was used as the reference standard, meaning it is not possible for there to be any false positives. N=70 were positive on both modalities (of these, n=1 clinically insignificant on CT but significant on MRI), n=11 were positive on CT but negative on MRI, n=150 were negative on both CT and MRI and n=12 were negative on CT but positive for clinically significant injury on MRI. An additional n=34 were negative on CT and positive on MRI, but with clinically insignificant injuries that did not form part of the calculation of sensitivity for clinically significant injuries.
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unlikely that index test (for MRI) or reference standard (for CT) were interpreted without knowledge of reference standard/index test, and mean time interval between index test and reference standard possibly at least 3 days – applies to both index tests

Reference	Fisher 2013 ¹⁷
	Indirectness: serious – all included were obtunded representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury
Comments	Study notes the intention was not to compare the accuracy of CT and MRI as a solo modality but to assess the added value of MRI to more safely clear the cervical spine.
	Not possible to calculate specificity based on the reference standard used.

Reference	Friesen 2014 ¹⁸
Study type	Retrospective
Study methodology	Data source: retrospective search of radiology information system database containing radiological information from three major public emergency departments
	Recruitment: retrospective review of people matching inclusion criteria from database of three major public emergency departments between 12 th January 2010 and 22 nd June 2012 – another time-period mentioned but not relevant to review protocol
Number of patients	n = 206 undergoing both MRI and CT of cervical spine
	(n=783 identified as relevant to the study and n=206 subsequently included in analysis as they had both CT and MRI of cervical spine)
Patient characteristics	Note: characteristics only reported for whole group (n=783) and not the relevant group analysed (n=206)
	Age, mean (SD): 60 (25) years
	Gender (male to female ratio): 55.0% males

Reference	Friesen 2014 ¹⁸
	Ethnicity: not reported
	Head injury: 76% with head CT as well as cervical spine CT
	Mechanism of injury: not reported
	GCS, mean (SD): not reported
	Injury Severity Score, mean (range): not reported
	Setting: secondary care – database of emergency department data
	Country: Australia
	Inclusion criteria: aged ≥16 years; and CT and MRI performed for suspected blunt acute cervical spine trauma between 12 th January 2010 and 22 nd June 2012
	Exclusion criteria: not reported

Reference	Friesen 2014 ¹⁸
	Adults with blunt trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – 76% said to have had brain CT alongside cervical spine imaging, therefore not downgraded for indirectness
Index test(s) and	Index test
reference standard	CT of cervical spine
	MRI of cervical spine
	Reference standard
	MRI reported in paper to be reference standard, but not in line with this review protocol. Therefore, data available used to calculate sensitivity of CT and MRI using CT + MRI as combined reference standard. This means specificity could not be calculated.
	Images from MRI and CT examinations retrospectively reviewed by consultant radiologist and radiology registrar. Each case reviewed by both authors and consensus determination of no traumatic abnormality, stable traumatic abnormality or unstable traumatic abnormality noted. CT helical acquisition made from above C1 to below T2. Reconstructions performed by 2 mm contiguous slices in axial, coronal and sagittal planes. Six different CT scanners from three manufacturers of 16, 64 and 128 slice were used. MRI performed with sagittal short T1 inversion recovery, T1 and T2 weighted imaging, axial 3D T2 weighted imaging and gradient recalled echo Siemens Magnetom Symphony 1.5T.
	Time between measurement of index test and reference standard: time between CT and MRI unclear

Reference	Friesen 2014 ¹⁸				
Outcome	Any cervical spinal cord injury (stable and unstable)				
	Unstable injuries only also reported but not extracted as would require MRI to be used as reference stand results not given for CT in terms of classifying into stable/unstable injuries based on CT. Unstable injuries by Denis 3 column definition as well as any cases requiring urgent surgery (within 5 days of injury) or urg- immobilisation (e.g. halo-traction ring). Other specific unstable injuries also included: flexion teardrop frac bilateral locked facets, hangman's fracture, Jefferson fracture and Type 2 dens fracture.			on CT. Unstable injuries defined 5 days of injury) or urgent ed: flexion teardrop fracture,	
2×2 table	Any cervical spi	nal cord injury – CT a	s index test		
		Reference standard +	Reference standard	Total	Note false positives were not possible using CT + MRI as a reference standard meaning specificity could not be calculated.
	Index test +	115	NA	115	
	Index test -	24	67	91	
	Total	139	67	206	
	Any cervical spinal cord injury – MRI as index test				
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a reference standard meaning specificity could not be calculated.
	Index test +	98	NA	98	
	Index test -	41	67	108	
	Total	139	67	206	

Reference	Friesen 2014 ¹⁸
Statistical measures	Any cervical spinal cord injury – CT as index test: calculated using excel sheet
	Sensitivity: 83.0
	Specificity: NA
	PPV: NA
	NPV: NA
	Any cervical spinal cord injury – MRI as index test: calculated using excel sheet
	Sensitivity: 71.0
	Specificity: NA
	PPV: NA
	NPV: NA
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unlikely that index test (for MRI) or reference standard (for CT) were interpreted without knowledge of reference standard/index test, and mean time interval between index test and reference standard unclear – applies to both index tests
	Indirectness: none – considered to represent head injury population as >75% said to have had brain CT at same time
Comments	Not possible to calculate specificity based on the reference standard used.

Reference	Gale 2005 ¹⁹
Study type	Retrospective
Study methodology	Data source: retrospective review of patients evaluated by Trauma Service Activation at Hospital of Pennsylvania
	Recruitment: retrospective inclusion of those matching inclusion criteria from single hospital between December 2002 and July 2003
Number of patients	n = 400
	(n=848 with blunt trauma, of which n=716 had a CT of the cervical spine and n=640 had both CT and plain radiography/X-ray of cervical spine; population was further reduced to n=400 having plain radiography/X-ray and supplemental CT)
Patient characteristics	Note: characteristics below given for whole group only (n=1151) and not those analysed (n=400)
	Age, mean (SD): 44.00 (22.08) years
	Gender (male to female ratio): 64.0% males
	Ethnicity: not reported
	Head injury: 84.4% of n=848 blunt trauma patients (excluding those with penetrating injuries) had head CT
	Mechanism of injury:

Reference	Gale 2005 ¹⁹
	 Motor vehicle collision, 42.3% Fall, 30.5% Assault, 11.9% Pedestrian vs. auto, 6.6% Other, 8.6%
	GCS, mean (SD): not reported
	Injury Severity Score, mean (SD): 9.37 (10.06)
	Setting: secondary care – those arriving in hospital with blunt trauma
	Country: USA
	Inclusion criteria: blunt trauma patients evaluated by Trauma Service activation between December 2002 and July 2003 – those relevant to review also had X-ray and CT of cervical spine that were complete.
	Exclusion criteria: penetrating injuries
	People with blunt trauma and suspected cervical spine injury

Reference	Gale 2005 ¹⁹
Target condition(s)	Suspected cervical spine injury – all underwent head CT so considered a direct population
Index test(s) and	Index test
reference standard	X-ray of cervical spine
	Reference standard
	CT of cervical spine – gold standard for fractures
	Intranet-based electronic medical records reviewed to ascertain which radiographic studies were obtained during their trauma evaluation. Reports of all studies were reviewed and anatomic adequacy of each and its results were recorded in a spreadsheet. Pain cervical spine radiography considered anatomically inadequate if evaluating radiologist dictation included any of the following: study limited to level <t1, a="" anatomic="" as="" attain="" because="" cervical="" cervicothoracic="" completion="" ct="" defined="" entire="" finding="" if="" inadequate="" it="" junction="" limited="" non-visualisation="" not="" observe="" obtained="" of="" on="" or="" performed="" plain="" radiography.<="" region="" specific="" spine="" spine.="" statement="" study="" supplemental="" suspicious="" t1.="" th="" the="" to="" visualised="" was=""></t1,>
	Time between measurement of index test and reference standard: unclear time interval between index test and reference standard
Outcome	Cervical spine fracture
	Plain cervical spine radiography considered positive if a specific feature identified or an area on plain films was interpreted as suspicious and warranting further imaging. Negative if no fracture identified and no further imaging recommended. CT considered positive only if a specific fracture was identified.
2×2 table	Cervical spine fracture – X-ray as index test

Reference	Gale 2005 ¹⁹					
		Reference standard +	Reference standard	Total		
	Index test +	6	3	9		
	Index test -	13	378	391		
	Total	19	381	400		
Statistical measures	Cervical spine fi	Cervical spine fracture – X-ray as index test: reported in paper				
	Sensitivity: 31.6%					
	Specificity: 99.2%					
	PPV: 66.7%					
	NPV: 96.7%					
Source of funding	Not reported	Not reported				
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if reference standard interpreted without knowledge of index test and unclear time interval between index test					
	Indirectness: serious – outcome limited to fractures rather than any cervical spine injury					
Comments	None					

Reference	Gharekhanloo 2021 ²⁰		
Study type	Prospective		
Study methodology	Data source: prospective study of trauma patients at an emergency department.		
	Recruitment: not reported.		
Number of patients	n = 220		
	(n=210 had normal CT scans, n=10 had cervical spine injury on CT scans).		
Patient characteristics	Age, mean (SD): 38.25 (5.13) years (35% between 26 and 35 years)		
	Gender (male to female ratio): 157/63		
	Ethnicity: not reported		
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head.		
	Mechanism of injury: car accidents (64%) and falls from height (17.7%).		
	GCS at initial trauma, mean (SD): not reported		
	Injury Severity Score, median: 5		

Reference	Gharekhanloo 2021 ²⁰
	Setting: Emergency department of a Hospital
	Country: Iran
	Inclusion criteria: low-risk status based on international NEXUS criteria.
	Exclusion criteria: penetrating trauma.
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	Plain radiography
	Reference standard
	Cervical CT scan
	Plain radiographs were obtained in anteroposterior, lateral and odontoid views. Cervical CT was performed using a 16-slice multidetector CT scanner in a supine position. Images started with lateral scout images from the foramen magnum to the junction of the C7-T1 vertebral Junction. The standard scan protocols included the

Reference	Gharekhanloo 2021 ²⁰				
	voltage of 130kV, collimation of 1mm, pitch of 0.66, and tube current-time product of 200mAs. Coronal and sagittal reformation images were reconstructed using 1.5-mm intervals from an axial source on a standard workstation.				
	Time between measurement of index test and reference standard: unclear time interval between index test and reference standard.				
Outcome	Cervical spine in	jury was defined as sub	luxation/dislocation or a	acute fracture or bo	oth.
	Interpretation of plain radiographs and cervical CT images preformed by 2 experienced, board-certified radiologists who were blinded to the results. A clinically significant injury was determined based on the neurosurgical recommendation of one or more interventions, operation and rigid cervical collar or halo application.				
2×2 table	Cervical spine i	njury – Plain radiogra	phy		
		Reference standard +	Reference standard -	Total	
	Index test +	4	6	10	
	Index test -	6	204	210	
	Total	10	210	220	
Statistical measures	Cervical spine injury				
	Sensitivity: 40%	Sensitivity: 40%			
	Specificity: 100%				

Reference	Gharekhanloo 2021 ²⁰
	PPV: 60%
	NPV: infinity
Source of funding	None
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear time interval between index test and reference standard.
	Indirectness: unclear if head injury patients.
Comments	Only 10 patients had cervical spine injury on the reference standard.

²¹ 22

Reference
 Goodnight 2008²¹

 Study type
 Retrospective

 Study methodology
 Data source: retrospective review of trauma registry (general database of all trauma admission) of a single American College of Surgeons verified Level 1 trauma centre

 Recruitment: retrospective chart review from a single trauma centre of admissions between 2003 and 2004

 Number of patients
 n = 379

 (n=1809 with trauma had CT of cervical spine, with n=379 subsequently included as they also had flexion-extension radiography performed once fracture had been ruled out on CT)

Reference	Goodnight 2008 ²¹
Patient characteristics	Age, mean (SD): 39 (19) years
	Gender (male to female ratio): 63% males
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: 53.0% motor vehicle crash
	GCS at initial trauma, mean (SD): not reported
	Injury Severity Score, median: 5
	Setting: secondary care – trauma centre database review
	Country: USA

Reference	Goodnight 2008 ²¹
	Inclusion criteria: blunt mechanism of injury; and received both CT of cervical spine and follow-up flexion/extension radiographs for continued cervical spine pain
	Exclusion criteria: neurologic deficits consistent with cervical cord injury; obtunded patients; penetrating mechanism of injury; and age <18 years
	Adults with blunt trauma and suspected cervical spine injury – continued cervical spine pain specifically
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	Flexion-extension X-ray of cervical spine
	Helical CT of cervical spine
	Reference standard All available evidence, including MRI in some patients. Unclear follow-up for those that did not have MRI. For CT, helical CT technique used with 1.5 mm collimation helical scanning at pitch of 1.5 from T1 to occiput performed in two acquisitions. Axial images reconstructed with bone algorithm at 1.5 mm intervals with sagittal and coronal reconstructions. MRI considered gold standard for ligamentous injuries. Obtained as confirmatory study in each patient with negative CT and positive flexion/extension radiograph. Radiologists routinely assessed CT scans for ligamentous injury.

Reference	Goodnight 2008 ²¹				
	Time between measurement of index test and reference standard: unclear time interval between index test and reference standard, unclear follow-up for those that did not have MRI.				
Outcome	Ligamentous injury of the cervical spine				
	Suspicion of ligamentous injury on CT based on interpretation of board-certified radiologists. Findings raising suspicion included paravertebral soft tissue swelling, widening or subluxation of facet joints, focal kyphosis with splaying of spinolaminar distances and abnormal widening of articulations at cranio-cervical junction. Report data obtained from radiology department database. CT findings classified into negative of cervical spine injury, suspicious for ligamentous injury and technically inadequate based on original reports.				
2×2 table	Ligamentous ce	Ligamentous cervical spine injury – Flexion-extension X-ray as index test			
		Reference standard +	Reference standard	Total	
	Index test +	6	10	16	
	Index test -	0	363	363	
	Total	6	373	379	
	Ligamentous cervical spine injury – Helical CT as index test				
		Reference standard +	Reference standard	Total	
	Index test +	6	13	19	
	Index test -	0	360	360	

Reference	Goodnight 2008	21			
	Total	6	373	379	
Statistical measures	Ligamentous ce	rvical spine injury – F	lexion-extension X-ray	y as index test: re	eported in paper
	Sensitivity: 100.0	%			
	Specificity: 97.3%)			
	PPV: 37.5%				
	NPV: 100.0%	NPV: 100.0%			
	Ligamentous cervical spine injury – Helical CT as index test: reported in paper				
	Sensitivity: 100.0%				
	Specificity: 96.5%)			
	PPV: 31.6%				
	NPV: 100.0%				
Source of funding	Not reported				
Limitations	without knowledg	e of index test, unclear		ndex test and refe	rence standard was interpreted rence standard and likely that a tests

Reference	Goodnight 2008 ²¹
	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, population where those with confirmed fractures were excluded meaning may differ from population presenting without any prior imaging, outcome limited to ligamentous injuries and unclear if reference standard included a 2 week follow-up period
Comments	Population had already been ruled out for cervical fracture so may represent different population to those initially presenting with no imaging.

Reference	Griffen 2003 ²²
Study type	Retrospective
Study methodology	Data source: TRACS database from single level 1 trauma centre
	Recruitment: retrospective review of database from single level 1 trauma centre between November 2000 and October 2001
Number of patients	n = 1199(n=3018 blunt trauma patients with risk of cervical spine injury identified, with n=1199 subsequently included as they had both plain radiography/X-ray and CT of cervical spine)
Patient characteristics	Average age: 39.4 years
	Gender (male to female ratio): 65.0% males Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head

Reference	Griffen 2003 ²²
	Mechanism of injury: not reported
	Average GCS: 13
	Average Injury Severity Score: 8.4
	Setting: secondary care – review of trauma centre database
	Country: USA
	Inclusion criteria: adults with blunt trauma between November 200 and October 2001; and having X-ray and CT of cervical spine
	Exclusion criteria: inadequate radiographs or a recommendation for cervical CT scan
	Adults with blunt trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test

Reference	Griffen 2003 ²²			
reference standard	X-ray of cervical spine			
	CT of cervical spine			
	Reference standard			
	Unclear, possibly all imaging/follow-up			
	Institutional protocol included initial physical exam of cervical spine. Those with reliable examinations and no neck pain or tenderness were clinically cleared by physical exam alone. Stabilisation collars removed and no further evaluation of cervical spine performed. Patients with neck tenderness, neurologic deficits, altered mental status or distracting pain from other injuries all underwent standard three-view cervical spine radiography and cervical spine CT scan. If these indicate negative results, cervical collar left in place until a reliable physical examination can be performed. Patients returning to clinic with continued cervical tenderness have flexion-extension radiographs to rule out ligamentous injuries. Those with persisting tenderness and negative radiography/CT including flexion-extension views after 1 month and those that develop any neurologic deficit referred to spine service for final clearance.			
Outcome	Cervical spine injury – poorly defined			
2×2 table	Cervical spine injury – X-ray as index test			
	Reference standard Reference standard Total + - -			

Reference	Griffen 2003 ²²				
	Index test +	75	NR	NR	Limited reporting of raw data means only sensitivity can be calculated.
	Index test -	41	NR	NR	
	Total	116	1083	1199	Of the 41 injuries missed, most were managed with cervical collar for 6 weeks, n=9 had an external stabilisation device, n=3 required surgical stabilisation and =2 died of associated injuries before full evaluation and treatment of cervical
					spine.
	Cervical spine in	njury – CT as index te	st		
		Reference standard +	Reference standard -	Total	Limited reporting of raw data means only sensitivity can be
	Index test +	116	NR	NR	calculated.
	Index test -	0	NR	NR	
	Total	116	1083	1199	
Statistical measures	Cervical spine ir	njury – X-ray as index	test: calculated using e	excel sheet	
	Sensitivity: 65.0%				

Reference	Griffen 2003 ²²
	Specificity: NR
	PPV: NR
	NPV: NR
	Cervical spine injury – CT as index test: calculated using excel sheet
	Sensitivity: 100.0%
	Specificity: NR
	PPV: NR
	NPV: NR
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if index test and reference standard interpreted without knowledge of the other, unclear time interval between index test and reference standard and likely that reference standard components differed between patients – applies to both index tests
	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, and unclear if reference standard included a 2 week follow-up period
Comments	None

Reference	Inaba 2016 ²⁷
Study type	Prospective
Study methodology	Data source: multi-centre prospective observational trial performed at 18 level 1 and 2 trauma centres
	Recruitment: multi-centre across 18 level 1 and 2 trauma centres in North America through Western Trauma Association Multi-institutional Trials group
Number of patients	n = 10,276
	(n=10,765 patients matched entry criteria, with n=489 subsequently excluded due to previous spinal surgery, outside hospital transfer or both; leaving n=10,276 analysed in the study)
Patient characteristics	Age, mean (range): 48.1 (18.0-110.0) years
	Gender (male to female ratio): 66.7% males
	Ethnicity: not reported
	Head injury: 3.6% said to be unevaluable due to a TBI – unclear if/proportion of others that were evaluable and had suspected or confirmed head injury
	Mechanism of injury:
	 Motor vehicle collision, 30.0% Ground level fall, 20.9%

Reference	Inaba 2016 ²⁷
	 Fall from height, 11.9% Other, 10.2%
	 Automobile vs. pedestrian, 9.0%
	Assault, 7.0%
	 Motorcycle collision, 6.9% Bicycle vs. automobile, 3.8%
	GCS at admission, median (IQR): 15 (14-15)
	Injury Severity Score, median (IQR): 9 (4-16)
	Neurological exemination:
	Neurological examination:
	 Unevaluable, 45.3% TBI, 3.6%
	 Distracting injury, 4.3%
	 Intoxicated/intubated, 11.4% Combination 26.0%
	 Combination, 26.0%
	• Evaluable, 54.7%
	 No deficit, 49.0% Motor deficit, 2.4%
	 Sensory deficit, 1.8%
	 Motor/sensory deficit, 1.5%
	Type of imaging:
	 ○ CT, 100.0%

Reference	Inaba 2016 ²⁷
	 MRI, 9.2% Plain X-ray, 1.4%
	 Flexion-extension X-ray, 0.4%
	Hospital length of stay, median (IQR): 2 (1-6) days
	ICU length of stay, median (IQR): 0 (0-1.4) days
	Setting: secondary care – multiple trauma centres
	Country: USA
	Inclusion criteria: blunt trauma; ≥18 years; and failing NEXUS 2 low risk criteria
	Exclusion criteria: transferred from an outside facility; had a history of spinal instrumentation; did not undergo
	diagnostic imaging with CT Scan of their C-spine; and cervical spine imaging from outside hospitals
	Adults with blunt trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	CT scan of cervical spine

Reference	Inaba 2016 ²⁷				
	Reference standard				
	Final diagnosis a	t discharge, including re	esults of all imaging and	d operative findings	8.
	-	8 8 8		8	History and physical exam
	(NEXUS criteria, presence or absence of midline C-spine tenderness and results of neurological examination) performed by senior resident or faculty member using structured form. All imaging interpreted by attending radiologist blinded to study case report form contents and final attending radiologist read was used in analysis.				
		easurement of index tes period than 2 weeks as		•	of stay was 2 (IQR 1-6) days –
Outcome	Clinically significant cervical spine fracture – an abnormal or equivocal finding observed on either CT or MRI consistent with acute traumatic injury was necessary, along with one of three active interventions: surgical stabilization, Halo Orthotic placement or use of a Cervical-Thoracic Orthotic.				
2×2 table	CT scan of cervical spine as index test – clinically significant cervical spine fracture				
		Reference standard +	Reference standard	Total	Raw data incompletely reported but calculated from
	Index test +	195	907	1102	sensitivity/specificity etc. reported in the paper
	Index test -	3	9171	9174	
	Total	198	10,078	10,276	

Reference	Inaba 2016 ²⁷
Statistical measures	CT scan of cervical spine as index test – clinically significant cervical spine fracture: reported in the paper
	Sensitivity: 98.5%
	Specificity: 91.0%
	PPV: 17.8%
	NPV: 99.97%
Source of funding	Reported that there were no funding disclosures
Limitations	Risk of bias: very serious – convenience sampling rather than consecutive patients enrolled, reference standard of final diagnosis at discharge does not involve a period of at least 2 weeks since admission, unclear if reference standard was interpreted without knowledge of index test and likely that reference standard different slightly between patients (e.g. any further tests performed) Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, and reference standard indirectness as outcome only includes fractures and does not involve a period of 2 weeks follow-up as specified in the protocol
Comments	None
Reference	Lau 2018 ²⁸
Study type	Retrospective
Study methodology	Data source: retrospective review of data from level 1 trauma centre
	Recruitment: retrospective review of level 1 trauma centre data between 1 st January 2008 and 31 st December 2012

Reference	Lau 2018 ²⁸
Number of patients	n = 63 (n=66 met inclusion criteria, with n=3 of these excluded based on exclusion criteria; leaving n=63 included in the analysis)
Patient characteristics	Age, mean (SD): 42.3 (18.2) years Gender (male to female ratio): 90.5% males
	Ethnicity: • Chinese, 63.5% • Malaysian, 11.1% • Indian, 19.0% • Other, 6.4%
	Head injury: unclear if all or most had head injury but suggests all may have undergone assessment for brain injuries (limited information)
	Mechanism of injury:
	 Fall from height, 17.5% Fall from standing height, 19.0% Road traffic accident: Motorcyclist, 31.7% Car, 15.9%

Reference	Lau 2018 ²⁸
	 Lorry/van, 7.9% Cyclist/pedestrian, 3.2%
	Direct blunt force, 4.8%
	GCS at initial trauma, mean (SD): not reported
	Injury Severity Score, mean (range): not reported
	Suspected injury level:
	 Cervical spine, 81.0% Thoracic spine, 22.2% Lumbar spine, 4.8%
	Neurology:
	 Normoreflexia, 36.4% Upper limb areflexia, 50.9% Lower limb areflexia, 61.8% Lax anal tone, 41.8% Unable to assess, 12.7%
	Setting: secondary care – trauma centre
	Setting: secondary care – trauma centre

Reference	Lau 2018 ²⁸
	Country: Singapore
	Inclusion criteria: blunt trauma; obtunded (GCS ≤8); and admitted to ICU unit.
	Exclusion criteria: incomplete data due to electronic downtime during admission; transferred from another hospital with CT or MRI scans already performed; requiring emergency surgery following CT scan as a form of resuscitation; and would not be able to perform an MRI scan due to medical reasons
	Obtunded people with blunt trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury but suggests all may have undergone assessment for brain injuries (limited information)
Index test(s) and	Index test
reference standard	CT of cervical spine
	MRI of cervical spine
	Reference standard
	MRI reported in paper to be reference standard, but not in line with this review protocol. Therefore, data available used to calculate sensitivity of CT and MRI using CT + MRI as combined reference standard. This means specificity could not be calculated.

Reference	Lau 2018 ²⁸
	Based on clinical workflow at the institution, all suffering blunt injuries and that are obtunded are evaluated with CT on emergency basis once initial resuscitation performed. CT performed as non-contrast study for head, cervical spine, thorax, abdomen and pelvis with 10 mm axial cuts. Purpose is to assess for cervical spine injuries at the same time as brain and visceral injuries. All apart for those requiring emergency surgery will be scheduled for interval MRI scan of cervical spine for clearance of injuries as part of standard clinical workflow. MRI performed without contrast within 48 h of admission after patient condition has stabilised. Cervical immobilisation only removed after CT and MRI image reviewed by attending spine or trauma consultant and following confirmation of the final report issued by senior radiologist.
	Time between measurement of index test and reference standard: CT performed when initial resuscitation performed and MRI performed within 48 h of admission after condition has stabilised.
Outcome	Cervical spine injuries – poorly defined but appears to include bony and soft tissue injuries
2×2 table	Data insufficiently reported to complete 2x2 tables.
	7 patients reported to have findings on MRI with no positive findings on CT. Reported that no patients within findings on CT were negative on MRI.
Statistical measures	Cervical spine injury – CT as index test: reported in paper
	Sensitivity: 87.2%
	Specificity: NA
	PPV: NA
	NPV: NA

Reference	Lau 2018 ²⁸
	Any cervical spinal cord injury – MRI as index test: calculated using statement that none that were positive on CT were negative on MRI
	Sensitivity: 100%
	Specificity: NA
	PPV: NA
	NPV: NA
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unlikely that index test (for MRI) or reference standard (for CT) were interpreted without knowledge of reference standard/index test, and mean time interval between index test and reference standard possibly at least 48 h – applies to both index tests
	Indirectness: serious – all included were at obtunded and admitted to ICU representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury
Comments	Not possible to calculate specificity based on the reference standard used.
Reference	Lee 2001 ²⁹
Study type	Retrospective
Study methodology	Data source: retrospective review of patients from single level 1 trauma centre emergency room

Reference	Lee 2001 ²⁹
	Recruitment: retrospective review of single level 1 trauma centre data between January 1999 and June 2000
Number of patients	n = 604(n=3684 adult trauma patients underwent screening of cervical spine, with n=604 included in the analysis as they had both conventional radiography and helical CT)
Patient characteristics	Age, mean (SD): not reported
	Gender (male to female ratio): 50.7% males
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported
	GCS at initial trauma, mean (SD): not reported
	Injury Severity Score, mean (SD): not reported
	Setting: secondary care – data from a trauma centre

Reference	Lee 2001 ²⁹
	Country: USA
	Inclusion criteria: adults suffering trauma presenting to ED; and underwent cervical spine imaging with both plain radiography and helical CT
	Exclusion criteria: those who only had plain radiography or only had helical CT of cervical spine, or where imaging were not available for comparison
	Adults with trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	X-ray of cervical spine
	Reference standard
	Helical CT scan
	Plain radiographs included standard four views (antero-posterior, lateral, and swimmers and open-mouthed odontoid views). Helical CT involved 1 mm collimation helical scanning from foramen magnum level to C3 vertebral body and 3 mm collimation from C3 to T1. Contiguous axial images obtained with bone and soft tissue

Reference	Lee 2001 ²⁹					
	algorithms. Sagittal and coronal reconstructions also obtained. All studies reviewed by radiology resident and neuroradiologist.					
	Time between measurement of index test and reference standard: unclear time interval between index test and reference standard					
Outcome	Cervical spine fra	acture				
2×2 table	Cervical spine f	racture – X-ray as ind	ex test			
		Reference standard +	Reference standard	Total	Raw data given only for individual fractures (with each	
	Index test +	12	NR	NR	patient possibly having more than one fractures – 36 fractures on CT in 30 patients)	
	Index test -	24	NR	NR		
	Total	36	NR	604 patients (raw data given for	Incompletely reported meaning specificity not	
			fractures – some	reported and calculation not possible.		
				patients had more		
				than one fracture)		
Statistical measures	Cervical spine fracture – X-ray as index test (individual fractures identified not patients with fractures): reported in paper					

Reference	Lee 2001 ²⁹
	Sensitivity: 33.3%
	Specificity: NR
	PPV: NR
	NPV: NR
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if reference standard interpreted without knowledge of index test and unclear time interval between index test and reference standard
	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, outcome limited to fractures rather than any cervical spine injury, and results interpreted at fracture level not patient level (patients could have more than one fracture and these included individually in analysis)
Comments	Analyses sensitivity at fracture level and not patient level (each patient could have more than one fracture and these were included separately in calculation of sensitivity)
Reference	Malhotra 2018 ³⁰

Reference	Malhotra 2018 ³⁰
Study type	Retrospective
Study methodology	Data source: radiology database of single tertiary health system and level 1 trauma centre
	Recruitment: retrospective review of database of single tertiary health system and level 1 trauma centre between
	February 2013 and November 2015

Reference	Malhotra 2018 ³⁰
Number of patients	n = 1080
	(n=1271 with blunt cervical spine trauma underwent both a CT and MRI of cervical spine, with n=191 subsequently excluded based on incomplete medical record information, limited CT studies or absent CT reports usually from transferred patients; leaving n=1080 included in the analysis)
Patient characteristics	Age, mean (range): 57 (18-93) years
	Gender (male to female ratio): 55.0% males
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury:
	 Fall from standing, 43.6% Motor vehicle collision-auto, 20.8% Fall from height, 19.5% Motor vehicle collision-pedestrian, 6.7% Assault, 4.7% Motor vehicle collision-motorcycle, 1.3% Sport injury, 2.6% Falling object, 0.7%

Reference	Malhotra 2018 ³⁰
	GCS at initial trauma, mean (SD): not reported
	Injury Severity Score, mean (range): not reported
	Setting: secondary care – trauma centre database
	Country: USA
	Inclusion criteria: adults with suspected blunt trauma to cervical spine; and underwent CT followed by MRI of cervical spine within 48 h
	Exclusion criteria: CT study was non-diagnostic due to patient motion or their medical record was incomplete
	Adults with suspected blunt trauma to cervical spine
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	CT of cervical spine
	MRI of cervical spine

Reference	Malhotra 2018 ³⁰
	Reference standard
	MRI reported in paper to be reference standard, but not in line with this review protocol. Therefore, data available used to calculate sensitivity of CT and MRI using CT + MRI as combined reference standard. This means specificity could not be calculated.
	CT cervical spine images acquired on 64-detector scanners with 1.25 mm slice helical acquisition without intravenous contrast and reformatted in coronal and sagittal planes. Siemens 1.5T and 3T magnets used for MRI scanning, without intravenous contrast using trauma protocol sequences including sagittal T1 FSE, axial and sagittal T2 FSE, sagittal STIR and sagittal GRE sequences.
	CT studies reviewed by neuroradiology fellow to classify into negative or positive for acute traumatic injury based on final report given by ED at time of scan. Studies that were unequivocally negative for injury were classified as negative CT. Studies were positive on CT if impressions included any of the following features: fractures of occipital condyles or C1-C7 vertebral bodies, disc space widening, vertebral subluxation, prevertebral or paravertebral oedema and haematoma, epidural haematoma, cord haematoma or new disc herniation.
	On MRI, studies were positive if contained any of the following features: fractures of occipital condyles or C1-C7 vertebral bodies, osseous oedema or contusion, ligamentous injury or paravertebral muscle strain, spinal cord oedema or haemorrhage, epidural/subdural haematoma, new or acute disc herniation, and prevertebral oedema or haematoma. MRI studies interpreted unequivocally as negative for any of the above findings were classified as negative MRI. MRI findings were confirmed by neuroradiology faculty with 8 years' experience blinded to patient characteristics, outcome, management and report contents other than the impression.
	Time between measurement of index test and reference standard: CT performed and MRI performed within 48 h.
Outcome	Any cervical spine injury – including osseous and ligamentous injuries
2×2 table	Any cervical spine injury – CT as index test

Reference	Malhotra 2018 ³⁰				
		Reference standard +	Reference standard	Total	Note false positives were not possible using CT + MRI as a reference standard meaning specificity could not be calculated.
	Index test +	368	NA	368	
	Index test -	149	563	712	
	Total	517	563	1080	
	Any cervical sp	ine injury – MRI as inc	lex test		
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a reference standard meaning specificity could not be calculated.
	Index test +	427	NA	427	
	Index test -	90	563	653	
	Total	517	563	1080	
Statistical measures	Any cervical spine injury – CT as index test: calculated using excel sheet				
	Sensitivity: 71.0				
	Specificity: NA				
	PPV: NA				
	NPV: NA				

Reference	Malhotra 2018 ³⁰
	Any cervical spine injury – MRI as index test: calculated using excel sheet
	Sensitivity: 83.0
	Specificity: NA
	PPV: NA
	NPV: NA
Source of funding	Stated that no funding was received
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unlikely that index test (for MRI) or reference standard (for CT) were interpreted without knowledge of reference standard/index test, and mean time interval between index test and reference standard likely at least 48 h – applies to both index tests
	Indirectness: serious – head injury not mentioned and unclear if all or most had head injury
Comments	Not possible to calculate specificity based on the reference standard used.

Reference	Mathen 2007 ³¹
Study type	Prospective
Study methodology	Data source: performed at single level 1 trauma centre
	Recruitment: unblinded prospective consecutive series design at a single level 1 trauma centre between October 2004 and February 2005. All presenting to institution prospectively enrolled into study protocol.
Number of patients	n = 667(n=682 matching inclusion criteria, with n=6 dying before cervical spine evaluation and n=9 only having CT rather than both X-ray and CT excluded from the analysis; leaving n=667 included)

Reference	Mathen 2007 ³¹
Patient characteristics	Average age: 35.4 years
	Gender (male to female ratio): not reported
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury:
	 Motor vehicle collision, 48.7% Pedestrian hit by auto, 14.4% Falls, 13.5%
	Average GCS score: 13.2
	Injury Severity Score, mean (SD): not reported
	Setting: secondary care – trauma centre

Reference	Mathen 2007 ³¹
	Country: USA
	Inclusion criteria: not meeting NEXUS low-risk criteria; and undergoing multi-slice CT and 3-view plain radiography of cervical spine
	Exclusion criteria: death before completion of both CT and plain radiography of cervical spine
	People with trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	X-ray of cervical spine
	Multi-slice CT of cervical spine
	Reference standard
	Final diagnosis of cervical spine injury based on all prospectively collected clinical data and imaging results, unclear follow-up duration
	All CTs (occiput to T1) performed using four-channel CT scanner with collimation of 2 mm. Coronal and sagittal reformation images using 1.5 mm to 2 mm intervals reconstructed from axial source images. Three-view plain radiography (X-ray) included anterior-posterior, lateral and odontoid views. Additional views (swimmers,

Reference	Mathen 2007 ³¹					
	obliques) performed at discretion of attending radiologist. Final radiographic interpretation of CT and plain films performed by board-certified radiologists.					
	Time between measurement of index test and reference standard: unclear time interval between index tests and subsequent tests/final confirmed diagnosis					
Outcome	Any acute cervical spine injury – acute fracture or subluxation, or both					
	and					
	Clinically signification	ant cervical spine injury	- requiring surgery or I	ong-term stabilisat	ion with a collar or halo	
2×2 table	Any acute cervi	Any acute cervical spine injury – X-ray as index test				
		Reference standard +	Reference standard	Total		
	Index test +	27	16	43		
	Index test -	33	591	624		
	Total	60	607	667		
	Any acute cervical spine injury – CT as index test					
	Reference standard Reference standard Total					

Reference	Mathen 2007 ³¹				
	Index test +	60	3	63	
	Index test -	0	604	604	
	Total	60	607	667	
	Clinically significant cervical spine injury – X-ray as index test				
		Reference standard +	Reference standard -		
	Index test +	12	31	43	
	Index test -	15	609	624	
	Total	27	640	667	
	Clinically significant cervical spine injury – CT as index test				
		Reference standard +	Reference standard -		
	Index test +	27	36	63	
	Index test -	0	604	604	
	Total	27	640	667	
Statistical measures	Any acute cervical spine injury – X-ray as index test: reported in paper				

Reference	Mathen 2007 ³¹
	Sensitivity: 45.0%
	Specificity: 97.4%
	PPV: 62.8%
	NPV: 94.7%
	Any acute cervical spine injury – CT as index test: reported in paper
	Sensitivity: 100.0%
	Specificity: 99.5%
	PPV: 95.2%
	NPV: 100.0%
	Clinically significant cervical spine injury – X-ray as index test: calculated using excel sheet
	Sensitivity: 44.0%
	Specificity: 95.0%
	PPV: 28.0%
	NPV: 98.0%
	Clinically significant cervical spine injury – CT as index test: calculated using excel sheet

Reference	Mathen 2007 ³¹
	Sensitivity: 100.0%
	Specificity: 94.0%
	PPV: 43.0%
	NPV: 100.0%
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if reference standard interpreted without knowledge of index test, unclear interval between index test and reference standard and unclear if reference standard contained the same components for all patients – applies to both index tests
Comments	None
Reference	Nguyen 2005 ³³
Study type	Prospective
Study methodology	Data source: observational study at level 1 trauma centre hospital with data recorded prospectively

Recruitment: prospective of all patients with trauma and undergoing imaging across a 70-day period

Number of patients n = 112 (n=78 in low risk group and n=34 in high risk group)

Reference	Nguyen 2005 ³³
	(n=219 patients meeting inclusion criteria for the paper, with n=112 analysed as they had both plain radiography and CT of the cervical spine)
Patient characteristics	Age, range: 2-89 years for low risk group and 11-88 years for high risk group
	Gender (male to female ratio): 47.4% males in low risk group and 64.7% males in high risk group
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported
	GCS at initial trauma, mean (SD): not reported
	Injury Severity Score, mean (SD): not reported
	Setting: secondary care – trauma centre hospital
	Country: USA

Reference	Nguyen 2005 ³³
	Inclusion criteria: patients with blunt trauma; and underwent imaging of cervical spine Exclusion criteria: not reported
	Patients with blunt trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	X-ray of cervical spine
	CT of cervical spine
	Reference standard
	Diagnosis based on final reports including all imaging, unclear duration of follow-up
	Treating physicians ordered films at their discretion. All major trauma patients screened with standard 3-view cervical spine radiography (cross-table lateral, antero-posterior and odontoid views) and CT. Cervical spine CT performed using Siemens Somatom CT scanner (3 mm slices, four detector rows) with soft tissue window and bone window with sagittal and coronal reconstructions. Injury status determined based on all radiographic studies reviewed and final report.

Reference	Nguyen 2005 ³³	Nguyen 2005 ³³					
		Time between measurement of index test and reference standard: unclear duration between index tests and other imaging/final diagnosis being confirmed					
Outcome	Cervical spine f risk in review p	•	lts separately for risk wl	nich are extracted	separately as strata based on		
2×2 table	Cervical spine	Cervical spine fracture – X-ray as index test (low-risk group)					
		Reference standard +	Reference standard	Total	Sensitivity not applicable as none in this group had		
	Index test +	0	0	0	fracture.		
	Index test -	0	78	78			
	Total	0	78	78			
	Cervical spine	fracture – CT as index	test (low-risk group)				
		Reference standard +	Reference standard	Total	Sensitivity not applicable as none in this group had		
	Index test +	0	0	0	fracture.		
	Index test -	0	78	78			
	Total	0	78	78			
	Cervical spine	fracture – X-ray as inde	ex test (high risk grou	n)			

Cervical spine fracture – X-ray as index test (high risk group)

Reference	Nguyen 2005 ³³				
		Reference standard +	Reference standard	Total	Raw data incompletely reported but missing values
	Index test +	14	1	15	calculated using excel sheet and reported sensitivity/specificity in paper
	Index test -	1	18	19	
	Total	15	19	34	Missed injury was non- displaced fracture through C7 left facet. No soft tissue abnormality associated with it and no misalignment.
	Cervical spine f	racture – CT as index	test (high risk group)		
		Reference standard +	Reference standard -	Total	Raw data incompletely reported but missing values
	Index test +	15	0	15	calculated using excel sheet and reported
	Index test -	0	19	19	sensitivity/specificity in paper
	Total	15	19	34	
Statistical measures	Cervical spine fracture – X-ray as index test (low risk group): reported in paper (apart from NPV which was calculated using excel sheet)				
	Sensitivity: NA				
	Specificity: 100.0	%			

Reference	Nguyen 2005 ³³
	PPV: NA
	NPV: 100.0%
	Cervical spine fracture – CT as index test (low risk group): reported in paper (apart from NPV which was calculated using excel sheet)
	Sensitivity: NA
	Specificity: 100.0%
	PPV: NA
	NPV: 100.0%
	Cervical spine fracture – X-ray as index test (high risk group): reported in paper (apart from NPV and PPV which were was calculated using excel sheet)
	Sensitivity: 93.3%
	Specificity: 95.0%
	PPV: 94.0%
	NPV: 95.0%
	Cervical spine fracture – CT as index test (high risk group): reported in paper (apart from NPV and PPV which were was calculated using excel sheet)
	Sensitivity: 100.0%

Reference	Nguyen 2005 ³³
	Specificity: 100.0%
	PPV: 100.0%
	NPV: 100.0%
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if index test and reference standard interpreted without knowledge of the other, unclear time interval between index test and reference standard and likely that reference standard components differed between patients – applies to both index tests
	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, outcome focuses only on fractures and unclear if reference standard included a 2 week follow-up period
Comments	None

Reference	Novick 2018 ³⁵
Study type	Retrospective
Study methodology	Data source: retrospective review of data from single level 1 trauma centre serving population of 1.4 million. Trauma registry queried for data and cross-referenced with radiology record system and main hospital medical record system.
	Recruitment: retrospective review of those matching inclusion criteria from single level 1 trauma centre between 1 st January 2008 and 31 st December 2015.
Number of patients	n = 241

Reference	Novick 2018 ³⁵
	(included n=241 that had both CT and MRI of cervical spine – flow of patients and those excluded unclear)
Patient characteristics	Age, mean (range): 43.9 (5-93) years
	Gender (male to female ratio): 60.2% males
	Ethnicity: not reported
	Head injury: unclear if all or most had head injury – 17% reported to have closed head injury, but unclear for others if head injury was part of the injury mechanism Mechanism of injury:
	 Assault, 3.3% Cyclist, 2.1% Fall from standing, 20.3% Fall >1 m, 6.2% Fall stairs, 6.6% Hanging, 1.2% Motorcycle crash, 2.1% Motor vehicle crash, 45.6% Sports-related, 3.3% Struck in head, 0.8% Pedestrian struck, 7.9% Gunshot wound to neck, 0.4%

Reference	Novick 2018 ³⁵
	GCS at initial trauma, mean (SD): not reported
	Injury Severity Score, mean (SD): not reported
	Indication for MRI:
	 Neck pain, 57.7% Abnormal neurologic exam, 34.0% Unable to assess due to: Closed head injury, 17.4% Drugs/alcohol, 9.1% Post-ictal, 2.1% Abnormal CT, 36.5% No signs or symptoms, 2.9%
	Setting: secondary care – trauma centre
	Country: USA
	Inclusion criteria: underwent both CT and MRI of cervical spine; and history of trauma
	Exclusion criteria: not reported

Reference	Novick 2018 ³⁵
	People with trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury – 17% reported to have closed head injury, but unclear for others if head injury was part of the injury mechanism
Index test(s) and	Index test
reference standard	CT of cervical spine
	MRI of cervical spine
	Reference standard
	Reference standard not reported in the paper but possible to calculate sensitivity of both imaging tests using CT + MRI as combined reference standard. This means specificity could not be calculated.
	Cervical spine clearance protocol involved clinical confrontational exam for neck pain, neurologic examination and CT of cervical spine. MRI was obtained immediately after CT if neurologic examination was abnormal or if CT indicated an abnormality. If clinical exam identified neck pain or could not be performed, or if the CT interpreted by radiologist as equivocal for abnormality or injury, an MRI was obtained within 48 h of admission.
	CT images obtained using 1320 or 16-slice machine. Routine trauma protocol consisted of multiple contiguous non-contrast axial sections are obtained from the posterior fossa to the cervical-thoracic junction without the intravenous administration of contrast. Multiplanar reformation was uniformly performed in the coronal and sagittal planes. Screening CT cervical spine slice thickness (acquired at 0.5 mm) with coronal and sagittal reformations was same on both CT scanners with 1-mm cuts and 3-mm reconstruction for coronal and sagittal images.

Reference	Novick 2018 ³⁵					
	MRI images obtained with 1.5 T magnet, performed in multiple planes and sagittal T1-weighted, T2-weighted, and short-tau inversion-recovery sequences (3 mm thick), as well as an axial T2*-weighted sequence (3 mm thick).					
	vertebra and artit	Studies assessed as technically adequate if images were obtained from the base of the skull to the first thoracic vertebra and artifact or motion did not markedly limit the evaluation. Studies assesses as not technically adequate by the radiology technician or physician were immediately repeated.				
	Time between measurement of index test and reference standard: CT performed followed by MRI where indicated, duration between the two differed depending on presentation but could be up to 48 h.					
Outcome		Cervical spine injuries – ligamentous or bony injury of the cervical vertebral spine, disc injuries, or spinal cord injuries as assessed by imaging				
2×2 table	Cervical spine i	njury – CT as index te	st			
		Reference standard +	Reference standard	Total	Note false positives were not possible using CT + MRI as a	
	Index test +	88	NA	88	reference standard meaning specificity could not be	
	Index test -	13	140	153	calculated.	
	Total	101	140	241		
	Cervical spine injury – MRI as index test					
	Reference standard + Reference standard Total Note false positives we possible using CT + MF					

Reference	Novick 2018 ³⁵						
	Index test +	78	NA	78	reference standard meaning specificity could not be		
	Index test -	23	140	163	calculated.		
	Total	101	140	241			
Statistical measures	Cervical spinal of	Cervical spinal cord injury – CT as index test: calculated using excel sheet					
	Sensitivity: 87.0						
	Specificity: NA	Specificity: NA					
	PPV: NA NPV: NA						
	Cervical spinal cord injury – MRI as index test: calculated using excel sheet						
	Sensitivity: 77.0						
	Specificity: NA	Specificity: NA					
	PPV: NA						
	NPV: NA						
Source of funding	Not reported						
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unlikely that index test (for MRI) or reference standard (for CT) were interpreted without knowledge of reference standard/index test, and mean time interval between index test and reference standard likely at least 48 h – applies to both index tests						

cted head injury/head
С

Reference	Parmar 2018 ³⁶
Study type	Prospective
Study methodology	Data source: clinical audit of single hospital ED. Data obtained from Department of Radiology PACS computer- based database, ICU electronic records and trauma registry.
	Recruitment: consecutive patients matching inclusion criteria between 9 th October 2015 and 8 th May 2016 from a single hospital ED
Number of patients	n = 27 analysed
	(n=100 unconscious patients identified, with n=27 analysed as they had both CT and MRI of cervical spine)
Patient characteristics	Note: characteristics only given for n=100 in whole study not the n=27 analysed
	Age, median (IQR): 38.5 (25-53) years
	Gender (male to female ratio): 81% males

Reference	Parmar 2018 ³⁶
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported
	GCS at initial trauma, mean (SD): not reported
	Injury Severity Score, median (IQR): 26 (12-33)
	Setting: secondary care – ED of hospital
	Country: Australia
	Inclusion criteria: adults that were unconscious/obtunded and admitted to the ED; and requiring artificial airway and mechanical ventilation – those included in analysis had to have CT and MRI of cervical spine
	Exclusion criteria: not reported

Reference	Parmar 2018 ³⁶
	Unconscious/obtunded adults admitted to the ED and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided (assumed to have head injury based on severity of injuries – all obtunded)
Index test(s) and	Index test
reference standard	CT of cervical spine
	MRI of cervical spine
	Reference standard
	MRI reported in paper to be reference standard, but not in line with this review protocol. Therefore, data available used to calculate sensitivity of CT and MRI using CT + MRI as combined reference standard. This means specificity could not be calculated.
	In the study centre, hard neck collar used routinely to immobilise cervical spine until cervical spine injuries can be excluded based on CT scan as primary imaging modality. If no bony injury or mal-alignment identified on CT, cervical spine is considered radiologically cleared with no further spinal precautions needed. Time-point <48 h used to confirm injury status of cervical spine as benchmark as Australian ICU clearance protocols recommend 48 h or less. Consultant radiologists were on-call and not on-site during off-office ours and were not informed of the audit.
	Time between measurement of index test and reference standard: unclear duration between CT and MRI.
Outcome	Any cervical spine injury – poorly defined but includes osseous and ligamentous injuries
2×2 table	Any cervical spine injury – CT as index test

Reference	Parmar 2018 ³⁶				
		Reference standard +	Reference standard	Total	Note false positives were not possible using CT + MRI as a reference standard meaning specificity could not be calculated.
	Index test +	20	NA	20	
	Index test -	7	0	7	
	Total	27	0	27	
	Any cervical sp	ine injury – MRI as ind	lex test		
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a
	Index test +	26	NA	26	reference standard meaning specificity could not be calculated.
	Index test -	1	0	1	
	Total	27	0	27	
Statistical measures	Any cervical sp	inal cord injury – CT a	s index test: calculate	d using excel shee	t
	Sensitivity: 74.0				
	Specificity: NA				
	PPV: NA				
	NPV: NA				

Reference	Parmar 2018 ³⁶
	Any cervical spinal cord injury – MRI as index test: calculated using excel sheet
	Sensitivity: 96.0
	Specificity: NA
	PPV: NA
	NPV: NA
Source of funding	Funding support from WA Health and Raine Medical Research Foundation through Raine Clinical Research Fellowship.
Limitations	Risk of bias: very serious – unlikely that index test (for MRI) or reference standard (for CT) were interpreted without knowledge of reference standard/index test, and mean time interval between index test and reference standard likely at least 48 h – applies to both index tests
	Indirectness: serious – all included were obtunded representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury
Comments	Not possible to calculate specificity based on the reference standard used.
Reference	Ptak 2001 ³⁷
Study type	Retrospective
Study methodology	Data source: retrospective cross-sectional analysis from single general hospital, using radiology report database.
	Recruitment: retrospective review of those matching inclusion criteria from single general hospital between 1 st July 1997 and 31 st August 1998.

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Reference	Ptak 2001 ³⁷
Number of patients	n = 676
	(n=2466 cervical spine CT studies identified, of which a subgroup of n=1182 cervical spine studies in trauma patients was selected; within this trauma subgroup, n=676 conforming to screening trauma cervical spine protocol were included)
Patient characteristics	Age, mean (SD): 47.2 (24.1) years, range 1-104 years
	Gender (male to female ratio): 66.0% males
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported
	GCS at initial trauma, mean (SD): not reported
	Injury Severity Score, mean (SD): not reported
	Setting: secondary care – records from a general hospital

Reference	Ptak 2001 ³⁷
	Country: USA
	Inclusion criteria: presenting to emergency radiology division for CT evaluation of cervical spine injuries following trauma (Massachusetts General Hospital procedure code #644 – having CT of cervical spine); and CT was initial screening evaluation of cervical spine following trauma
	Exclusion criteria: non-traumatic injuries; cases where CT was preceded by a plain film series (more than one portable lateral film) of cervical spine; and cases not performed according to standardised ED helical CT protocol for cervical spine screening or those where protocol could not be confirmed
	People sustaining trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	Helical CT of cervical spine
	Reference standard
	Final clinical diagnosis (including operative and discharge notes), possibly incorporating CT results

Reference	Ptak 2001 ³⁷				
	After initial clinical evaluation and portable plain film trauma series including anteroposterior supine chest, pelvis and cross-table lateral view of cervical spine while immobilised in the trauma bay, patients transferred to CT suite. Screening cervical spine images acquired on CT scanner using helical technique with 3 mm beam collimation and pitch of 1.5. Images reconstructed to 3 mm spacing using high spatial frequency bone algorithm. Images acquired from skull base to T2 vertebral body. Images immediately post-processed into 1 mm spacing using detail spatial frequency algorithm, from which 2D coronal and sagittal reformations constructed. Optional 3D reformations available.				
Outcome	Cervical spine fracture – no further definition provided Positive CT cases for fracture taken as reported in the radiological report.				
2×2 table	Cervical spine fracture – Helical CT as index test				
		Reference standard +	Reference standard -	Total	
	Index test +	59	0	59	
	Index test -	1	616	617	
	Total	60	616	676	
Statistical measures	Cervical spine f	racture – Helical CT as	s index test: reported i	n paper	

Reference	Ptak 2001 ³⁷
	Sensitivity: 98.3%
	Specificity: 100.0%
	PPV: 100.0%
	NPV: 99.8%
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if reference standard interpreted without knowledge of index test, unclear time interval between index test and reference standard and likely that reference standard components differed between patients
	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, outcomes focuses only on fractures and unclear if reference standard included a 2 week follow-up period
Comments	None

Reference	Raza 2013 ⁴⁰
Study type	Retrospective
Study methodology	Data source: review of medical records of people presenting to ED of single hospital retrospectively
	Recruitment: people presenting to ED of single hospital in London between October 2007 and December 2008 retrospectively reviewed
Number of patients	n = 53

Reference	Raza 2013 ⁴⁰
	(n=108 presenting to ED were reviewed, with n=53 included as they matched inclusion criteria)
Patient characteristics	Age, mean (SD): not reported
	Gender (male to female ratio): not reported
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported
	GCS at initial trauma, mean (SD): not reported
	Injury Severity Score, mean (SD): not reported
	Setting: secondary care – ED of hospital
	Country: UK

Reference	Raza 2013 ⁴⁰
	Inclusion criteria: adult blunt trauma patients with GCS ≤14 (altered sensorium/obtunded); intoxicated with alcohol or drugs; and cervical spine multidetector CT obtained on admission Exclusion criteria: fracture identified on initial cervical spine multidetector CT; became examinable before additional CS imaging; died before cervical spine clearance; discharge records not available; and those presenting prior to October 2007
	Adults with altered sensorium/obtunded following blunt trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided (assumed to have head injury based on severity of injuries – all with altered sensorium/obtunded)
Index test(s) and	Index test
reference standard	CT of cervical spine
	Reference standard
	Final diagnosis of injury at hospital discharge, follow-up appointments or any readmissions, possibly includes follow-up of at least 2 weeks specified in the protocol given readmissions and follow-up appointments considered
	PACS and electronic patient records reviewed for patient records and imaging reports in addition to hand search of hospital notes.

Reference	Raza 2013 ⁴⁰						
	Time between measurement of index test and reference standard: unclear						
Outcome	Clinically significant cervical spine injury – poorly defined						
2×2 table	Insufficient reporting of data to be able to calculate 2x2 tables.						
Statistical measures	Clinically significant cervical spine injury – CT as index test: reported in paper Sensitivity: 100% Specificity: NR PPV: NR NPV: NR						
Source of funding	Not reported						
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if reference standard of final diagnosis includes a period of at least 2 weeks follow-up, reference standard likely not interpreted without knowledge of index test and likely that reference standard different slightly between patients (e.g. any further tests performed)						
	Indirectness: serious – all included were obtunded/had altered sensorium representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury						
Comments	Poor reporting in the paper means specificity is not reported and it is not possible to calculate it.						

Reference	Resnick 2014 ⁴¹
Study type	Prospective
Study methodology	Data source: prospective study of patients from single level 1 trauma centre
	Recruitment: prospective observational study of consecutive adults matching inclusion criteria between 1 st January 2010 and 31 st May 2011 at single level 1 trauma centre
Number of patients	n = 830
	(n=3801 matching inclusion criteria, with n=830 patients requiring imaging as they could not be cleared clinically subsequently included in the analysis)
Patient characteristics	Age, mean (SD): 42.6 (18.0) years
	Age >55 years, 22.4%
	Gender (male to female ratio): 70.6% males
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury:
	 Motor vehicle collision, 39.8% Fall, 31.6%
	 Auto vs. pedestrian, 15.5%

Reference	Resnick 2014 ⁴¹
	 Assault, 8.4% Motorcycle collision, 4.0% Other, 0.7%
	GCS at initial trauma, mean (SD): not reported
	Injury severity indices:
	 Injury Severity Score (ISS), mean (SE): 3.3 (2.5) ISS >25, 0.0% Chest AIS ≥3, 0.0% Abdomen AIS ≥3, 0.0% Extremities AIS ≥3, 0.0%
	Setting: secondary care – trauma centre
	Country: USA
	Inclusion criteria: adults (>18 years) sustaining blunt trauma; deemed eligible for evaluation (GCS 15, not intoxicated and with no distracting injury); underwent CT evaluation of cervical spine; and admitted to centre between 1 st January 2010 and 31 st May 2011.
	Exclusion criteria: deemed ineligible for evaluation (GCS <15; intoxicated; or with a distracting injury)

Reference	Resnick 2014 ⁴¹
	Adults with blunt trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	CT of cervical spine
	Reference standard
	Final diagnosis at time of discharge (including all imaging and operative findings) – unclear duration of follow-up incorporated into this reference standard
	Standardised physical examination of cervical spine performed. Collar removed with in-line immobilisation maintained in supine position. Resident or attending surgeon examined for deformities and midline bony tenderness to palpation. Complete peripheral neurologic exam also performed. Those that were awake, alert and able to be examined and had persistent midline pain, tenderness to palpation or focal neurologic deficit enrolled and had CT of cervical spine. MRI was ordered at discretion of attending surgeon or neurosurgeon. All patients monitored on day of discharge.
	Multidetector-row helical CT performed. Images obtained through occiput to T4. 64-slice scanner variables included no intravenous contrast, 120 kV (p), 100-250 mA, gantry revolution speed 0.5 seconds, beam pitch 0.95 and beam collimation of 64x0.5 mm. Reconstruction performed with 2 mm section thickness in axial, coronal and sagittal planes. Images reviewed in multiple window width and level settings.
	All MRI scans obtained on 1.5T system, including sagittal T1 fast spin echo (FSE), sagittal T2 FSE, sagittal short tau inversion recovery, axial T2 FSE and axial T1 sequences. Images reviewed at 3 megapixel resolution. Final radiology reading provided by board-certified radiologist used for analysis.

Reference	Resnick 2014 ⁴¹	Resnick 2014 ⁴¹				
	(Time between measurement of index test and reference standard: unclear how long between initial CT and final diagnosis at discharge and whether at least 2 weeks as in protocol					
Outcome	Any cervical spin	Any cervical spine injury – any abnormal finding observed on CT or MRI consistent with acute traumatic injury				
	and Clinically significant cervical spine injury – those requiring surgical intervention for stabilisation or halo placement, as well as unstable injuries requiring a hard collar					
2×2 table	Any cervical spi	ine injury – CT as inde	ex test			
		Reference standard +	Reference standard -	Total	Raw data incompletely reported but missing numbers	
	Index test +	149	0	149	calculated using excel sheet and sensitivity/specificity	
	Index test -	15	666	681	values reported in paper	
	Total	164	666	830		
	Clinically significant cervical spine injury – CT as index test					
		Reference standard +	Reference standard –	Total	Raw data incompletely reported but missing numbers	
	Index test +	23	0	23	calculated using excel sheet	

Reference	Resnick 2014 ⁴¹	Resnick 2014 ⁴¹					
	Index test -	0	807	807	and sensitivity/specificity values reported in paper		
	Total	23	807	830			
Statistical measures	Any cervical sp	bine injury – CT as	s index test: reporte	d in paper			
	Sensitivity: 90.9	Sensitivity: 90.9%					
	Specificity: 100.	Specificity: 100.0%					
	PPV: 100.0%	PPV: 100.0%					
	NPV: 97.8%						
	Clinically significant cervical spine injury – CT as index test: reported in paper						
	Sensitivity: 100.	Sensitivity: 100.0%					
	Specificity: 100.	Specificity: 100.0%					
	PPV: 100.0%						
	NPV: 100.0%						
Source of funding	Not reported						
Limitations	follow-up, uncle	ar if reference stan	dard interpreted with		cludes a period of at least 2 weeks x test and likely that reference		

Reference

Resnick 2014⁴¹

	Indirectness: very serious – head injury not mentioned and unclear if all or most had head injury, and unclear if reference standard incorporates 2 week follow-up period specified in the protocol
Comments	None

Reference	Schoenfeld 2018 ⁴²
Study type	Retrospective
Study methodology	Data source: obtained from Partners Health System Research Patient Data Registry which gathers clinical data, demographics, radiology results and operative reports on all patients treated at Brigham and Women's Hospital and Massachusetts General Hospital – two academic level 1 trauma centres
	Recruitment: retrospectively reviewed database containing data from two level 1 trauma centres between 2007 and 2014
Number of patients	n = 668
	(n=8753 deemed eligible, with n=8060 having CT of cervical spine and n=693 having both CT and MRI of cervical spine; number analysed further reduced to n=668 for those with CT and MRI based on propensity matching process)
Patient characteristics	Age, mean (SD): 52.6 (22.7) years in CT group and 54.8 (21.7) years in CT-MRI group
	Gender (male to female ratio): 60.0% in both groups Ethnicity: 72% white in CT group and 76% white in CT-MRI group

Reference	Schoenfeld 2018 ⁴²
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported
	GCS at initial trauma, mean (SD): 13.3 (3.6) in both groups
	Injury Severity Score, mean (SD): 6.0 (9.7) in CT group and 6.2 (9.4) in CT-MRI group
	Setting: secondary care – trauma centres
	Country: USA
	Inclusion criteria: adults receiving CT alone or CT-MRI for primary evaluation of cervical spine injury following trauma between 2007 and 2014
	Exclusion criteria: patients with initial evaluations at outside centres that were transferred for care; prior history of spine surgery or spinal metastases; penetrating trauma; those without clear history of trauma despite cervical spine imaging ordered for other reasons; and those lacking complete radiologist reports, emergency room evaluation and/or surgical reports

Reference	Schoenfeld 2018 ⁴²
	Adults following trauma with suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	CT of cervical spine
	MRI of cervical spine
	Reference standard
	No specific reference standard reported in but data available used to calculate sensitivity of CT and MRI using CT + MRI as combined reference standard. This means specificity could not be calculated.
	CT performed using 128-slice scanner with reference mAs of 180 and 2 mm slice thickness. MRI performed with 1.5 T scanners with axial and sagittal sequences. Imaging results recorded directly from radiologist reports and injury characteristics taken from clinical notes and operative reports.
	Time between measurement of index test and reference standard: unclear duration between CT and MRI
Outcome	Cervical spine injury – poorly defined
	Cervical spine injury (for example) fracture, dislocation, traumatic disc herniation and ligamentous disruption on MRI) with associated change in clinical management or surgical intervention was used as primary outcome.
2×2 table	Cervical spine injury – CT as index test

Reference	Schoenfeld 2018 ⁴²						
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a reference standard meaning specificity could not be calculated.		
	Index test +	195	NA	195			
	Index test -	53	420	473			
	Total	248	420	668	Of the 53 with injury on MRI but not CT, a change of management occurred as a result in n=47 (surgery n=5 and non-operative n=42)		
	Cervical spine i	Cervical spine injury – MRI as index test					
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a reference standard meaning specificity could not be calculated.		
	Index test +	248	NA	248			
	Index test -	0	420	420			
	Total	248	420	668			
Statistical measures	Cervical spine i	njury – CT as index te	st: calculated using exc	cel sheet			
	Sensitivity: 79.0						
	Specificity: NA						
	PPV: NA						

Reference	Schoenfeld 2018 ⁴²
	NPV: NA
	Cervical spine injury – MRI as index test: calculated using excel sheet
	Sensitivity: 100.0
	Specificity: NA
	PPV: NA
	NPV: NA
Source of funding	No funding reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unlikely that index test (for MRI) or reference standard (for CT) were interpreted without knowledge of reference standard/index test, and time interval between index test and reference standard unclear – applies to both index tests
	Indirectness: serious – head injury not mentioned and unclear if all or most had head injury
Comments	Not possible to calculate specificity based on the reference standard used.
Reference	Songur 2020 ⁴⁴
Study type	Retrospective
Study methodology	Data source: cross-sectional study using data from a single ED department at tertiary healthcare centre. Data

obtained from hospital electronic medical records.

Reference	Songur 2020 ⁴⁴
	Recruitment: retrospective inclusion from a single centre between June 2014 and June 2018
Number of patients	n = 195 for any injury and n=88 for unstable injuries
	(n=14,795 with relevant injury codes identified, with n=57 excluded based on a coding error; n=198 identified as having both CT and MRI of the cervical spine, with n=3 of these excluded due to missing data and leaving n=195 for the 'any injury' analysis)
Patient characteristics	Age, mean (SD): 47.34 (21.90) years
	Gender (male to female ratio): 71.8% males Ethnicity: not reported Head injury: no details reported, unclear if all or most suffered concomitant injury to the head Mechanism of injury: • Fall from height, 51.3% • Motor vehicle accident, 33.3% • Pedestrian, 8.7% • Assault, 2.6% • Other mechanism, 4.1% • Unknown, 1.5%

Reference	Songur 2020 ⁴⁴
	GCS at initial trauma, mean (SD): 97.9% had GCS >13 and were alert
	Injury Severity Score, mean (SD): not reported
	Setting: secondary care – ED department
	Country: Turkey
	Inclusion criteria: ICD-10 codes S17 (crushing injury of the neck) or S12 (fracture of cervical vertebra and other parts of the neck); and underwent CT scan followed by MRI within 48 h of admission
	Exclusion criteria: other diagnoses (coding error); non-diagnostic CT results and/or incomplete medical records
	People with trauma to the neck and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided
Index test(s) and	Index test
reference standard	CT of cervical spine
	MRI of cervical spine

Reference	Songur 2020 ⁴⁴
	Reference standard
	MRI reported in paper to be reference standard, but not in line with this review protocol. Therefore, data available used to calculate sensitivity of CT and MRI using CT + MRI as combined reference standard. This means specificity could not be calculated.
	CT scans performed with GE Discovery HD 750 machine. MRI performed either either 1.5 Tesla Siemens Symphony or 3 Tesla Siemens Avanto CMR scanner. All images assessed by emergency medicine specialist, a neuroradiologist and a neurosurgeon to classify interpretations as 'negative' or 'positive' for acute traumatic injury patients. CT was positive if impressions included any of the following features: major fractures of vertebrae, disc space widening, vertebral subluxation, epidural hematoma, and prevertebral or paravertebral oedema/hematoma. MRI was positive if they had any of the following features: ligamentous injury, posttraumatic spinal cord pathological signal changes or haemorrhage, epidural/subdural hematoma, new or acute disc herniation and prevertebral oedema or haematoma. MRI that were unequivocally negative for aforementioned findings were classified as MRI-negative.
	admission.
Outcome	All cervical spine injuries
	And
	Unstable cervical spine injury – based on neurological status of the patient, degree of spinal canal stenosis and degree of instability. Denis' 1983 definition of single-level ligamentous injury extending to two of three columns.

Reference	Songur 2020 ⁴⁴						
2×2 table	Any cervical sp	Any cervical spine injury – CT as index test					
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a		
	Index test +	64	NA	64	reference standard meaning specificity could not be		
	Index test -	20	111	131	calculated.		
	Total	84	111	195			
	Any cervical spine injury – MRI as index test						
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a reference standard meaning specificity could not be		
	Index test +	83	NA	83			
	Index test -	1	111	112	calculated.		
	Total	84	111	195			
	Unstable cervio	cal spine injury – CT as	s index test				
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a		
	Index test +	63	NA	63	reference standard meaning specificity could not be		
	Index test -	18	7	25	calculated.		

Reference	Songur 2020 ⁴⁴					
	Total	81	7	88		
	Unstable cervic	al spine injury – MRI a	is index test			
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a	
	Index test +	81	NA	81	reference standard meaning specificity could not be	
	Index test -	0	7	7	calculated.	
	Total	81	7	88		
Statistical measures	Any cervical sp	ine injury – CT as inde	ex test: calculated using	g excel sheet		
	Sensitivity: 76.0					
	Specificity: NA					
	PPV: NA					
	NPV: NA					
		Any cervical spine injury – MRI as index test: calculated using excel sheet				
	Any cervical sp	ine injury – MRI as inc	lex test: calculated usir	ng excel sheet		
	Any cervical sp Sensitivity: 99.0	ine injury – MRI as inc	lex test: calculated usir	ng excel sheet		
		ine injury – MRI as inc	lex test: calculated usir	ng excel sheet		

Reference	Songur 2020 ⁴⁴
	NPV: NA
	Unstable cervical spine injury – CT as index test: calculated using excel sheet
	Sensitivity: 78.0
	Specificity: NA
	PPV: NA
	NPV: NA
	Unstable cervical spine injury – MRI as index test: calculated using excel sheet
	Sensitivity: 100.0
	Specificity: NA
	PPV: NA
	NPV: NA
Source of funding	Stated that no financial support was received.
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unlikely that index test (for MRI) or reference standard (for CT) were interpreted without knowledge of reference standard/index test, and time interval between index test and reference standard unclear. In addition, for unstable injuries, fewer participants are analysed compared to any injury and unclear why – applies to both index tests

Reference	Songur 2020 ⁴⁴
	Indirectness: serious – head injury not mentioned and unclear if all or most had head injury
Comments	Not possible to calculate specificity based on the reference standard used.

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Reference	Takami 2014 ⁴⁶
Study type	Prospective
Study methodology	Data source: data prospectively collected from people at single emergency outpatient service
	Recruitment: prospective collection people transported to emergency outpatient services at single university from September 2007 to May 2009
Number of patients	n = 179
	(n=179 identified as matching inclusion criteria and n=179 analysed – no further details about exclusion reasons)
Patient characteristics	Age, average (range): 54.3 (4-94) years
	Gender (male to female ratio): 74.9%
	Ethnicity: not reported

Head injury: of n=54 with spinal fra feature), unclear proportion of who	ctures, n=8 had head trauma (intracranial haemorrhage or cranial or facial e group
Mechanism of injury: not reported	
GCS at initial trauma, mean (SD): r	not reported
Injury Severity Score, average (ran	ge): for n=54 with spinal fractures, this was 20.2 (4-70)
Setting: secondary care – emergen	cy outpatient service
Country: Japan	
	nergy trauma as determined by emergency personnel on-site and immobilised emergency outpatient services by an ambulance or air ambulance
Exclusion criteria: not reported	
People with high-energy trauma an	d suspected cervical spine injury

Reference	Takami 2014 ⁴⁶						
Target condition(s)	Suspected cervical spine injury – proportion had concomitant head injury but unclear how many, reported to be 15% in those with fractures						
Index test(s) and	Index test						
reference standard	X-ray of cervical	X-ray of cervical spine					
	Defense et en d						
	Reference stand	ard					
	Full CT of spine						
	Full spine CT performed on same day as arrival. Effective dose of full spine CT calculated in a person with standard body weight. 3D reconstructed image produced using multi-planar construction and presence/absence of fractures determined by two orthopaedic specialists. Plain X-rays of cervical spine examined for all during primary care.						
	Time between measurement of index test and reference standard: unclear but during same admission, possible that reference standard performed prior to index test						
Outcome	Cervical spine fra	acture – no further defin	ition provided				
2×2 table	Cervical spine f	racture – X-ray as inde	ex test				
		Reference standard +	Reference standard -	Total	Raw data only sufficient to		
	Index test +	10	NR	NR	calculate sensitivity		
	Index test -	6	NR	NR			

Reference	Takami 2014 ⁴⁶	Takami 2014 ⁴⁶					
	Total	16	163	179			
Statistical measures	Cervical spine fracture – X-ray as index test: calculated using excel sheet						
	Sensitivity: 63.0%						
	Specificity: NR						
	PPV: NR						
	NPV: NR						
Source of funding	Reported that no benefits or funding was received to support this study.						
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if reference standard and index test were interpreted without knowledge of the other and unclear time interval between index test and reference standard						
	Indirectness: very serious – head injury mentioned for a small proportion of participants but unclear if head injury was part of the injury mechanism for all or most, and outcome focuses specifically on fractures rather than any cervical spine injury						
Comments	Only provides suff	ficient data to calculate	sensitivity and not spe	cificity			
Reference	Tan 2014 ⁴⁷						
Study type	Retrospective						
Study methodology	Data source: elec	tronic medical record d	atabase at a single univ	versity medical cer	ntre		

Reference	Tan 2014 ⁴⁷
	Recruitment: retrospective review of records for those matching inclusion criteria at a single medical centre between January 2008 and December 2010
Number of patients	n = 83
	(n=83 identified as matching inclusion criteria and n=83 analysed – no further details about exclusion reasons)
Patient characteristics	Age, mean (SD): not reported
	Gender (male to female ratio): not reported
	Ethnicity: not reported
	Head injury: all had head injury to be included
	Mechanism of injury: not reported
	GCS, mean: 12.09
	 GCS 3-10, 24.0% GCS 11-14, 76.0%
	Injury Severity Score, mean (SD): not reported

Reference	Tan 2014 ⁴⁷
	Setting: secondary care – university medical centre including those with trauma
	Country: USA
	Inclusion criteria: obtunded patients admitted to centre with diagnosis of intracranial haemorrhage and concomitant history of minor cervical spine trauma; and had both CT and MRI of cervical spine
	Exclusion criteria: not reported
	People with confirmed intracranial haemorrhage and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – all had head injury to be included
Index test(s) and	Index test
reference standard	CT of cervical spine
	MRI of cervical spine
	Reference standard

Reference	Tan 2014 ⁴⁷					
	MRI reported in paper to be reference standard, but not in line with this review protocol. Therefore, data av used to calculate sensitivity of CT and MRI using CT + MRI as combined reference standard. This means specificity could not be calculated. CT performed using Siemens 64-slice machine and MRI using Siemens 1.5T or 3T. Both read by board-ce attending neuroradiologist and an attending neurosurgeon.					
	Time between me	easurement of index tes	st and reference standa	rd: unclear duratio	on between CT and MRI	
Outcome	Any cervical spin	e injury – no definition				
	and Unstable cervical spine injury – no definition					
2×2 table	Any cervical spi	ne injury – CT as inde	ex test			
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a	
	Index test +	28	NA	28	reference standard meaning specificity could not be	
	Index test -	4	51	55	calculated.	
	Total	32	51	83	All of those with missed injuries on CT scan had	

Reference	Tan 2014 ⁴⁷					
					intermedullary T2 hyper intensity consistent with possible central cord syndrome, described as not being unstable.	
	Any cervical sp	oine injury – MRI as inc	lex test			
		Reference standard +	Reference standard -	Total	Note false positives were not possible using CT + MRI as a	
	Index test +	32	NA	32	reference standard meaning specificity could not be	
	Index test -	0	51	51	calculated.	
	Total	32	51	83		
Statistical measures	Any cervical sp	oine injury – CT as inde	ex test: calculated usin	g excel sheet		
	Sensitivity: 76.0					
	Specificity: NA					
	PPV: NA					
	NPV: NA					
	Any cervical spine injury – MRI as index test: calculated using excel sheet					
	Sensitivity: 99.0					
	Specificity: NA					

Reference	Tan 2014 ⁴⁷
	PPV: NA
	NPV: NA
	Unstable cervical spine injury – CT as index test: determined using statements in paper as exact numbers with unstable injuries not provided
	Sensitivity: 100.0
	Specificity: NA
	PPV: NA
	NPV: NA
	Unstable cervical spine injury – MRI as index test: determined using statements in paper and the fact that sensitivity was 100% for any injuries as exact numbers with unstable injuries not provided
	Sensitivity: 100.0
	Specificity: NA
	PPV: NA
	NPV: NA
Source of funding	Stated that there was no funding
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unlikely that index test (for MRI) or reference standard (for CT) were interpreted without knowledge of reference standard/index test, and time interval between index test and reference standard unclear = applies to both index tests

Reference	Tan 2014 ⁴⁷
	Indirectness: serious – all included were obtunded representing a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury
Comments	Not possible to calculate specificity based on the reference standard used. Unstable injuries described but numbers not given to complete 2x2 tables.

Reference	Vanguri 2014 ⁴⁸
Study type	Retrospective
Study methodology	Data source: retrospective review of data from single level 1 trauma centre
	Recruitment: retrospective inclusion from single level 1 trauma centre between January 2008 and December 2012
Number of patients	n = 5676
	(n=5676 identified as matching inclusion criteria and n=5676 analysed – no further details about exclusion reasons)
Patient characteristics	Age, median (range): 39.0 (18-103) years
	Gender (male to female ratio): not reported
	Ethnicity: not reported

Reference	Vanguri 2014 ⁴⁸
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported
	GCS median (range): 15 (3-15)
	Injury Severity Score, median (range): 5 (1-75)
	Duration of stay, median (range): 2 (1-175) days
	Setting: secondary care – trauma centre
	Country: USA
	Inclusion criteria: adults with blunt trauma; meeting triage criteria for trauma team activation (included vital signs such as GCS, obvious anatomic injury and mechanism); and underwent CT of cervical spine
	Exclusion criteria: not reported

Reference	Vanguri 2014 ⁴⁸			
	Adults with blunt trauma and suspected cervical spine injury			
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided			
Index test(s) and	Index test			
reference standard	CT of cervical spine			
	Reference standard Unclear, possibly including other imaging such as MRI and flexion-extension depending on patient Siemens Sensation 16 mm multidetector CT used for all patients. Standard protocol included 2 mm thick axial cuts performed at 2 mm increments with sagittal multiplanar reformatted images. Scan extended from base of skull to level of third thoracic vertebra. Findings suggesting ligamentous injury included increased space between ligamentous columns or other contiguous structures, prevertebral haematoma or oedema and abnormal alignment of vertebra. For MRIs, Siemens Avanto 1.5 used, with protocol including 3 mm thick sagittal cuts and 3 mm thick axial cuts with 0.3 mm standard of error. For all imaging only the final attending radiologist reads of scans were considered to determine pathology. Time between measurement of index test and reference standard: unclear at what time-point reference standard			
Outcome	Cervical spine injury – poorly defined			
2×2 table	Cervical spine injury – CT as index test			
	Reference standard Reference standard Total			

Reference	Vanguri 2014 ⁴⁸				
	Index test +	420	0	420	
	Index test -	0	5256	5256	
	Total	420	5256	5676	
Statistical measures	Cervical spine in	Cervical spine injury – CT as index test: calculated using excel sheet			
	Sensitivity: 1.00				
	Specificity: 1.00				
	PPV: 1.00				
	NPV: 1.00				
Source of funding	Not reported	Not reported			
Limitations	were interpreted	without knowledge of th val between index test	ne other, reference stan	dard poorly descril	x test and reference standard bed and unclear if appropriat it not all patients received the
	-	v serious – head injury i rd included a 2 week fo		ear if all or most ha	ad head injury, and unclear i
Comments	Reference standa	ard poorly described			

Reference	Widder 2004 ⁴⁹
Study type	Prospective
Study methodology	Data source: prospective collection of data from those presenting to a single centre
	Recruitment: 3-year convenience sample used obtained prospectively presenting to single centre between July 1999 and July 2002
Number of patients	n = 102
	(n=113 met inclusion criteria, with n=11 excluded as they have incomplete plain film series; leaving n=102 included in the analysis)
Patient characteristics	Note: characteristics given only for n=113 matching inclusion criteria and not the n=102 analysed
	Age, average: 35.0 years
	Gender (male to female ratio): 77.0% males
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported
	GCS, average: 7.8

Reference	Widder 2004 ⁴⁹
	Injury Severity Score, average: 33.2
	Setting: secondary care – those with severe injuries
	Country: Canada
	Inclusion criteria: ≥16 years; ISS ≥16; GCS <9 or intubated with motor score ≤5 at presentation to trauma centre and at 24 h – high-risk subpopulation of severely injured patients
	Exclusion criteria: diagnosis of cervical cord injury or cervical spine injury at admission; and death within 72 h of arrival at trauma centre
	Adults with trauma and suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if all or most had head injury as no details provided (assumed to have head injury based on severity of injuries – high risk subpopulation of those severely injured)
Index test(s) and	Index test
reference standard	X-ray of cervical spine (data not included further in review as not relevant as initial imaging in severely injured)
	CT of cervical spine

Reference	Widder 2004 ⁴⁹
	Reference standard
	For X-ray: CT used as reference standard
	For CT: final diagnosis at discharge and any readmissions used as reference standard – time-point not mentioned but including readmissions suggests likely >2 weeks (not just limiting to discharge diagnosis)
	Radiological evaluation consisted of three-view (anteroposterior, lateral and odontoid) cervical spine radiographs. If inadequate films, further views (swimmer's) were taken. Helical CT then performed (3 mm cuts) from skull base to T1. Axial images and sagittal and coronal reconstruction reviewed. Concomitantly taken to CT scanner for scans of head and other injured areas. Images obtained using 2.5 mm thickness cuts with 3.75 mm rotations. Blinded radiology review of all plain radiographs and CT images performed separately by two independent radiologists. One radiologist reviewed CT scans and the other reviewed plain films. Blinded reviews performed at least 3 months following admission. Complete plain films adequate if all levels visualised including odontoid and C7-T1.
	Clinical follow-up performed using trauma quality improvement process. All initially admitted to ICU. Once weaned from mechanical ventilation, transferred to one of two acute care services (trauma or neurosurgery). Weekly reviews performed to document missed injuries and complications. All charts reviewed at discharge by trauma service registrars to document all injuries. After discharge, patients referred to trauma clinic and/or brain injury clinic for follow-up. Clinics routinely refer missed injuries back to trauma services for review.
	Time between measurement of index test and reference standard: unclear time interval between tests and follow- up duration for CT reference standard of final diagnosis/readmissions
Outcome	Cervical spine abnormality – poorly defined
2×2 table	Cervical spine abnormality – X-ray as index test (CT as reference standard)

Reference	Widder 2004 ⁴⁹				
		Reference standard +	Reference standard -	Total	
	Index test +	7	2	9	
	Index test -	11	82	93	
	Total	18	84	102	
	Cervical spine a standard)	bnormality – CT as in	dex test (final diagnos	sis at discharge/re	eadmissions as reference
		Reference standard +	Reference standard -	Total	Does not report raw data to calculate specificity or report
	Index test +	18	NR	NR	specificity in the paper.
	Index test -	0	NR	NR	
	Total	18	84	102	
Statistical measures	Cervical spine a PPV and NPV wh		index test (CT as refe	rence standard):	reported in paper (apart from
	calculated using	excel sheet)			
	Sensitivity: 39.0%	, 0			
	Specificity: 98.0%				
	PPV: 78.0%				

Reference	Widder 2004 ⁴⁹
	NPV: 88.0%
	Cervical spine abnormality – CT as index test: reported in paper
	Sensitivity: 100.0%
	Specificity: NR
	PPV: NR
	NPV: NR
Source of funding	Not reported
Limitations	Risk of bias: serious – unclear if consecutive sample enrolled and some concerns about exclusion criteria, and unclear duration between index test and reference standard – applies to both index tests. In addition, for CT as an index test the reference standard may have different in components between patients.
	Indirectness: serious – all included were within a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury
Comments	None
Comments	

D₄2 Children

Reference	AI-Sarheed 2020 ²
Study type	Retrospective cohort study

Reference	AI-Sarheed 2020 ²
Study methodology	Data source: retrieved data from trauma registry (medical records files from January 2005 to March 2015 and hospital's electronic system from April 2015 to July 2018).
	Recruitment: retrospective review of patients between January 2005 and July 2018 at King Abdulaziz Medical City, Saudi Arabia. Level 1 trauma centre serving national guard military staff, employees and their families.
Number of patients	n = 62 (n=65 based on raw data reported)
	(N=62 said to meet inclusion criteria, though this appeared to be n=65 based on raw data reported – no further details about exclusion reasons provided)
Patient characteristics	Age, mean (SD): 8 (3.9) years, range 6 months to 15 years
	Gender (male to female ratio): 48:14 (77.4% male)
	Ethnicity: not reported
	Head injury: unclear if all or most suffered concomitant injury to the head. 17.4% with confirmed intra-axial/extra- axial brain haemorrhage, 5.33% with skull/face laceration, 4.33% with brain oedema, 3.88% with brain injury, 3.39% with brain herniation and 23.30% with skull fracture, of those that had associated injuries.
	Arrived from:
	• Scene, 82.3%

Reference	AI-Sarheed 2020 ²
	Referring hospital, 17.7%
	Mechanism of injury:
	• Fall, 14.5%
	Motor vehicle accident, 59.7%
	Pedestrian, 21.0%
	• Others, 4.8%
	Injury type:
	• Blunt, 98.4%
	Penetrating, 1.6%
	GCS at initial trauma, mean (SD): 7.6 (3.7)
	Injury Severity Score, mean (range): 24.1 (0-68)
	Duration of stay, mean (SD):
	• ICU, 18.2 (36.9) days

Reference	AI-Sarheed 2020 ²
	• Ward, 29.2 (45.8) days
	Setting: arrived at emergency department
	Country: Saudi Arabia
	Inclusion criteria: aged ≤15 years; sustained trauma (motor vehicle accident, fall, struck by falling heavy object, pedestrian, all-terrain vehicle accident and sports injuries); and were intubated at scene or in emergency department.
	Exclusion criteria: patients that were awake or were extubated before clearance.
	Paediatric patients with suspected cervical spine injury. Reports separately for children <8 and ≥8 years, but not relevant for this review protocol.
Target condition(s)	Suspected cervical spine injury – unclear if most/all had head injury (assumed to have head injury based on the severity of injuries – all intubated/unconscious)
Index test(s) and	Index test
reference standard	CT – institution's protocol for clearance of cervical spine in obtunded paediatric trauma patient is to perform CT of cervical spine for all patients. MRI considered if there are any abnormal findings on CT scan, significant mechanism of injury or high clinical suspicion.

Reference	Al-Sarheed 2020 ²				
	Reference standard				
	Radiology/clinical examination, including MRI for some where performed. Cases where CT detected cervical spine injury requiring stabilisation were 'true positives'. Those where CT scan failed to detected cervical spine injury in those that were cervically cleared were classified 'true negatives'. False negatives were those where CT was negative but patients had evidence of cervical spine injuries either clinically or radiologically. False positives were those with abnormal radiological findings but who were cervically cleared by examination/radiology.				
	Time between measurement of index test and reference standard: for those that had MRI, mean (SD) time from CT to MRI was 5.1 (5.7) days (not all had MRI).				
Outcome	Cervical spine injury mandating stabilisation – no further details provided				
2×2 table	CT vs. radiology/clinical examination – cervical spine injury mandating stabilisation				
		Reference standard +	Reference standard	Total	Note: numbers given do not match those said to be
	Index test +	28	0	28	included (n=62)
	Index test -	5	32	37	
	Total	33	32	65	
Statistical measures	<u>CT scan</u>				
	Sensitivity: 84.8% (95% CI 68.1-94.8%) – reported in paper				
	Specificity: 100.0% (95% CI 89.1-100.0%) – reported in paper				
	PPV: 100.0 (calculated using spreadsheet)				

Reference	AI-Sarheed 2020 ²
	NPV: 86.49 (calculated using spreadsheet)
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, reference standard likely not interpreted without knowledge of index test and index test may have affected extent of testing/examination used as reference standard, not all had the same reference standard and unclear duration between index test and final diagnosis used as reference standard
	spine injury, and unclear if the reference standard matches protocol as definition provided is limited to 'radiology/clinical examination'
Comments	None
Reference	Brockmeyer 2012⁵

Study type	Prospective
Study methodology	Data source: prospective study from single centre
	Recruitment: prospective enrolment of patients in continuous fashion between November 2005 and September 2009
Number of patients	n = 24
	(n=24 enrolled in the study and no details about exclusion reasons provided)

Reference	Brockmeyer 2012⁵
Patient characteristics	Age, range: 4 months to 16 years
	Gender (male to female ratio): 66.7% males
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury:
	 Auto/pedestrian, n=2 Fall, n=3 Skiiing, n=1 Scooter, n=1 Kicked by a horse, n=1 Snowmobile, n=1 Non-accidental trauma, n=6 All terrain vehicle, n=3 Motor vehicle accident, n=4 Auto/bicycle, n=1 Motorcycle, n=1
	GCS at initial trauma, mean (SD): not reported

Reference	Brockmeyer 2012⁵
	Injury Severity Score, mean (SD): not reported
	Setting: secondary care – those with severe injuries admitted to ICU
	Country: USA
	Inclusion criteria: GCS ≤8 after haemodynamic stabilisation; admitted to ICU; >2 weeks old and <17 years; and suspected or known traumatic cervical spine injury
	Exclusion criteria: inability to obtain plain radiographs, CT or MR imaging within 7 days of admission (later amended to 10 days); inability to obtain follow-up plain cervical spine radiographs 3-4 months after injury; and isolated gunshot or penetrating wound to head with little chance of cervical spine injury.
	Children with severe injury admitted to ICU with suspected cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if most/all had head injury (assumed to have head injury based on severity of injuries – all severely injured and admitted to ICU)
Index test(s) and	Index test
reference standard	X-ray of cervical spine
	CT of cervical spine

Reference	Brockmeyer 201	Brockmeyer 2012 ⁵			
	MRI of cervical s	MRI of cervical spine			
	Reference standa	ard			
	Clinical outcome/diagnosis of early instability – undergoing surgical correction. Unclear how confirmed but possibly mixture of all available data. Follow-up possibly >2 weeks as mentions plain radiographs at follow-u 3-4 months post-injury.				
	For each patient a plain lateral cervical spine radiograph, complete cervical spine CT with 2D sagittal and coronal reconstructions, cervical spine MRI imaging with T1 and T2 weighted and short tau inversion recovery images and cervical spine flexion/extension films with fluoroscopy were acquired. Follow-up radiographic imaging consisted of plain lateral flexion-extension radiographs at 3-4 months post-injury. All images obtained using passive motion in an awake patient and read by paediatric neuroradiologist.				u inversion recovery images up radiographic imaging All images obtained using
Outcome	Early cervical spine instability – required surgical correction				
2×2 table	Early cervical spine instability (surgical correction) – X-ray (plain radiography) as index test				
		Reference standard +	Reference standard	Total	
	Index test +	1	1	2	
	Index test -	0	22	22	

Reference	Brockmeyer 2012⁵					
	Total	1	23	24		
	Early cervical spine instability (surgical correction) – CT as index test					
		Reference standard +	Reference standard	Total		
	Index test +	1	0	1		
	Index test -	0	23	23		
	Total	1	23	24		
	Early cervical spine instability (surgical correction) – MRI as index test					
		Reference standard +	Reference standard -	Total		
	Index test +	1	6	7		
	Index test -	0	17	17		
	Total	1	23	24		
Statistical measures					as) as index test: calculated aw data throughout study	
	Sensitivity: 100.0	%				

Reference	Brockmeyer 2012 ⁵
	Specificity: 96.0%
	PPV: 50.0%
	NPV: 100.0%
	Early cervical spine instability (surgical correction) – CT as index test: calculated using excel sheet as numbers in paper don't quite match those calculated using raw data throughout study
	Sensitivity: 100.0%
	Specificity: 100.0%
	PPV: 100.0%
	NPV: 100.0%
	Early cervical spine instability (surgical correction) – MRI as index test: calculated using excel sheet as numbers in paper don't quite match those calculated using raw data throughout study
	Sensitivity: 100.0%
	Specificity: 74.0%
	PPV: 14.0%
	NPV: 100.0%
Source of funding	Technical fees for MR imaging portion of study funded by a grant from Primary Children's Foundation.

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Reference	Brockmeyer 2012⁵
Limitations	Risk of bias: very serious – unclear if index tests were interpreted without knowledge of other tests and whether reference standard interpreted without knowledge of index test, unclear time interval between index test and reference standard and unclear if reference standard components were the same for all patients – applies to all three index tests
	Indirectness: serious – all included were at within a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury
Comments	None

Reference	Derderian 2019 ¹²
Study type	Retrospective study
Study methodology	Data source: hospital trauma registry (from children's hospital with level 1 trauma centre) used to identify those matching inclusioncriteria. Patients with traumatic brain, spine or multiorgan injury identified and radiology records subsequently reviewed to narrow itdown to those that underwent both CT and MRI following injury. Electronic medical records also queried to obtain a list of all childrenthat underwent MRI of the cervical spine, which were cross-referenced with those from the trauma registry.
	Recruitment: those matching inclusion criteria between January 2001 and November 2015.
Number of patients	n = 221
	(n=222 trauma patients had both CT and MRI to evaluate the cervical spine, but n=1 was excluded due to major upper thoracic spinal cord transection with extension into the cervical region; leaving n=221 included in the analysis)

Reference	Derderian 2019 ¹²
Patient characteristics	Age, median (IQR): 9 (3-14) years
	Gender (male to female ratio): 64.7% male and 35.3% female
	Ethnicity: not reported
	Head injury: unclear if all or most had concomitant head injury as part of the injury mechanism – 15.8% said to have isolated headinjury, with multiorgan injury including 66.5% which may include head injury
	Mechanism of injury:
	Motor vehicle accident, 30.8%
	Non-accidental trauma, 10.9%
	• Fall, 10.0%
	Ski-ing/snowboarding/sledding, 5.4%
	Other sports-related, 17.6%
	• Other, 8.6%
	Type of traumatic injury:

Reference	Derderian 2019 ¹²
	 Isolated head injury, 15.8%
	 Isolated spine injury, 17.6%
	• Multiorgan, 66.5%
	Treatment of C-spine:
	Cleared prior to or during hospitalisation, 68.8%
	Collar at time of discharge, 15.8%
	Deceased prior to clearance, 0.4%
	• Halo, 1.4%
	 Fusion surgery (with or without halo), 13.6%
	Intensive care unit admission: 73.8%
	Intensive care unit length of stay, median (IQR): 8 (3-13) days
	Lisewitz Lisewitz of start marking (LOD), 45 (4.20) date
	Hospital length of stay, median (IQR): 15 (4-36) days
	GCS score median (IQR): 11 (5-15)

Reference	Derderian 2019 ¹²
	Setting: secondary care – trauma patients at children's hospital
	Country: USA
	Inclusion criteria: children that received CT and MRI of cervical spine following trauma
	Exclusion criteria: one child excluded with a major upper thoracic spinal cord transection with extension into the cervical region.
	Paediatric trauma patients undergoing cervical spine CT and MRI scan
Target condition(s)	Suspected cervical spine injury – unclear if most/all had head injury
Index test(s) and	Index test
reference standard	CT – stable or unstable injury on CT used as CT-positive
	Or
	MRI – stable or unstable injury on MRI used as MRI-positive

Reference	Derderian 2019 ¹²
	Reference standard
	Study not specifically reported as a diagnostic accuracy study. Calculated accuracy from data provided for CT and MRI alone in predicting clinical instability (defined as undergoing intervention by halo placement or spinal fusion). Unclear follow-up period covered by the data.
	Time between measurement of index test and reference standard: unclear time period between CT and MRI, unclear follow-up period in terms of intervention being performed.
	Institutional protocol was as follows: cervical spine imaging in those with risk factors for cervical spine injury. Those that could cooperate with a clinical examination also had a supplemental 3-view radiograph if >5 years. For those ≤5 years, odontoid view replaced by right and left oblique cervical radiography. CT reserved for those with risk factors who were unable to participate in a clinical examination or had significant distracting injury. If no CT findings of unstable spinal column or a spinal cord injury and the suspicion for one was low, MRI was deferred until the patient demonstrated improved clinical status sufficient for extubation and thorough neurological examination. If this did not occur within 72 h then the MRI was obtained and collar removed if no unstable cervical spine injury observed.
	Abnormal imaging findings included fracture, translation, angulation, herniated nucleus pulposus, intraspinal haematoma or ligamentous injury. Ligamentous injury considered column-disrupting if normal linear T2 hypointense structure of the anterior longitudinal ligament, posterior longitudinal ligament or ligamentum flavum was interrupted to suggest a complete ligamentous tear. Isolated oedema in interspinous ligaments not considered column-disrupting injury. If any of these abnormalities were not associated with disruption of two or more spinal columnts then they were considered stable.
	Imaging results for CT and MRI categorised into one of three groups: no injury, stable injury and unstable injury.

Reference	Derderian 2019 ¹²				
Outcome	Clinical instability – those undergoing a surgical intervention (spinal fusion or halo placement) were defined as clinically unstable in the study.				
2×2 table	CT vs. reference standard (clinical instability – surgical spinal fusion or halo placement)				
		Reference standard +	Reference standard -	Total	Note that these tables and subsequent accuracy results
	Index test +	33	28	61	provided under 'statistical measures' were calculated
	Index test -	0	160	160	from data provided in the paper.
	Total	33	188	221	L - L
	MRI vs. reference standard (clinical instability – surgical spinal fusion or halo placement)				
		Reference standard +	Reference standard –	Total	Note that these tables and subsequent accuracy results provided under 'statistical measures' were calculated
	Index test +	33	104	137	
	Index test -	0	84	84	from data provided in the paper.
	Total	33	188	221	μαροι.
Statistical measures	CT vs. reference data in paper Sensitivity: 1.00	e standard (clinical ins	stability – surgical spi	nal fusion or hal	o placement) – calculated from
	Specificity: 0.85				

Reference	Derderian 2019 ¹²
	PPV: 0.54
	NPV: 1.00
	MRI vs. reference standard (clinical instability – surgical spinal fusion or halo placement) – calculated from data in paper
	Sensitivity: 1.00
	Specificity: 0.45
	PPV: 0.24
	NPV: 1.00
Source of funding	Reported to be no funding
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, likely that results of index tests affected decisions about considering intervention (reference standard) and unclear duration of follow-up for intervention and whether follow-up was similar for all patients – applies to both index tests
	Indirectness: serious – unclear if all of the majority also sustained a head injury
Comments	Not formally described as a diagnostic accuracy study and no sensitivity etc. reported, but data available to calculate sensitivity and specificity for clinically unstable injuries.

Reference	Henry 2013-2 ²⁵
Study type	Retrospective study
Study methodology	Data source: retrospective review of patients treated for cervical spine injury using paediatric trauma database at Tufts Medical Center (level 1 paediatric trauma centre)
	Recruitment: retrospective review of database.
Number of patients	n = 84
	(n=84 said to match inclusion criteria and n=84 analysed – no details about exclusion reasons)
Patient characteristics	Age, mean (SD): 9.0 (5.8) years
	Gender (male to female ratio): 56% male and 44% female Ethnicity: not reported Head injury: no mention of head injury, unclear if all or majority had concomitant head injury as part of the injury mechanism.
	Mechanism of injury:
	Sports/physical activity, 28%
	Motor vehicle accident, 32%

Reference	Henry 2013-2 ²⁵
	Domestic violence, 6%
	• Fall, 29%
	Self-inflicted, 1%
	Non-specific/unclear, 4%
	GCS, mean (range): 12.8 (3-15)
	Setting: secondary care – trauma centre
	Country: USA
	Inclusion criteria: aged ≤18 years at time of injury; involved in a trauma; and had CT and MRI scans of the cervical spine performed
	within 48 h of injury.
	Exclusion criteria: not reported.
	Paediatric patients evaluated for cervical spine injury.

Reference	Henry 2013-2 ²⁵				
Target condition(s)	Suspected cervical spine injury – unclear if most/all had head injury				
Index test(s) and	Index test and ret	ference standard			
reference standard					
	CT and MRI were both an index test as well as a reference standard, depending on the type of injury. CT was used as the reference standard for osseous injury (bony injuries – fractures, locked facets, subluxations and dislocations) and MRI was used as the reference standard for soft tissue injuries (compression fractures, soft tissue oedema, ligamentous injury, muscular injury and spinal cord injury). Time between measurement of index test and reference standard: mean (range) time between CT and MRI was 0.4 (0-2) days. 80/84 had CT prior to MRI scan, while 4 had MRI prior to CT scan. Time between injury and CT				
	, , ,	s and between injury a			
Outcome	Osseous	injury (reference standa	rd as CT): fractures, lo	cked facets, sublux	ations and dislocations
		 Soft tissue injury (reference standard as MRI): compression fractures, soft tissue oedema, ligamentous injury, muscular injury and spinal cord injury 			
2×2 table	MRI vs. CT (refe	rence standard) for os	sseous injury		
		Reference standard +	Reference standard -	Total	
Index test + 6 2 8					
	Total	6	78	84	

Reference	Henry 2013-2 ²⁵				
	CT vs. MRI (reference standard) for soft tissue injury				
		Reference standard +	Reference standard	Total	
	Index test +	3	0	3	
	Index test -	10	71	81	
	Total	13	71	84	
Statistical measures		MRI vs. CT (reference standard) for osseous injury – values reported in paper			
	Sensitivity: 100% Specificity: 97%				
	PPV: 75% NPV: 100%				
	CT vs. MRI (reference standard) for soft tissue injury – values reported in paper				
	Sensitivity: 23%				
	Specificity: 100%	Specificity: 100%			
	PPV: 100%				
	NPV: 88%				
Source of funding	Not reported				

Reference	Henry 2013-2 ²⁵
Limitations	Risk of bias: very serious - unclear if consecutive sample enrolled and depending on the type of injury detected (osseousor soft tissue), index or reference standard was likely interpreted with knowledge of the other as CT performed first in most cases – applies to both index tests Indirectness: very serious - unclear if all or the majority also sustained a head injury and outcome limited to fractures or ligamentous injury depending on the index test
Comments	None

Reference	Henry 2013-1 ²⁶
Study type	Retrospective study
Study methodology	Data source: retrospective review of patients treated for cervical spine injury using paediatric trauma database at Tufts Medical Center (level 1 paediatric trauma centre)
	Recruitment: those between 2002 and 2011 in the database and matching inclusion criteria were included
Number of patients	n = 73
	(n=146 meeting inclusion criteria identified, with n=12 excluded due to lack of information about cervical spine clearance in medical charts, n=23 prescribed a rigid collar and cleared at follow-up and n=38 without follow-up information on record excluded; leaving n=73 included in the analysis)
Patient characteristics	Age, mean (SD): 8.3 (5.8) years

Reference	Henry 2013-1 ²⁶
	Gender (male to female ratio): 65% male and 35% female
	Ethnicity: not reported
	Head injury: no mention of head injury, unclear if all or majority had concomitant head injury as part of the injury mechanism.
	Mechanism of injury:
	Sports/physical activity, 22%
	Motor vehicle accident, 40%
	Domestic violence, 3%
	• Fall, 28%
	Self-inflicted, 0%
	Non-specific/unclear, 7%
	Indications for imaging:
	Neck pain, 12%
	Neurological deficit, 3%

Reference	Henry 2013-1 ²⁶
	Neurological symptom, 8%
	Distracting injuries, 44%
	Sedation/intubation, 16.5%
	• Pain + ≥1 other factor, 16.5%
	GCS at admission, mean (SD): 12.1 (5.0)
	Duration of stay, mean: 4.6 days (range, 0-28 days)
	Setting: secondary care – paediatric trauma centre
	Country: USA
	Inclusion criteria: aged ≤18 years at the time of injury; could not be cleared by means of clinical criteria; and underwent MRI of the cervical spine with a STIR (short T1 inversion recovery) sequence within 48 h of injury.
	Exclusion criteria: excluded a group that required a hard collar and clearance at follow-up, focusing on those that were cleared prior to discharge (may affect diagnostic accuracy results?); and lack of information about cervical spine clearance in medical charts.

Reference	Henry 2013-1 ²⁶
	Paediatric patients being assessed for cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if most/all had head injury
Index test(s) and	Index test
reference standard	MRI with STIR sequence
	Reference standard
	Follow-up or flexion/extension radiographs – injury requiring surgical intervention or presenting with clinical (significant pain or neurological compromise) or radiographic evidence of instability upon follow-up. Flexion-extension radiographs used to identify false positive findings on MRI. Mean follow-up 10.0 (18.4) months, range 4 days – 7.6 years.
	MRI diagnosis of cervical injury requiring surgical stabilisation considered true positives and cases where MRI findings were negative and patients cleared as in-patients with follow-up information on record were true negatives (no surgical intervention and no instability or pain at follow-up). False negatives defined as cases where patient displayed clinical (significant pain or neurological compromise) or radiographic evidence of instability during follow-up with an initially negative MRI. False positive defined as case where MRI showed abnormal findings but patient was cleared by flexion-extension radiographs during admission (only used if patient had reasonable range of cervical motion).
	Time between measurement of index test and reference standard: unclear time between MRI and flexion- extension where this was used, mean follow-up of 10.0 months in terms of cases where follow-up was used as reference standard.

Reference	Henry 2013-1 ²⁶					
Outcome	Cervical spine injury – instability: requiring surgical stabilisation – either undergoing it or demonstrating signs of instability, pain or neurological compromise during follow-up).					
2×2 table	MRI STIR vs. ref	MRI STIR vs. reference standard (follow-up with/without flexion-extension radiographs during admission)				
		Reference standard +	Reference standard -	Total		
	Index test +	1	2	3		
	Index test -	0	70	70		
	Total	1	72	73		
Statistical measures		MRI STIR vs. reference standard (follow-up with/without flexion-extension radiographs during admission) – reported in paper				
	Sensitivity: 100%					
	Specificity: 97%					
	PPV: 33%					
	NPV: 100%	NPV: 100%				
Source of funding	Not reported					
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, excluded a group that were discharged in a hard collar and cleared later, unlikely that reference standard was interpreted without knowledge of index test results and reference standard varied between patients					

Reference Henry 2013-1²⁶

IVEIGLEHICE	116111 y 2013-1
	Indirectness: serious – unclear if all or the majority also sustained a head injury
Comments	None

Reference	Qualls 2015 ³⁸	
Study type	Retrospective cohort study	
Study methodology	Data source: trauma registry at St. Louis Children's Hospital (academic children's hospital with level 1 trauma certification) queried to identify all patients matching inclusion criteria.	
	Recruitment: those admitted between 1 st January 2002 and 31 st December 2012.	
Number of patients	n = 63	
	(n=64 met inclusion criteria, with n=1 excluded due to a previous cervical spine injury; leaving n=63 included in the analysis)	
Patient characteristics	Age, median (range): 9.6 (0.1-17.8) years	
	Gender (male to female ratio): 52.4% male and 47.6% female	
	Ethnicity: not reported	
	Head injury: all had severe traumatic brain injury to be included	

Reference	Qualls 2015 ³⁸
	Mechanism of injury:
	Motor vehicle occupant, 47.6%
	Non-accidental trauma, 20.6%
	Other motorised transport crash, 14.3%
	Pedestrian vs. automobile, 14.3%
	Fall from elevation, 3.2%
	GCS, median (range):
	 In ED, 5 (3-13)
	At neurosurgical consult, 6 (3-13)
	At admission to paediatric intensive care unit, 6 (3-14)
	Injury Severity Score, median (range): 30 (11-75)
	Paediatric trauma score, median (range): 3 (-3 to 10)
	Duration of stay, median (range): 34 (5-129) days

Reference	Qualls 2015 ³⁸
	Duration of paediatric intensive care unit stay, median (range): 13 (2-34) days
	Intubated:
	• In ED, 90.5%
	At neurosurgical consult, 96.8%
	At admission to paediatric intensive care unit, 98.4%
	Setting: secondary care – trauma injuries admitted to hospital
	Country: USA
	Inclusion criteria: severe traumatic brain injury admitted to hospital; and received cervical spine MRI and cervical spine CT.
	Exclusion criteria: those receiving cervical spine MRI but that had a history of previous cervical spine injury; GCS score >8 at admission to ED, initial neurosurgery evaluation and admission to paediatric intensive care unit (all three time-points) were excluded.

Reference	Qualls 2015 ³⁸
	Children with severe traumatic brain injury and assessed for cervical spine injury
Target condition(s)	Suspected cervical spine injury in children with severe traumatic brain injury
Index test(s) and	Index test
reference standard	CT alone
	MRI alone
	Reference standard
	CT followed by MRI (CT + MRI)
	Presence of injury on CT or MRI determined by review of all imaging reports by a paediatric neurosurgeon and a paediatric emergency physician. They determined whether patients had an injury, whether it was unstable and which imaging modalities were able to detect the injuries seen. Institutional protocol did not require plain radiography of the cervical spine but where this had been performed the imaging was evaluated in the same way as CT and MRI and determined to demonstrate evidence of injury or no evidence of injury.
	Time between measurement of index test and reference standard: reference standard was the process of using both rather than only a single imaging test. Institutional protocol was to perform MRI if no improvement in mental status or intubation >72 h.
	Institutional protocol was as follows for patients with significant altered mental status or intubation following blunt trauma: axial cervical spine CT for all children immediately on presentation. Cervical spine precautions continued for those with normal CT. If mental status and intubation normalised within 72 h, children were cleared clinically.

Reference	Qualls 2015 ³⁸				
	Otherwise, MRI obtained if persisted >72 h. If MRI was normal, patients were cleared and cervical spine precautions discontinued.				
Outcome	operative obtained	 Unstable cervical spine injury – injury resulting in neurological deficit localised to cervical spinal cord, operative stabilisation, halo placement or cervical immobilisation of 3 months of greater (duration obtained from patient records including follow-up appointments by neurosurgery service). Any cervical spine injury, including those that were not considered unstable as well as unstable injuries. 			
2×2 table	Unstable cervic	al spine injury – CT al	one vs. CT followed b	y MRI (reference	standard)
		Reference standard +	Reference standard	Total	Raw data calculated from diagnostic accuracy measures provided in paper.
	Index test +	5	9	14	
	Index test -	0	49	49	
	Total	5	58	63	
	Unstable cervical spine injury – MRI alone vs. CT followed by MRI (reference standard)				
		Reference standard +	Reference standard	Total	Raw data calculated from diagnostic accuracy
	Index test +	4	11	15	measures provided in paper.
	Index test -	1	47	48	
	Total	5	58	63	

Reference	Qualls 2015 ³⁸		
	Any cervical spine injury – CT alone vs. CT followed by MRI (reference standard)		
	Not possible to calculate raw data from accuracy measures provided as only sensitivity and NPV were reported.		
	Any cervical spine injury – MRI alone vs. CT followed by MRI (reference standard)		
	Not possible to calculate raw data from accuracy measures provided as only sensitivity and NPV were reported.		
Statistical measures	Unstable cervical spine injury – CT alone vs. CT followed by MRI (reference standard) – as reported in paper		
	Sensitivity: 100% (95% CI 48-100%)		
	Specificity: 84.5% (95% CI 73-93%)		
	PPV: 35.7% (95% CI 13-65%)		
	NPV: 100% (95% CI 93-100%)		
	Unstable cervical spine injury – MRI alone vs. CT followed by MRI (reference standard) – as reported in paper		
	Sensitivity: 80% (95% CI 29% to 97%)		
	Specificity: 81% (95% CI 79-90%)		
	PPV: 26.7% (95% CI 8-55%)		
	NPV: 98% (95% CI 89% to 100%)		

Reference	Qualls 2015 ³⁸
	Any cervical spine injury – CT alone vs. CT followed by MRI (reference standard) – as reported in paper
	Sensitivity: 63.2% (95% CI 38-84%)
	Specificity: NR
	PPV: NR
	NPV: 86.3% (95% CI 74-94%)
	Any cervical spine injury – MRI alone vs. CT followed by MRI (reference standard) – as reported in paper
	Sensitivity: 68.4% (95% CI 43-87%)
	Specificity: NR
	PPV: NR
	NPV: 88% (95% CI 76-95%)
Source of funding	No funding was received for the study
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, either index test (for MRI alone) or reference standard (for CT alone) unlikely to have been interpreted without knowledge of the other and appears to be 72 h between CT and MRI being performed – applies to both index tests
	Indirectness: serious – all included were within a more severely injured subgroup which may be less applicable to general population of those attending ED with suspected cervical spine injury
Comments	2x2 data not reported so calculated from the diagnostic accuracy measures reported (sensitivity etc.)

Reference	Rana 2009 ³⁹
Study type	Retrospective
Study methodology	Data source: retrospective review of data from single children's hospital which has level 1 paediatric trauma centre status
	Recruitment: retrospective review of children from single level 1 trauma centre between 2004 and 2006
Number of patients	n = 54
	(n=318 had imaging for cervical spine injury, with n=54 with both CT and plain films included in the analysis)
Patient characteristics	Note: characteristics reported only for n=318 patients identified not specifically those analysed with both CT and MRI (n=54)
	Age, mean: 10.2 years
	Gender (male to female ratio): 64.0% males
	Ethnicity: not reported
	Head injury: no details reported, unclear if all or most suffered concomitant injury to the head
	Mechanism of injury: not reported

Reference	Rana 2009 ³⁹
	GCS at initial trauma, mean: 13
	Injury Severity Score, mean: 14.2
	Intubated, 24.0%
	Setting: secondary care – those with trauma at children's hospital
	Country: USA
	Inclusion criteria: paediatric trauma patients (<18 years); and cervical spine imaging and/or a confirmed cervical spine injury
	Exclusion criteria: those without cervical spine imaging or a cervical spine injury.
	Children sustaining trauma and with suspected or confirmed cervical spine injury
Target condition(s)	Suspected cervical spine injury – unclear if most/all had head injury
Index test(s) and	Index test
reference standard	X-ray of cervical spine

Reference	Rana 2009 ³⁹
	CT of cervical spine
	<u>Reference standard</u> CT used as 'gold standard' for X-ray, but clinical outcome used as gold standard for CT, which includes subsequent imaging where performed. Follow-up unclear for reference standard of clinical outcome.
	Institutional protocol included initial physical examination of cervical spine. Those with reliable exams and who were fully aware without motor/sensory deficits, neck pain, evidence of intoxicating agents and distracting injuries were cleared clinically in the trauma centre. No further evaluation of cervical spine was required (excluded from study). If there were complaints of neck tenderness, neurologic deficits, abnormal GCS or distracting pain from another imaging, cervical spine imaging was performed (plain radiographs, CT or both at discretion of trauma team leader). If radiographs negative for injury, initial stabilisation collar changed to padded collar until reliable examination performed. Flexion and extension views performed in those with continued cervical tenderness or if prolonged intubation required. If pain persisted, discharged home with cervical spine collar and followed up by neurosurgery team for clearance.
	Time between measurement of index test and reference standard: duration between index tests and confirmed diagnosis unclear.
Outcome	Cervical spine injury – poorly defined
2×2 table	Raw data reported difficult to follow in paper (unclear how many analysed for X-ray) and attempts to work out do not match sensitivity/specificity values reported in paper. Values reported in paper therefore used and 2x2 tables not completed.

Reference	Rana 2009 ³⁹
Statistical measures	Cervical spine injury – X-ray as index test: reported in paper
	Sensitivity: 61.5%
	Specificity: 1.6%
	PPV: 61.5%
	NPV: NR
	Cervical spine injury – CT as index test: reported in paper
	Sensitivity: 100.0%
	Specificity: 97.6%
	PPV: 79.4%
	NPV: NR
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if reference standard interpreted without knowledge of index test and unclear interval between index test and reference standard – applies to both index tests. In addition, for CT as an index test, it is possible that the reference standard components differed between patients
	Indirectness:
	X-ray as index test – serious - unclear if all or the majority also sustained a head injury

Reference	Rana 2009 ³⁹
	CT as index test – very serious - unclear if all or the majority also sustained a head injury, and unclear if follow- up of at least 2 weeks as part of the reference standard
Comments	None

Reference	Somppi 2018 ⁴³
Study type	Retrospective cohort study
Study methodology	Data source: those matching inclusion criteria and presenting to urban tertiary care centre (designated ACS level 1 paediatric trauma centre). Suspected cervical spine injury identified from radiology database using ICD codes 70490, 70492 and 87.22 of ninth revision. These correspond to cervical CT without contrast, soft tissue cervical CT with and without contrast, and other X-ray of cervical spine. Also reviewed hospital trauma registry for study period to identify those with neck injuries that may have been missed by automated search.
Number of patients	n = 574 (n=671 in the base dataset, with n=27 with no cervical spine injury suspected, n=10 with no consent for retrospective studies, n=49 with CT imaging not performed at the institution and n=11 with unavailable records/not admitted to ER excluded; leaving n=574 included in the analysis)
Patient characteristics	Age, median (IQR): 9.70 (4.78-13.83) years

Reference	Somppi 2018 ⁴³
	Gender (male to female ratio): 57.5% male and 42.5% female
	Ethnicity: not reported
	Head injury: unclear proportion that also sustained a head injury, n=230 (40%) underwent head CT
	Disposition:
	• Discharged, 78.6%
	Admitted, 21.4%
	Mechanism of injury:
	• Fall, 50.0%
	• Sports, 22.2%
	Motor vehicle accident, 12.7%
	Other/unknown, 15.0%
	GCS: not reported

Reference	Somppi 2018 ⁴³
	Cervical spine imaging performed:
	• X-ray, 86.6%
	• CT, 47.9%
	• MRI, 4.9%
	Single imaging study, 51.7%
	Two imaging studies, 40.9%
	Three imaging studies, 6.6%
	• Four or more imaging studies, 0.5%
	Setting: secondary care
	Country: USA
	Inclusion criteria: children and adolescents (aged ≤19 years); presenting with possible neck injury to urban tertiary care centre
	Exclusion criteria: underwent CT imaging as part of a diagnostic procedure (i.e. for abscess drainage or interventional radiology); patients receiving CT imaging before transfer to the trauma centre the study was performed at; no cervical spine injury suspected (e.g. imaging performed for fever with neck pain and no related trauma); no consent for retrospective studies; and record unavailable or not admitted to the emergency room.

Reference	Somppi 2018 ⁴³
	Children and adolescents presenting with possible neck injury
Target condition(s)	Suspected cervical spine injury – unclear if most/all had head injury (40% had head CT)
Index test(s) and	Index test
reference standard	X-ray (n=495)
	CT (n=130)
	MRI (n=21)
	Reference standard
	Unclear, possibly all imaging and follow-up? To ensure complete identification of spinal cord injuries, medical records for all patients with a negative imaging study (CT or MRI) were reviewed for up to 1 month after the index
	ED visit to assess for cervical spine pain on ED or outpatient visits to the institution.
	Time between measurement of index test and reference standard: time between different types of imaging unclear, up to 1 month follow-up for those with negative imaging CT or MRI.
Outcome	Cervical spine injury – ligamentous or osseous injury documented by attending radiologist in their report. Patients
	with spinal cord injury without radiograph evidence were defined as those with MRI abnormalities indicating spinal cord injury but with a negative X-ray or CT OR any patient re-presenting after initial index ED visit with

Reference	Somppi 2018 ⁴³
	persistent cervical spine pain or neurologic abnormalities (e.g., tingling, numbness) that then underwent MRI which revealed ligamentous or osseous injuries.
2×2 table	Raw data incompletely reported and could not calculate raw data from diagnostic accuracy measures provided, meaning 2x2 tables could not be completed. A total of 8 patients had confirmed cervical spine injury.
Statistical measures	X-ray vs. reference standard (imaging/follow-up?) – reported in study (n=495)
	Sensitivity: 0.83 (95% CI 0.36-0.99)
	Specificity: 0.97 (95% CI 0.96-0.99)
	PPV: 0.31 (95% CI 0.12-0.59)
	NPV: 0.99 (95% CI 0.98-0.99)
	CT vs. reference standard (imaging/follow-up?) – reported in study (n=130)
	Sensitivity: 1.00 (95% CI 0.52-1.00)
	Specificity: 1.00 (95% CI 0.96-1.00)
	PPV: 1.00 (95% CI 0.52-1.00)
	NPV: 1.00 (95% CI 0.96-1.00)
	MRI vs. reference standard (imaging/follow-up?) – reported in study (n=21)
	Sensitivity: 1.00 (95% CI 0.51-1.00)
	Specificity: 1.00 (95% CI 0.75-1.00)

Reference	Somppi 2018 ⁴³
	PPV: 1.00 (95% CI 0.52-1.00)
	NPV: 1.00 (95% CI 0.75-1.00)
Source of funding	Not reported
Limitations	Risk of bias: very serious – unclear if consecutive sample enrolled, unclear if index tests all interpreted without knowledge of reference standard, reference standard used for each index test unclear as is whether reference standard was interpreted without knowledge of index test and unclear time interval between index test and reference standard and whether all received the same reference standard – applies to all three index tests
	Indirectness: very serious – unclear if all or the majority also sustained a head injury (40% said to have had head CT) and reference standard poorly defined so unclear if matches protocol.
Comments	2x2 data not reported. Attempted to calculate based on diagnostic accuracy measures provided (sensitivity etc.), but numbers were not consistent with the number analysed.

58 Appendix E – Forest plots

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Ead Coupled sensitivity and specificity forest plots

- E.6.1 Adults all having index test and not limited to those that were admitted
- E.1821 X-ray as index test

Figure 2: Those with blunt trauma with all having head CT, CT as reference standard, cervical spine fracture as outcome

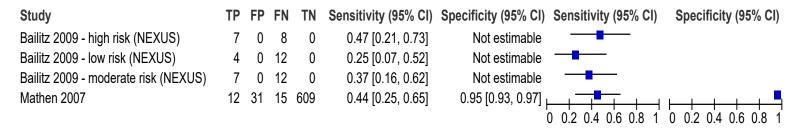
63

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Figure 3: Those with trauma and low risk (one NEXUS criterion), unclear if head injury, CT as reference standard, clinically significant cervical spine injury as outcome

Study	ΤР	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Gharekhanloo 2021	4	6	6	204	0.40 [0.12, 0.74]		0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1

Figure 4: Those meeting at least one NEXUS criterion, unclear if head injury, final diagnosis as reference standard, clinically significant injury as outcome



Note that for the three Bailitz 2009 analyses, FP and TN were not reported and values were not necessarily zero. Only sensitivity could be calculated for these analyses.

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Figure 5: Any following trauma, unclear if head injury, CT as reference standard, cervical spine fractures as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Lee 2001	12	0	24	0	0.33 [0.19, 0.51]	Not estimable		
Takami 2014	10	0	6	0	0.63 [0.35, 0.85]			

Note that for both studies, FP and TN were not reported and values were not necessarily zero. Only sensitivity could be calculated for these analyses.

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Figure 6: Any with blunt trauma, unclear if head injury, MRI as reference standard, ligamentous cervical spine injury as outcome

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Figure 7: Any with blunt injury and cervical spine assessed, unclear if head injury, reference standard unclear/final diagnosis, cervical spine injuries as outcome

Study TP FP FN TN Sensitivity (95% CI) Specificity (95% CI) Sensitivity	y (95% CI) Specificity (95% CI)
	0.6 0.8 1 0 0.2 0.4 0.6 0.8 1

Note that for this study, FP and TN were not reported and values were not necessarily zero. Only sensitivity could be calculated for this analysis.

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Figure 8: Any with blunt injury and cervical spine assessment, unclear if head injury, reference standard unclear/final diagnosis, cervical spine fractures as outcome

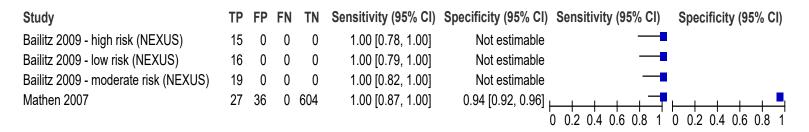
Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Nguyen 2005 - high risk	14	1	1	18	0.93 [0.68, 1.00]	0.95 [0.74, 1.00]		
Nguyen 2005 - Iow risk	0	0	0	78	Not estimable			0 0.2 0.4 0.6 0.8 1

Note that for the low-risk analysis, sensitivity could not be calculated as there were no references standard positives in this group.

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E.1.752 CT as index test

Figure 9: Those meeting at least one NEXUS criterion, unclear if head injury, final diagnosis as reference standard, clinically significant injury as outcome



Note that for the three Bailitz 2009 analyses, FP and TN were not reported and values were not necessarily zero. Only sensitivity could be calculated for these analyses.

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Figure 10: Those failing NEXUS low risk criteria, unclear if head injury, final diagnosis as reference standard, clinically significant fractures as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Inaba 2016	195	907	3	9171	0.98 [0.96, 1.00]			0 0.2 0.4 0.6 0.8 1

Figure 11: Any with blunt injury and cervical spine assessed, unclear if head injury, reference standard unclear/final diagnosis, cervical spine injuries as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Duane 2016	561	6	0	8660	1.00 [0.99, 1.00]	1.00 [1.00, 1.00]	•	•
Griffen 2003	116	0	0	0	1.00 [0.97, 1.00]	Not estimable	•	l
Vanguri 2014	420	0	0	5256	1.00 [0.99, 1.00]	1		0 0.2 0.4 0.6 0.8 1

Note that for the Griffen 2003 study, FP and TN were not reported and values were not necessarily zero. Only sensitivity could be calculated for this analysis.

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81

Figure 12: Any with blunt injury and cervical spine assessment, unclear if head injury, reference standard unclear/final diagnosis, cervical spine fractures as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Nguyen 2005 - high risk	15	0	0	19	1.00 [0.78, 1.00]	1.00 [0.82, 1.00]		
Nguyen 2005 - Iow risk	0	0	0	78	Not estimable	1.00 [0.95, 1.00]		-
Ptak 2001	59	0	1	616	0.98 [0.91, 1.00]		0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1

Note that for the Nguyen 2005 low-risk analysis, sensitivity could not be calculated as there were no references standard positives in this group.

82

83

Figure 13: Any with suspected blunt cervical spine injury, >75% with head CT, CT + MRI as reference standard (only sensitivity possible), any cervical spine injury as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Friesen 2013	115	0	24	0	0.83 [0.75, 0.89]	Not estimable	*	
							0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1

Note that false positives are not possible when the index test forms part of reference standard, meaning specificity cannot be calculated (n=67 were negative on both CT and MRI, representing true negatives but not input into plot as false specificity of 100% calculated)

84

85

Figure 14: Any with suspected blunt cervical spine injury, unclear if head injury, CT + MRI as reference standard (only sensitivity possible), unstable cervical spine injury as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Songur 2020	63	0	18	0	0.78 [0.67, 0.86]	Not estimable _I	_	
								0 0.2 0.4 0.6 0.8 1

Note that false positives are not possible when the index test forms part of reference standard, meaning specificity cannot be calculated (n=7 were negative on both CT and MRI, representing true negatives but not input into plot as false specificity of 100% calculated)

86

Figure 15: Any with suspected blunt cervical spine injury, unclear if head injury, CT + MRI as reference standard (only sensitivity possible), any cervical spine injury as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Malhotra 2018	368	0	149	0	0.71 [0.67, 0.75]	Not estimable		
Novick 2018	88	0	13	0	0.87 [0.79, 0.93]	Not estimable	-	
Schoenfeld 2018	195	0	53	0	0.79 [0.73, 0.84]	Not estimable	0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1

Note that false positives are not possible when the index test forms part of reference standard, meaning specificity cannot be calculated (n=563, n=140 and n=420 were negative on both CT and MRI in Malhotra 2018, Novick 2018 and Schoenfeld 2018, respectively,, representing true negatives but not input into plot as false specificity of 100% calculated)

88

E.1893 MRI as index test

Figure 16: Any with suspected blunt cervical spine injury, >75% with head CT, CT + MRI as reference standard (only sensitivity possible), any cervical spine injury as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Friesen 2013	98	0	41	0	0.71 [0.62, 0.78]	Not estimable _P		
								0 0.2 0.4 0.6 0.8 1

Note that false positives are not possible when the index test forms part of reference standard, meaning specificity cannot be calculated (n=67 were negative on both CT and MRI, representing true negatives but not input into plot as false specificity of 100% calculated)

90

Figure 17: Any with suspected blunt cervical spine injury, unclear if head injury, CT + MRI as reference standard (only sensitivity possible), unstable cervical spine injury as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Songur 2020	81	0	0	0	1.00 [0.96, 1.00]	Not estimable _I	-+++++	
							0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1

Note that false positives are not possible when the index test forms part of reference standard, meaning specificity cannot be calculated (n=7 were negative on both CT and MRI, representing true negatives but not input into plot as false specificity of 100% calculated)

92

93

Figure 18: Any with suspected blunt cervical spine injury, unclear if head injury, CT + MRI as reference standard (only sensitivity possible), any cervical spine injury as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI) Specificity (95% CI)
Malhotra 2018	427	0	90	0	0.83 [0.79, 0.86]	Not estimable	
Novick 2018	78	0	23	0	0.77 [0.68, 0.85]	Not estimable	-
Schoenfeld 2018	248	0	0	0	1.00 [0.99, 1.00]		0 0.2 0.4 0.6 0.8 1 0 0.2 0.4 0.6 0.8 1

Note that false positives are not possible when the index test forms part of reference standard, meaning specificity cannot be calculated (n=563, n=140 and n=420 were negative on both CT and MRI in Malhotra 2018, Novick 2018 and Schoenfeld 2018, respectively,, representing true negatives but not input into plot as false specificity of 100% calculated)

E.952 Adults – only including those admitted, not those subsequently discharged following index test

E.1.9261 X-ray as index test

Figure 19: Any admitted with blunt injury and cervical spine assessed, unclear if head injury, reference standard unclear/final diagnosis, cervical spine injuries as outcome

Note that for this study, FP and TN were not reported and values were not necessarily zero. Only sensitivity could be calculated for this analysis.

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98

Figure 20:	Α	ny a	adm	nitteo	following traum	na, unclear if hea	d injury, CT as re	ference standard	d, cervical spine fractures as outcome
Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)	
Duane 2008	16	7	68	913	0.19 [0.11, 0.29]				

E.1.202 CT as index test

Figure 21: Any admitted following injury and cervical spine assessed, unclear if head injury, final diagnosis as reference standard, clinically significant cervical spine injuries as outcome

101

- E1023 Adults only including those that are obtunded, unconscious and/or requiring intensive care unit admission
- E.1.0331 CT as index test
 - Figure 22: High risk trauma patients (pain, neurological symptoms or obtundation), unclear if head injury, final diagnosis as reference standard, any cervical spine injury as outcome
 - 104 Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Adams
 2006).

106

Figure 23: High risk severely injured patients, unclear if head injury, final diagnosis as reference standard, cervical spine abnormality as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Widder 2004	18	0	0	0	1.00 [0.81, 1.00]	Not estimable _I		
							0 0.2 0.4 0.6 0.8 1	

Note that for this study, FP and TN were not reported and values were not necessarily zero. Only sensitivity could be calculated for this analysis.

107

108

Figure 24: High risk blunt trauma and admission to intensive care unit, unclear if head injury, final diagnosis as reference standard, unstable cervical spine injury as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI) Sensit	tivity (95% CI)	Specificity (95% CI)
Berne 1999	8	0	0	50	1.00 [0.63, 1.00]			

109

110

Figure 25: Unconscious intubated trauma patients, unclear if head injury, final diagnosis as reference standard, unstable cervical spine injury as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Brohi 2005	29	4	0	348	1.00 [0.88, 1.00]			
						(0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1

111

Figure 26: Altered sensorium admitted following injury and cervical spine assessed, unclear if head injury, final diagnosis as reference standard, clinically significant cervical spine injuries as outcome

113 Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Raza2013).

115

Figure 27: Obtunded with intracranial haemorrhage diagnosis following non high-impact trauma, CT + MRI as reference standard (only sensitivity possible), unstable cervical spine injury as outcome

116 Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Tan 2014).

- 118
- 119

Figure 28: Obtunded patients, possibly all had brain assessment, CT + MRI as reference standard (only sensitivity possible), any cervical spine injuries as outcome

Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Lau 2018).

122

123

Figure 29: Obtunded patients (mostly adults), unclear if head injury, CT + MRI as reference standard (only sensitivity possible), clinically significant cervical spine injury as outcome

Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Fisher 2013).

Figure 30: Unconscious adults, unclear if head injury, CT + MRI as reference standard (only sensitivity possible), any cervical spine injury as outcome

Study	TP FI	P FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Parmar 2018	20 () 7	0	0.74 [0.54, 0.89]			0 0.2 0.4 0.6 0.8 1

Note that false positives are not possible when the index test forms part of reference standard, meaning specificity cannot be calculated (n=0 were negative on both CT and MRI, representing true negatives)

128

E.1292 MRI as index test

- Figure 31: Obtunded with intracranial haemorrhage diagnosis following non high-impact trauma, CT + MRI as reference standard (only sensitivity possible), unstable cervical spine injury as outcome
- Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Tan
 2014).
- 132
- 133
- Figure 32: Obtunded patients, possibly all had brain assessment, CT + MRI as reference standard (only sensitivity possible), any cervical spine injuries as outcome
- 134 Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Lau135 2018).

Figure 33: Obtunded patients (mostly adults), unclear if head injury, CT + MRI as reference standard (only sensitivity possible), clinically significant cervical spine injury as outcome

138 Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Fisher2013).

140

141

Figure 34: Unconcious adults, unclear if head injury, CT + MRI as reference standard (only sensitivity possible), any cervical spine injury as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Parmar 2018	26	0	1	0	0.96 [0.81, 1.00]			
						(0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1

Note that false positives are not possible when the index test forms part of reference standard, meaning specificity cannot be calculated (n=0 were negative on both CT and MRI, representing true negatives)

142

- E1434 Adults other very specific populations
- E.1.4441 X-ray as index test
 - Figure 35: Those with diffuse idiopathic skeletal hyperostosis with low energy trauma, unclear if head injury, whole spine CT as reference standard, acute fracture of cervical spine as outcome

Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Dan Lantsman 2020).

Figure 36: Those with blunt trauma with CT and flexion-extension radiographs for continued cervical pain (fractures already excluded), unclear if head injury, all available evidence as reference standard, ligamentous cervical spine injury as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Goodnight 2008	6	10	0	363	1.00 [0.54, 1.00]			0 0.2 0.4 0.6 0.8 1

148

E.1.4492 CT as index test

Figure 37: Intoxicated adults with blunt trauma, unclear if head injury, final/discharge diagnosis as reference standard, unstable cervical spine injury as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Bush 2016	13	0	1	617	0.93 [0.66, 1.00]			0 0.2 0.4 0.6 0.8 1

150

Figure 38: Those with blunt trauma with CT and flexion-extension radiographs for continued cervical pain, unclear if head injury, all available evidence as reference standard, ligamentous cervical spine injury as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Goodnight 2008	6	13	0	360	1.00 [0.54, 1.00]	•		0 0.2 0.4 0.6 0.8 1

152

- E1\$35 Children all having index test and not limited to those that were admitted
- E.1541 X-ray as index test
 - Figure 39: Children with possible neck injury, 40% had head CT, reference standard unclear/follow-up/other imaging, cervical spine injury as outcome
 - 155 Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Somppi 156 2018).

157

- Figure 40: Children with cervical spine imaging, unclear if head injury, reference standard as CT, cervical spine injury as outcome
- 158 Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Rana
 159 2009).

E.1.6512 CT as index test

Figure 41: Children with possible neck injury, 40% had head CT, reference standard unclear/follow-up/other imaging, cervical spine injury as outcome

162 Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Somppi 163 2018).

164

Figure 42: Children with cervical spine imaging, unclear if head injury, reference standard as other imaging findings (unclear), cervical spine injury as outcome

165 Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Rana2009).

167

Figure 43: Children following trauma, unclear if head injury, reference standard as final diagnosis/unclear, unstable cervical spine injuries as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Derderian 2019	33	28	0	160	1.00 [0.89, 1.00]			0 0.2 0.4 0.6 0.8 1

168

Figure 44: Children following trauma, unclear if head injury, reference standard as MRI, any soft tissue cervical spine injury as outcome

170

E.1.513 MRI as index test

Figure 45: Children with possible neck injury, 40% had head CT, reference standard unclear/follow-up/other imaging, cervical spine injury as outcome

Raw data not reported and Forest Plot could therefore not be generated. Results are provided in GRADE tables as reported in the paper (Somppi 2018).

172

Figure 46: Children following trauma, unclear if head injury, reference standard as final diagnosis/unclear, unstable cervical spine injuries as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Derderian 2019	33	104	0	84	1.00 [0.89, 1.00]	0.45 [0.37, 0.52]	-	I - ∎-
Henry 2013-1	1	2	0	70	1.00 [0.03, 1.00]	1		0 0.2 0.4 0.6 0.8 1

174

175

Figure 47: Children following trauma, unclear if head injury, reference standard as CT, any osseous cervical spine injury as outcome

Study	TP	FP	FN	TN	Sensitivity (95% CI)	Specificity (95% CI) Sensitivity (95% CI) Specificity (95% CI)	
Henry 2013-2	6	2	0	76	1.00 [0.54, 1.00]	0.97 [0.91, 1.00]	
						0 0.2 0.4 0.6 0.8 1 0 0.2 0.4 0.6 0.8 1	

E1176 Children – only including those that are obtunded, unconscious and/or requiring intensive care unit admission

E.1.7681 X-ray as index test

Figure 48: Children with severe injuries admitted to intensive care unit, unclear if head injury, reference standard as clinical outcome/diagnosis at time of discharge/latest follow-up, cervical spine instability requiring surgery as outcome

179

E.18602 CT as index test

Figure 49: Children with cervical spine assessment and confirmed severe traumatic brain injury, reference standard as CT + MRI possibly other imaging, unstable cervical spine injury as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Qualls 2015	5	9	0	49	1.00 [0.48, 1.00]	0.84 [0.73, 0.93] _I		
								0 0.2 0.4 0.6 0.8 1

181

Figure 50: Children with severe injuries/unconscious, unclear if head injury, reference standard as final diagnosis/all information, injuries requiring stabilisation/surgical correction as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Al-Sarheed 2020	28	0	5	32	0.85 [0.68, 0.95]	1.00 [0.89, 1.00]		
Brockmeyer 2012	1	0	0	23	1.00 [0.03, 1.00]			

183

E.18643 MRI as index test

Figure 51: Children with cervical spine assessment and confirmed severe traumatic brain injury, reference standard as CT + MRI possibly other imaging, unstable cervical spine injury as outcome

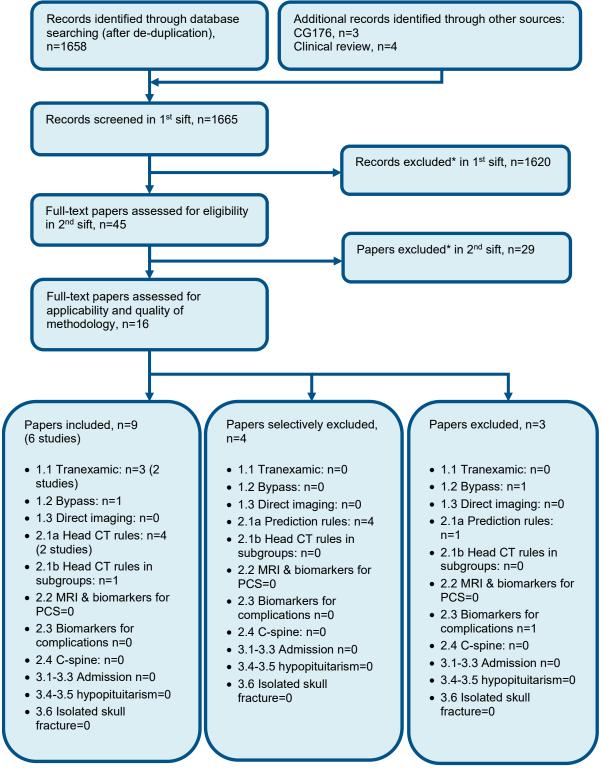
Study	TP FP	FN TN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Qualls 2015	4 11	1 47	0.80 [0.28, 0.99]	0.81 [0.69, 0.90]		
				(0 0.2 0.4 0.6 0.8 1	0 0.2 0.4 0.6 0.8 1

185

Figure 52: Children with severe injuries admitted to intensive care unit, unclear if head injury, reference standard as clinical outcome/diagnosis at time of discharge/latest follow-up, cervical spine instability requiring surgery as outcome

Study	TP	FP	FN	ΤN	Sensitivity (95% CI)	Specificity (95% CI)	Sensitivity (95% CI)	Specificity (95% CI)
Brockmeyer 2012	1	6	0	17	1.00 [0.03, 1.00]	•		0 0.2 0.4 0.6 0.8 1

1 Appendix F – Economic evidence study selection



* Non-relevant population, intervention, comparison, design or setting; non-English language

1 Appendix G – Economic evidence tables

2 None.

Appendix H – CG176 Health economic model (2014)

2 H.1 Methods

3 H.1.1 Model overview

Head injury (HI) patients can sustain bony and/or soft tissue injuries to the cervical (C) spine.
Whether patients experience a soft tissue injury becomes relevant after the initial imaging
shows a bony injury, or if the initial imaging is negative but the clinical picture still suggests
that there is a high risk of a cervical spine injury (CSI), in which case patients will experience
solely a soft tissue injury of the C-spine.

- 9 CG56 included a tentative cost analysis on this topic, with the comparison between the 10 NEXUS and the Canadian CT rule for CSI prediction. It was estimated that the Canadian rule 11 could save from £4 to £14 per patient to the NHS. However, this cost analysis had limited 12 validity due to the use of overseas data and simplified assumptions with regards to dealing 13 with indeterminate diagnostic imaging results.
- 14The management of patients with HI and suspected CSI is particularly challenging in terms of15resource implications. The main trade offs for this topic are represented by the cost of the16diagnostic tests (whether X-ray, CT scan and MRI) versus the failure to detect their CSI17(false negatives).
- The guidelines update of the CG56 literature review found no new economic evidence since
 the publication of CG56 on the cost-effectiveness of clinical prediction rules for any of the
 clinical questions for this topic.
- As a consequence, the GDG has identified this topic as a high priority for an original economic analysis.
- 23 The economic analysis will address the following clinical question:
- 24 Q1. What is the best clinical prediction rule for determining which patients with head injury 25 should be imaged (initial imaging with X-ray or CT) for cervical spine injury?

26 H.1.1.1 Comparators

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- 27 Seven clearance strategies for patients with HI and suspected CSI were devised to allow for 28 differential use of diagnostic imaging.
- 29 The strategies compared in this cost-effectiveness analysis are:
 - **CT on all:** In this strategy, no prediction rule is used. Everyone with HI and suspected CSI is given a CT scan.
 - **X-ray on all:** In this strategy, no prediction rule is used. Everyone with HI and suspected CSI is given an X-ray.
 - **CT according to NEXUS:** In this strategy, the NEXUS prediction rule is used to determine whether a CT scan is necessary. Only under the direction of the NEXUS prediction rule is a CT scan undertaken.
 - **CT according to Canadian C-Spine:** In this strategy, the Canadian C-spine prediction rule is used to determine whether a CT scan is necessary. Only under the direction of the Canadian C-spine prediction rule is a CT scan undertaken.
- **X-ray according to NEXUS:** In this strategy, the NEXUS prediction rule is used to determine whether an X-ray is necessary. Only under the direction of the NEXUS prediction rule is an X-ray undertaken.

- X-ray according to Canadian C-spine: In this strategy, the Canadian C-spine prediction rule is used to determine whether an X-ray is necessary. Only under the direction of the Canadian C-spine prediction rule is an X-ray undertaken.
- **No imaging:** In this strategy, patients with HI and suspected CSI do not receive any diagnostic imaging.
- 6 The CT on all, X-ray on all, and No imaging strategies were included as theoretical strategies 7 to explore the overall cost-effectiveness of diagnostic imaging. In practice, the first two 8 strategies are not feasible and the last is not acceptable.

9 H.1.1.2 Population

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10The population of the model consists of patients over the age of 16 with HI and suspected11CSI.

12 H.1.1.3 Time horizon, perspective, discount rates used

13 The analysis took the perspective of the UK NHS. The time horizon of the model was one in-14 hospital episode including diagnosis and treatment, discounting was therefore not applicable.

15 H.1.1.4 Deviations from NICE reference case

- 16 The search for quality of life evidence did not identify any data which the GDG felt applicable 17 to inform the expected health benefits for each diagnostic outcome. With long-term 18 management of CSI patients falling outside of the scope of this guideline, accurate data on 19 the long-term health outcomes and resource use associated with downstream management 20 were not available.
- 21 As a compromise, the GDG identified the cost of prevention of a false negative as the most 22 useful outcome for decision making and cautioned the interpretation of results due to the lack 23 of evaluation of all of the trade offs involved between the diagnostic outcomes (such as the benefit of true positives and negatives, and the health cost of the false positive, noting cost of 24 25 treating a false negative case was included in the analysis). To further assess the net cost of avoiding a false negative, a range of potential litigation costs of a false negative was 26 incorporated in a threshold sensitivity analysis. Also, a conservative hypothetical scenario 27 28 where minimal QALY gain was associated with a true positive and zero health or monetary cost associated with the false diagnostic outcomes was analysed. 29
- There is divergence from the NICE reference case as the main analysis is a costeffectiveness analysis (rather than a CUA) assessing the cost per diagnostic outcome in a time horizon limited to the diagnostic workup and short-term management. In addition, we employ the litigation cost which may be associated to a false negative and the underlying assumption that no clinical harm or cost (other than that of initial treatment) is associated to patients who have a false positive test result to assess cost-effectiveness. This further analysis is in essence a cost minimisation analysis.

37 H.1.1.5 Uncertainty

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The base case analysis employs expected values of costs, utilities and probabilities for model parameters and serves as base case analysis. If there are uncertainties about the values and assumptions used in the main cost-effectiveness analysis, sensitivity analyses are conducted. Results from base case and sensitivity analyses are compared. 1 There are two types of sensitivity analysis.

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8 9 **Deterministic Sensitivity Analysis** (DSA) is where the value of one of the parameters is changed to observe any effect on the results. This allows determination of the threshold at which a parameter's value is likely to change the conclusion. The GDG were uncertain about a number of parameters: the prevalence of CSI in a population, the cost of no procedure for patients with and without CSI, the clinical decision for further imaging after an initial X-ray / CT, and the specifications of initial imaging strategies (the probability of being given CT/X-ray/no imaging initially) and these uncertainties were tested by deterministic sensitivity analysis.

10 Probabilistic Sensitivity Analysis (PSA) is conducted to quantify parameter uncertainty. For every parameter subject to uncertainty (i.e. unit costs, sensitivities and specificities of the 11 12 prediction rules and clinician estimates), a distribution is assigned to reflect its uncertainty. Random draws across all parameter distributions are undertaken using Monte Carlo 13 14 methods. This process is repeated many times to build up a simulated sample of the expected value of the model output parameters, as well as a quantification of parameter 15 16 uncertainty. The PSA will determine the probability an intervention is cost-effective given a particular cost-effectiveness threshold. 17

18 H.1.2 Approach to modelling

19 The model is a decision tree which includes evidence on the prevalence of CSI among 20 patients with head injury as well as on intermediate outcomes (specificity and sensitivity) of all strategies being compared (for example X-rays, CT scans, MRI, prediction rules). The 21 combination of the prevalence of CSI with the specificity and sensitivity of each strategy 22 23 determines the proportion of patients who have abnormal, indeterminate and normal imaging 24 results. According to diagnostic imaging results, patients undergo a specific type of medical 25 management (observation, immediate discharge or surgical and non-surgical treatment). The 26 model tracks the number of patients for whom the clinical decision is appropriate (TP, TN) or inappropriate (FP, FN). 27

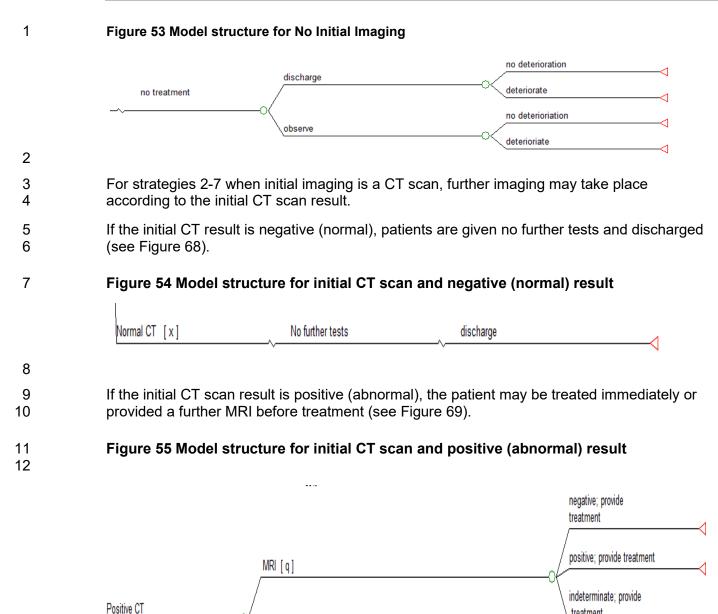
As there was limited data availability for survival and medical events (such as long term disability) following medical interventions received or not received by patients, the most important health outcome was considered to be the number of false negatives identified by each strategy.

32 H.1.2.1 Model structure

There are 7 clearance strategies for all patients with HI and suspected CSI regardless of the presence or absence of CSI. These seven strategies are described in M.1.1.1.

For Strategies 1 - 7 where no initial imaging is undertaken, patients are treated as normal,
 receive no treatment and are either discharged or observed in hospital for a period of 1 week
 (see

Figure 67). If patients have CSI, they may or may not experience deterioration. If patients do
 not have CSI, they do not experience deterioration.



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If the initial CT scan is indeterminate (Figure 70), the patient will undergo further diagnostic 15 imaging -- MRI or Flexion Extension (FE). If the second diagnostic imaging (MRI/FE) is 16 positive, the patient may be treated immediately or given a third diagnostic scan (MRI/FE). 17 Patients are treated if the third diagnostic scan is positive. Patients are observed in hospital 18 for a one week period if the third diagnostic scan is indeterminate. Patients receive no further 19 diagnostic imaging and are dischared if the third diagnostic scan is negative. 20

treat [f]

treatment

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Figure 56 Model structure for initial CT scan and indeterminate result

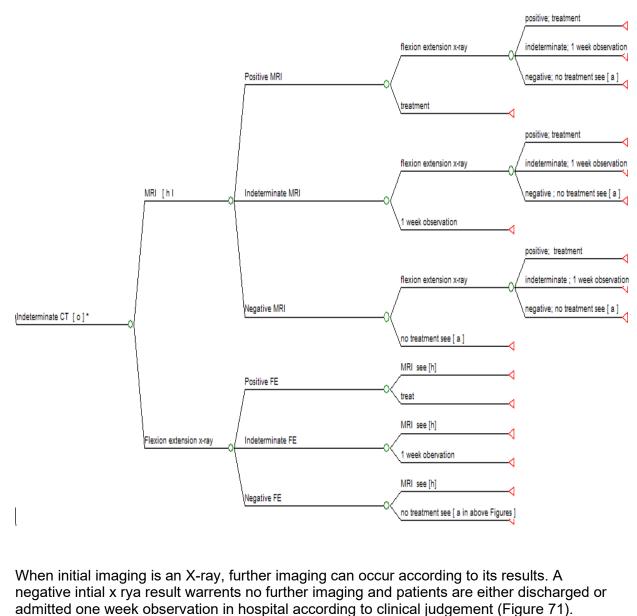
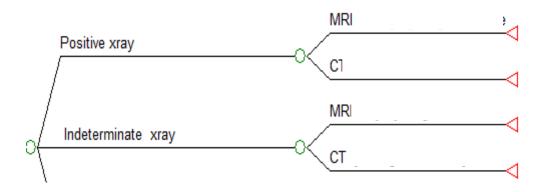


Figure 57 Model structure for initial X-ray and negative result



A positive or indeterminate X-ray result requires further imaging (MRI / CT). The model
structure following a MRI/CT scan is summarised in Figure 72. As the model structure here is
the same as those described and illustrated above, refer to Figure 68, Figure 69, and Figure
70, for details of the model structure following a CT scan and Figure 70, branch [h] for details
of the model structure following an MRI).

Figure 58 Model structure for initial X-ray and positive / indeterminate result



Patients who need treatment are provided specific procedures according to injury characterisics. Specifically, the GDG judged that the characteristics of a Cervical Spine Injury – bone; ligamentous; compression; stability; and presence of (Cervical Spinal Cord Injury) SCI— would determine the type of treatment required. The tree structure detailed in Figure 73 show the subcategorisation of injury characteristics and the appropriate corresponding procedure.

- Patients with complete or partial SCI and compression required a surgical or non-surgical
 procedure. Those who require surgery receive decompression and, where necessary, fusion.
 A collar could be provided in the case that a non-surgical procedure is deemed appropriate.
- Patients with partial or complete SCI and no compression were treated according to the
 stability of their injury. When the injury is stable, no procedure was necessary and instead,
 patients would receive a period of hospital observation. If the injury was unstable, a surgical
 or non-surgical procedure is required.
- Some patients with cervical spine injury will not have SCI. When these patients have stable
 injuries, then no procedure is required. Instead, they receive a period of hospital observation.
 However, if these patients sustain an unstable injury, surgical or non-surgical treatment is
 needed.

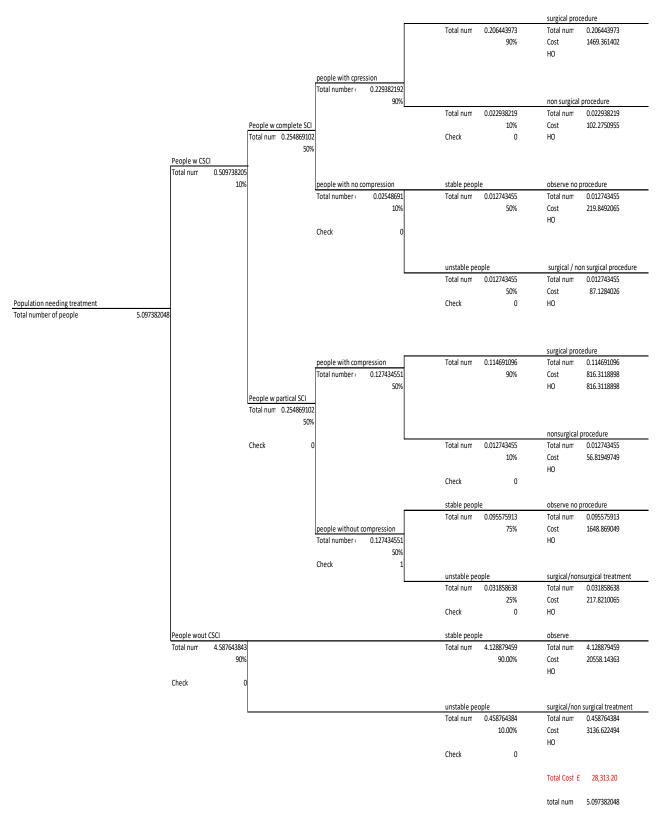
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Figure 59 Model structure for patients who require treatment



1 H.1.3 Model inputs

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2 H.1.3.1 Summary table of model inputs

Model parameters were based on clinical evidence identified in the systematic review undertaken for the guideline and supplemented by additional data sources when necessary. For example, a recent economic paper, Harlpen et al.24, was considered to be the best available source in the absence of a full systematic review on the diagnostic accuracy of the imaging modalities contained within the model. The authors had conducted a systematic search on these parameters, and several sources were used to inform the estimates used.

Model inputs were validated with clinical members of the GDG. In all but one instance only one source was identified in the clinical review to inform accuracy estimates of the clinical decision rules. In the case of the rule to x-ray by Canadian C-spine there was more than one source identified. In clinical validation of the sources in regards to their applicability and quality, the developers considered Coffey et al.8 to be the only appropriate source to inform the model for the following reasons. Throughout the guideline the developers placed more emphasis on recent UK studies, with Coffey et al. being the only source for this parameter to be both derived and validated in the UK context.

1718 A summary of the

A summary of the model inputs used in the base case (primary) analysis are provided in below. More details about sources, calculations and rationale for selection can be found in the sections following this summary table.

Parameter Deterministic Probability Distribution distribution description estimate parameters Source **Cohort Settings** Patients with HI 99.5%/0.5% GDG Expert Opinion and no CSI/with HI and CSI Cost of Prediction Rules (£) Canadian C-spine £0 Criteria are freely accessible **NEXUS** £0 Cost of Diagnostic Imaging (£) Calculated from 2011-2012 NHS Best fit distribution X-ray £30 identified according to reference cost codes DAPF methods described in Calculated from 2011-2012 NHS £60 Flexion, extension section M.1.4, reference cost codes DAPF and X-ray according to GDG Expert Opinion Table 70. CT £104 Calculated from 2011-2012 NHS reference cost codes RA08A. RA11Z and RA13Z MRI £182 Calculated from 2011-2012 NHS reference cost codes RA01A &RA04Z £0 No imaging N/A Cost of Treatment (£)

Table 30: Summary of base-case and sensitivity analysis model inputs

Parameter description	Deterministic estimate	Probability distribution	Distribution parameters	Source
Surgical procedure	£7,117	Best fit distri identified ac	bution	Calculated from 2011-2012 NHS reference cost code HC01-HC04
Surgical or Non- Surgical Procedure	£6,837	methods des section M.1.		Calculated from 2011-2012 NHS reference cost code HC01-HC06
Non –Surgical Procedure	£ 4,459	Table 70 .		Calculated from 2011-2012 NHS reference cost codes HC05-HC06
No procedure, (patients with SCI)	£17,252			Calculated from 2011-2012 NHS reference cost codes HC21B
No Procedure, (patients with no SCI)	£4,979			Calculated from 2011-2012 NHS reference cost code HC21C
Deterioration after treatment	£7,214			GDG Expert Opinion
Discharge	£0			
Performance of Pre	diction Tools			
Canadian C-spine X-ray - Sensitivity	1.00	Beta	α =8 , β =0	Clinical Paviaw, Coffay 20118
Canadian C-spine X-ray - Specificity	0.43	Beta	α = 605,β =807	Clinical Review- Coffey 2011 ⁸
Canadian C-spine CT - Sensitivity	1.00	Beta	α =192 , β =0	Clinical Review- Duane 2011A ¹⁵
Canadian C-spine CT - Specificity	0.06	Beta	α = 18,β =2991	
NEXUS X-ray - Sensitivity	0.91	Beta	α =147 , β =15	Clinical Review- Stiell 200345
NEXUS X-ray - Specificity	0.37	Beta	α = 2677,β =4599	
NEXUS CT - Sensitivity	0.90	Beta	α = 37, β =4	Clinical Review- Griffith 2011 ²³
NEXUS CT - Specificity	0.24	Beta	α = 364,β =1160	
Performance of X-ra	ау			
Sensitivity	0.568	Beta	α =334 ,β =254	Clinical Review- Halpern 2010 ²⁴
Normal results which are indeterminate	80%	Beta	α = 800, β =200	GDG Expert Opinion
Abnormal results which are indeterminate	10%	Beta	α = 100, β =900	
Specificity	0.997	Beta	α = 45822, β =138	Clinical Review- Halpern 2010 ²⁴
Normal results which are indeterminate	10%	Beta	α = 100, β =900	GDG Expert Opinion
Abnormal results which are indeterminate	80%	Beta	α = 800, β =200	

Parameter description	Deterministic estimate	Probability distribution	Distribution parameters	Source			
Performance of CT							
Sensitivity	0.832	Beta	α = 1545, β =312	Clinical Review- Halpern 2010 ²⁴			
Normal results which are indeterminate	90%	Beta	α = 900, β =100	GDG Expert Opinion			
Abnormal results which are indeterminate	10%	Beta	α = 100, β =900				
Specificity	0.999	Beta	α = 15335, β =15	Clinical Review- Halpern 2010 ²⁴			
Normal results which are indeterminate	10%	Beta	α = 100, β =900	GDG Expert Opinion			
Abnormal results which are indeterminate	90%	Beta	α = 900, β =100				
Performance of MRI							
Sensitivity	0.867	Beta	α = 386, β =59	Clinical Review- Halpern 2010 ²⁴			
Normal results which are indeterminate	10%	Beta	α = 100, β =900	GDG Expert Opinion			
Abnormal results which are indeterminate	0%						
Specificity	0.997	Beta	α = 565, β =2	Clinical Review- Halpern 2010 ²⁴			
Normal results which are indeterminate	10%	Beta	α =100 , β =900	GDG Expert Opinion			
Abnormal results which are indeterminate	0%						
Performance of FE-X	K-ray						
Sensitivity	0.568	Beta	α =334 , β =254	Clinical Review- Halpern 2010 ²⁴			
Normal results which are indeterminate	70%	Beta	α = 700, β =300	GDG Expert Opinion			
Abnormal results which are indeterminate	20%	Beta	α = 200, β = 800				
Specificity	0.997	Beta	α = 45822, β =138	Clinical Review- Halpern 2010 ²⁴			
Normal results which are indeterminate	90%	Beta	α = 900, β =100	GDG Expert Opinion			
Abnormal results which are indeterminate	50%	Beta	α = 500, β =500				
Clinical events (Positive Cases—Patients with CSI)							

Parameter	Deterministic estimate	Probability distribution	Distribution	Source
description	estimate	distribution	parameters	Source
After no imaging Probability clinician chooses immediate discharge	5%	Uniform	Min =4.5% , Max =5.5%	GDG Expert Opinion
Probability clinician chooses observation then discharge	95%	Uniform	Min = 85.5%, Max =100%	
After no imaging & discharge				
Probability deteriorate	95.0%	Uniform	Min = 85.5%, Max =100%	
Probability no deterioration	5.0%	Uniform	Min = 4.5%, Max =5.5%	
After no imaging & observe				
Probability deteriorate	20.0%	Uniform	Min = 18%, Max =22%	
Probability no deterioration	80.0%	Uniform	Min = 72%, Max =88%	
After abnormal initial CT result				
Probability clinician chooses MRI again	70%	Uniform	Min = 63%, Max =77%	
Probability clinician chooses to treat	30%	Uniform	Min = 27%, Max =33%	
After indeterminate initial CT result				
Probability clinician chooses MRI again	60%	Uniform	Min = 54%, Max =66%	
Probability clinician chooses flexion/extension x- ray	40%	Uniform	Min = 36%, Max =44%	
After indeterminate CT and abnormal MRI				

Descent	Determine	Duckskiller	Distribution	
Parameter description	Deterministic estimate	Probability distribution	Distribution parameters	Source
Probability clinician chooses flexion/extension x- ray	10%	Uniform	Min = 9%, Max =11%	
Probability clinician chooses to treat	90%	Uniform	Min = 81%, Max =99%	
After indeterminate CT and indeterminate MRI				
Probability clinician chooses flexion/extension x- ray	50%	Uniform	Min = 45%, Max =55%	
Probability clinician chooses to observe	50%	Uniform	Min = 45%, Max =55%	
After indeterminate CT and normal MRI				
Probability clinician chooses flexion/extension x- ray	20%	Uniform	Min = 18%, Max =22%	
Probability clinician chooses to discharge	70%	Uniform	Min = 63%, Max =77%	
Probability clinician chooses to observe 1 week	10%	Uniform	Min = 9%, Max =11%	
After indeterminate CT and abnormal flexion-extension				
Probability clinician chooses to treat	5%	Uniform	Min = 4.5%, Max =5.5%	
Probability clinician chooses MRI	95%	Uniform	Min = 85.5%, Max =100%	
After indeterminate CT and indeterminately flexion-extension				

-	B (1 · · /)	Dura ha hall the	Distribution	
Parameter description	Deterministic estimate	Probability distribution	Distribution parameters	Source
Probability clinician chooses to observe and discharge	1%	Uniform	Min = 0.9%, Max =1.1%	
Probability clinician chooses MRI	99%	Uniform	Min =89.1%, Max =100%	
After indeterminate CT and normal flexion-extension				
Probability clinician chooses to discharge	40%	Uniform	Min =36% , Max =44%	
Probability clinician chooses MRI	60%	Uniform	Min = 54%, Max =66%	
After first x-ray is abnormal				
Probability clinician chooses CT	95%	Uniform	Min =85.5%, Max =100%	
Probability clinician chooses MRI	5%	Uniform	Min = 4.5%, Max =5.5%	
After first x-ray is indeterminate				
Probability clinician chooses CT	99%	Uniform	Min =89.1%, Max =100%	
Probability clinician chooses MRI	1%	Uniform	Min = 0.9%, Max =1.1%	
After first x-ray is normal				
Probability clinician chooses discharge	95%	Uniform	Min =85.5%, Max =100%	
Probability clinician chooses observe	5%	Uniform	Min = 4.5%, Max =5.5%	
Clinical events (Neg	ative Cases—Pa	atients withou	t CSI)	
After no imaging				GDG Expert Opinion
Probability clinician chooses immediate discharge	95%	Uniform	Min =85.5%, Max =100%	

Parameter	Deterministic estimate	Probability distribution	Distribution
description Probability clinician chooses observation then discharge	5%	Uniform	parameters Min = 4.5%, Max =5.5%
After abnormal initial CT result			
Probability clinician chooses MRI again	99%	Uniform	Min =89.1%, Max =100%
Probability clinician chooses to treat	1%	Uniform	Min = 0.9%, Max =1.1%
After indeterminate initial CT result			
Probability clinician chooses MRI again	90%	Uniform	Min =89.1%, Max =100%
Probability clinician chooses flexion/extension x-ray	10%	Uniform	Min =9% , Max =11%
After indeterminate CT and abnormal MRI			
Probability clinician chooses flexion/extension x-ray	10%	Uniform	Min = 9%, Max =11%
Probability clinician chooses to treat	90%	Uniform	Min = 81%, Max =99%
After indeterminate CT and indeterminate MRI			
Probability clinician chooses flexion/extension x-ray	35%	Uniform	Min =31.5%, Max =38.5%
Probability clinician chooses to observe	65%	Uniform	Min =58.5%, Max =71.5%

Deveneter	Deterministic	Drobobility	Distribution	
Parameter description	Deterministic estimate	Probability distribution	Distribution parameters	Source
After indeterminate CT and normal MRI				
Probability clinician chooses flexion/extension x-ray	1%	Uniform	Min = 0.9%, Max =1.1%	
Probability clinician chooses to discharge	98%	Uniform	Min =88.2%, Max =100%	
Probability clinician chooses to observe 1 wee ⁵²⁰ k	1%	Uniform	Min =0.9% , Max =1.1%	
After indeterminate CT and abnormal flexion- extension				
Probability clinician chooses to treat	10%	Uniform	Min = 9%, Max =11%	
Probability clinician chooses MRI	90%	Uniform	Min = 81%, Max =99%	
After indeterminate CT and indeterminate flexion- extension				
Probability clinician chooses to observe and discharge	5%	Uniform	Min = 4.5%, Max =5.5%	
Probability clinician chooses MRI	95%	Uniform	Min =85.5%, Max =100%	
After indeterminate CT and normal flexion- extension				
Probability clinician chooses to discharge	50%	Uniform	Min = 45%, Max =55%	

Parameter description	Deterministic estimate	Probability distribution	Distribution parameters	Source			
Probability clinician chooses MRI	50%	Uniform	Min = 45%, Max =55%				
After first x-ray is abnormal							
Probability clinician chooses CT	95%	Uniform	Min = 85%, Max =100%				
Probability clinician chooses MRI	5%	Uniform	Min = 4.5%, Max =5.5%				
After first x-ray is indeterminate							
Probability clinician chooses CT	99%	Uniform	Min =89.1% Max =100%				
Probability clinician chooses MRI	1%	Uniform	Min = 0.9%, Max =1.1%				
After first x-ray is normal							
Probability clinician chooses discharge	95%	Uniform	Min = 85%, Max =100%				
Probability clinician chooses observe	5%	Uniform	Min = 4.5%, Max =5.5%				
Clinical events (Tre	atment Clinical J	udgements)					
Of all patients needing treatment,				GDG Expert Opinion			
percentage who have Cervical Spinal Cord Injury (CSCI)	10%	Uniform	Min =9% , Max =11%				
percentage who do not have CSCI	90%	Uniform	Min = 81%, Max =99%				
Of all patients with CSCI,							
percentage who have complete CSCI ?	50%	Uniform	Min = 45%, Max =55%				
percentage who have partial CSCI	50%	Uniform	Min = 45%, Max =55%				

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Parameter description	Deterministic estimate	Probability distribution	Distribution parameters
description	estimate	ulstilloution	parameters
Of the patients with complete CSCI,			
percentage who have compression	90%	Uniform	Min = 81%, Max =99%
percentage who do not have compression	10%	Uniform	Min = 9%, Max =11%
Of the patients with complete CSCI and compression,			
Percentage who have surgical treatment	90%	Uniform	Min = 81%, Max =99%
Percentage who have non-surgical treatment	10%	Uniform	Min = 9%, Max =11%
Of the patients with complete CSCI and no compression,			
percentage who are stabl	50%	Uniform	Min = 45%, Max =55%
percentage who are unstable	50%	Uniform	Min = 45%, Max =55%
Of the notionte			
Of the patients with partial CSCI,			
percentage who have compression	50%	Uniform	Min = 45%, Max =55%
percentage who do not have compression	50%	Uniform	Min = 45%, Max =55%
Of patients with partial CSCI and compression,			
percentage who have surgical procedure	90%	Uniform	Min = 81%, Max =99%
percentage who have non-surgical procedure	10%	Uniform	Min = 9%, Max =11%

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Parameter description	Deterministic estimate	Probability distribution	Distribution parameters
Of patients with partial CSCI and no compression,			
percentage who are stable	75%	Uniform	Min = 68%, Max =83%
percentage who are unstable	25%	Uniform	Min = 23%, Max =28%
Of patients with no CSCI,			
percentage who are stable?	90.0%	Uniform	Min = 81%, Max =99%
percentage who are unstable	10.00%	Uniform	Min = 9%, Max =11%

CSI = Cervical Spine Injury; CT = Computed Tomography; FE = Flexion Extension X-ray; HI = Head Injury; MRI = Magnetic Resonance Imaging; NEXUS = National Emergency X-Radiography Utilisation Study;

3 H.1.3.2 Resource use and cost

4 NHS reference costs 2011-2012¹¹ were used to identify cost estimates for diagnostic imaging 5 and treatment for CSI used in the base case analysis. Details are reported below.

6 Diagnostic Imaging

Diagnostic imaging costs are routinely incorporated in inpatient HRG codes. However,
 Multiple Trauma HRG codes and Emergency Medicine HRG codes relevant to our population
 were considered inadequate for our purposes as these cost codes were minimally influenced
 by differences in diagnostic imaging interventions and were largely derived from surgical and
 medical procedures.

As a result, unbundled costs for diagnostic imaging were used to allow for clear cost
 differentiations. The GDG judged this to be appropriate especially because a significant
 proportion the population could have diagnostic imaging without patient admittance into
 hospital.

16 The cost of CT and MRI diagnostic imaging techniques were calculated by taking a weighted 17 average of total activities and cost in outpatient, direct access and other settings. The GDG 18 judged that a CT or MRI scan requires a scan of two areas considering patients will need 19 their head and cervical spine areas examined (NHS Reference Cost Codes 2011-2012 20 RA11Z; RA04Z). The cost of a CT was £104 and the cost of a MRI was £182.

21The cost of diagnostic imaging with x-ray (Plain Film Radiograph) was £30.3 and was22derived from NHS Reference Costs 2011-2012 cost code DAPF. The GDG judged a flexion23extension investigation would require 2 plain film X-rays with a total cost of £61.

24 Cost of treatment

25 Costs for treatment were derived from NHS Reference Costs 2011-2012, HC codes (Spinal 26 Surgery and Disorders Chapter). There is a certain degree of double counting as each NHS 27 reference cost code (HC01-HC06) is applied to more than one treatment cost calculation. This was deemed appropriate as the GDG judged procedures within NHS reference codes HC01-HC06 were applicable to multiple treatment categories.

A patient who is discharged upon clinical impressions and diagnostic imaging results showing no abnormality does not require treatment and accrues a cost of £0. The GDG judged the cost of discharge to be similar across all patients who remain alive. Thus, the cost of discharge was not considered necessary for our incremental analysis.

Some patients with CSI and in need of treatment are inappropriately discharged and
experience deterioration. The GDG assumed that a patient who deteriorates will again
present to the hospital, undergo diagnostic imaging, and then receive treatment. Assuming a
worst-case scenario where the diagnostic investigation requires all types of diagnostic
imaging (a CT, MRI, FE X-ray and an X-ray) and the treatment requires a surgical and/or
non-surgical procedure, the maximum cost for deterioration is £7,214. Those patients who do
not experience deterioration did not accrue any additional costs.

14 In particular, where a surgical procedure was deemed appropriate, the cost was £7,117, the 15 weighted average of NHS cost codes HC01-HC04. The cost of a non-surgical procedure was £4,459 and was the weighted average of NHS cost codes HC05 and HC06. Using the NHS 16 Reference cost code HC21B weighted across settings, the cost of no procedure with SCI 17 was £17,252 for an average length of stay of 42 days. The cost of a surgical or non-surgical 18 procedure was £6,837 calculated as the weighted average of NHS reference cost codes 19 20 HC01-HC06. According to the NHS reference cost code HC21C weighted across settings, the cost of no procedure for patients without CSI was £4,979 for an average length of stay of 21 22 5.6 days.

23 H.1.3.3 Diagnostic mark-up

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For each strategy, the diagnostic mark-up provides the total cost and number of diagnostic images undertaken per diagnostic technique (X-ray, CT, MRI, and FE X-ray). The total number of diagnostic imagings was the sum of diagnostic imagings undertaken at initial and at further imaging stages.

28 Initial Imaging

29 The number of patients who received initial imaging (CT, X-ray, or no imaging) was different according to strategy. In blanket Strategies 1-3, the entire cohort received initial CT / X-ray 30 imaging or no imaging. In Strategies 4-7, the number of patients who received initial imaging 31 was determined by the sensitivity and specificity of prediction rules. These strategies did not 32 33 indicate diagnostic imaging (CT/X-ray) for all patients. For Example, in Strategy 4 (Canadian 34 C-spine for X-ray), the prediction rule did not recommend an X-ray for 58% of patients 35 without CSI. The GDG assumed that these patients might still be imaged. To determine the 36 proportion of patients who would receive the remaining diagnostic imaging alternatives, the 37 GDG estimated half of all remaing patients would receive no imaging and the other half of all remaining patients would receive the alternative diagnostic imaging technique (CT/X-ray). In 38 39 Strategy 4, the prediction ruled did not recommend an X-ray for 58% of patients without CSI and of these 58%, 29% received CT and 29% received no Imaging Details on the GDG 40 41 estimated apportioning of patients to all initial imaging alternatives for each strategy can be found in Figure 74. 42

Figure 60

Probability of having a given initial image	Initi	Initial clinical decision				
strategy	(for th	ose without injury)	(for those with injury)			
	No imaging	CT first	X ray first	No imaging	CT first	X ray first
Strategy 1: No imaging	100%			100%		
Strategy 2: CT all		100%			100%	
Strategy 3: x ray all			100%			100%
Strategy 4: Canadian C spine for Xray	29%	29%	43%	0%	0%	100%
Strategy 5: Canadian C Spine for CT	49.7%	0.6%	49.7%	0%	100%	0%
Strategy 6: NEXUS for Xray	32%	32%	37%	4.65%	4.65%	90.70%
Strategy 7: NEXUS for CT	38%	24%	38%	5%	90%	5%

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Further Imaging

The number of further diagnostic imaging performed was determined by the results from the initial diagnostic imaging technique. Results from a diagnostic imaging technique were categorised as normal (diagnostic imaging and clinical impression finds no abnormality), indeterminate (diagnostic imaging and clinical impression finds presence or absence of injury uncertain) or abnormal (abnormality is clear from diagnostic imaging and clinical impression). The number of normal and abnormal results were derived by from the sensitivity (abnormal) and specificity (normal) of diagnostic clearance strategies found in published literature (Halpern 2010)²⁴. However, there is no data available to inform the number of indeterminate results from diagnostic imaging. The GDG considered that a certain proportion normal and abnormal results would be considered 'indeterminate' and that these proportions would differ for a population with CSI and a population without CSI.

- Patients who did not receive initial imaging and patients with normal initial imaging results
 would not be given any further imaging or treatment.
- Patients with an indeterminate or abnormal initial imaging result could receive further
 diagnostic imaging. The type and number of further diagnostic imaging (maximum number =
 3) was determined by clinical judgement.
- Therefore, the cost of diagnostic imaging was the product of the total number of diagnostic
 imagings undertaken per diagnostic technique and the unit cost of each diagnostic
 technique.
- 23 Where there is indication of abnormality from diagnostic imaging results and clinical 24 impressions, further management is required.

25 H.1.3.4 Treatment component

The treatment component uses GDG clinical judgments to subcategorise patients requiring treatment according to injury characteristics so as to identify the type of treatment required and apply the correct weighting to costs. These GDG judgements are detailed in section M.1.2.1. The cost of treatment was calculated as the sum of the cost of each category of treatment. The cost of each category of treatment was the product of the number of treatments and the unit cost of treatment.

32 H.1.3.5 Computations

The analysis was undertaken using Microsoft Excel 2010. The model is a cohort decisiontree. The PSA was conducted using 7500 simulations (see M.1.4). Each strategy is made up of a diagnostic and treatment component. The prevalence of CSI combined with the performance of prediction rules and the performance of diagnostic imaging techniques

1determined the number of patients correctly provided treatment (TP), incorrectly provided2treatment (FP), correctly left untreated (TN), and incorrectly left untreated (FN).

For computations informing estimation of cost effectiveness please refer to sections M.1.5
and M.1.6.

5 H.1.4 Sensitivity analyses

 A number of deterministic sensitivity analysis were undertaken to investigate uncertain individual input parameters. The GDG wished to identify whether varying that individual input value would have an effect on results. The following inputs were investigated using DSA.

- Cost of no procedure for patients with and without CSI: there was uncertainty around the cost differentiation for no procedure in patients with (£17,252) and without CSI (£4,979). Hence, the cost for no procedure was made equal for both patients with and without CSI at £ 5,141. This was the weighted cost of HC21B and HC21C across NHS settings for a 10 day length of stay.
 - 2. Litigation cost associated with a FN: given the uncertainty around the average litigation cost for a missed CSI, the litigation cost was varied from £0 to £1,000,000.
- 3. Initial imaging decisions: there was uncertainty over the base case percentage of patients without CSI who would receive initial imaging (CT/X-ray) or no imaging according to clinical decision rules in Strategies 1-7. Primary analysis percentages were calculated based on the sensitivity and specificity of clinical decision rules and GDG estimates. The uncertainty was attributed to the low quality of specificity data for prediction rules in Duane¹⁵ and Griffith²³. This was explored by calculating percentages using different GDG estimates as indicated in Figure 75 (see percentages highlighted by red rectangle).
- 4. QALY pay-offs: in the absence of applicable Quality of Life information for this population, an extremely conservative QALY pay-off was assigned to each outcome (TP, FN, TN, and FP) in a hypothetical scenario. The QALY payoffs assigned (TP = 1.5 QALYs, TN & FP =2 QALYs, and FN = 1 QALY) served to incorporate the smaller pay-off associated with a FN in comparison to patients without CSI (TN) and patients who received treatment (TP and FP). Net monetary benefit was subsequently calculated using Equation 1, where 'Outcome' was equal the number of QALYs and D was equal to the threshold of £20,000 per QALY gained.
- 5. Prevalence of patients over the age of 16 with CSI: given the absence of information on the prevalence of CSI, the prevalence was varied between 0.5% (base case) to 5% in increments of 0.5%.
- 6. Clinical decision for further imaging of indeterminate and negative initial imaging results: Given the absence of clinical and economic evidence on the clinical and costeffectiveness identified for Strategies 1-7 and their application to further imaging scenarios, the following scenarios were compared
 - a. further imaging on indeterminate cases only (base case analysis)
 - b. no further imaging on negative or indeterminate cases
 - c. further imaging on all negative and indeterminate cases
- 41 In scenarios a. to c., positive initial imaging results receive further imaging.

Figure 61 GDG estimation of initial imaging probabilites for those without injury (Strategy 4-7)

Probability of having a given initial image strategy		clinical decision se without injury		Initial clinical decision (for those with injury)			
7	No imaging	CT first	X ray first	No imaging	CT first	X ray first	
Strategy 1: No imaging	100%			100%			
Strategy 2: CT all		100%			100%		
Strategy 3: x ray all			100%			100%	
Strategy 4: Canadian C spine for Xray	54%	3%	43%	0%	0%	100%	
Strategy 5: Canadian C Spine for CT	54%	46%	0%	0%	100%	0%	
Strategy 6: NEXUS for Xray	60%	3%	37%	4.65%	4.65%	90.70%	
Strategy 7: NEXUS for CT	22%	40%	38%	5%	90%	5%	

Probabilistic Sensitivity Analysis

For the probabilistic analysis, inputs were parameterised with distributions as described in

Table 70 below. To parameterise the reference costs probabilistically, three distributions (gamma, lognormal and normal) were fitted and the best-fit distribution was chosen. Each distribution was fit using the standard deviation of the trust cost (calculated using the reported mean and interquartile range), and where appropriate, the distribution's alpha and beta values. The distribution that provided the interquartile range closest in value to the interquartile range reported by the NHS reference cost was considered the best fit distribution. Estimates from the best-fit distribution were applied to the formulas listed below to calculate the standard error of the mean NHS cost and subsequently, the probabilistic value was drawn.

probabilistic sensitivity analysis								
	Probability	Properties of distribution						
Parameter	distribution							
Clinical Judgements	Uniform	Uniform distribution fitted between the minimum and maximum range allows an equal chance of any value within this range being selected in any simulated run of the probabilistic analysis. The minimum and maximum range for clinical judgements was ±10% of the base case value with a maximum of 100%.						
Performance of prediction rules (sensitivity and specificity)	Beta	Beta distribution fitted between 0 and 1. As the sample size and the number of events were specified alpha and beta values were calculated as follows: Alpha: (number of patients with CSI/without CSI) Beta=(Number of patients)-(number of patients with CSI/without CSI)						
Performance of diagnostic imaging techniques (sensitivity and specificity)	Beta	Beta distribution fitted between 0 and 1. Derived from mean of a domain or total quality of life score and its standard error, using the method of moments. Alpha and Beta values were calculated as follows: Alpha = mean ² *(1-(mean/SE ²)-mean Beta = Alpha *((1-mean)/mean)						
Number of indeterminate results after imaging technique	Beta	Beta distribution fitted between 0 and 1. The sample size and the number of events were specified by the cohort size and GDG estimations. Thus, alpha and beta values were calculated as follows: Alpha = (number of patients with indeterminate result) Beta = (Number of patients)-(number of patients with indeterminate results)						
NHS Reference Costs (diagnostic and treatment)	Gamma	Gamma distribution bounded at 0 and positively skewed. Derived from mean and its standard error. Alpha and Beta values were calculated as follows: Alpha = (mean/SE) ² Beta = SE ² /Mean						
NHS Reference Costs (diagnostic and treatment)	Lognormal	Where appropriate, the lognormal distribution may provide a better fit than the gamma distribution for costs. The natural log of the mean was calculated as follows: Natural log of the mean = [Ln(mean) – (InSE) ²]/2 Where the natural log of the standard error (InSE) was calculated by: $\sqrt{\ln \frac{SE^2 + mean^2}{mean^2}}$						
NHS Reference Costs (diagnostic and treatment)	Normal	Where appropriate, the normal distribution may provide a better first than the gamma and lognormal distribution for costs. The mean and standard error was calculated as follows: $Mean = \frac{sum \ of \ all \ values}{number \ of \ values}$ Standard Error = $\frac{standard \ deviation}{\sqrt{number \ of \ values}}$						

Table 31: Description of the type and properties of distributions used in the probabilistic sensitivity analysis

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With all distributions drawn, a simulation was run for each strategy independently and key results of each simulation were copied and stored. To compare the results generated for a single iteration, the starting seed for each random number selected for the probabilistic analysis was reset to original with each rerun of the probabilistic simulation. This assured, for example, the PSA referred to the same prevalence for all seven strategies in any given iteration and ensured the results for each iteration across the strategies were comparable.

7 H.1.5 Estimation of cost effectiveness

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8 The widely used cost-effectiveness metric is the incremental cost-effectiveness ratio (ICER). This is calculated by dividing the difference in costs associated with two alternatives by the 9 difference in QALYs. The decision rule then applied is that if the ICER falls below a given 10 cost per QALY threshold the result is considered to be cost effective. If both costs are lower 11 and QALYs are higher the option is said to dominate and an ICER is not calculated. 12

$$ICER = \frac{Costs(B) - Costs(A)}{QALYs(B) - QALYs(A)}$$

 Cost-effective if: ICER < Threshold

Where: Costs/QALYs(X) = total costs/QALYs for option X

13 When there are more than two comparators, as in this analysis, options must be ranked in order of increasing cost then options ruled out by dominance or extended dominance before 14 15 calculating ICERs excluding these options. An option is said to be dominated, and ruled out, if another intervention is less costly and more effective. An option is said to be extendedly 16 dominated if a combination of two other options would prove to be less costly and more 17 18 effective.

It is also possible, for a particular cost-effectiveness threshold, to re-express costeffectiveness results in term of net monetary benefit (NMB). This is calculated by multiplying the total QALYs for a comparator by the threshold cost per QALY value (for example, £20,000) and then subtracting the total costs (formula below). The decision rule then applied is that the comparator with the highest NMB is the most cost-effective option at the specified threshold. That is the option that provides the highest number of QALYs at an acceptable cost.

$$NMB(X) = (QALYs(X) \times \lambda) - Costs(X)$$
 • Costs(X)

Where: NMB= Net Monetary Benefit; Costs/QALYs(X) = total costs/QALYs for option X; λ = threshold

t-effective if: highest net monetary benefit

Both methods of determining cost effectiveness will identify exactly the same optimal strategy. For ease of computation, adaptations of the NMB formula are used in this analysis to identify the optimal strategy.

In the case of cost-effectiveness analysis where cost per QALY is not estimated, and rather an alternative outcome (i.e. cost per false negative avoided) is used, there is not a specific cost per effect threshold employed to assess cost effectiveness. However, these outcomes can still be used to identify dominated and extendedly dominated options. Further, an assumed cost and/or QALY weight can be attached to such outcomes to enable net monetary benefit calculations, as described in the below equations:

$NMB(X) = \left(Outcome(X) \times QALY_Weight \times \lambda\right) - Costs(X)$	 Cost-effect highest net n
Where: NMB = Net Monetary benefit; Outcome(x) = the diagnostic	benef

outcome for which the QALY weight applies; λ = threshold of £20,000

ctive if: monetary fit

$NMB(X) = (FN(X) \times -LitigationCost) - Costs(X)$

Where: NMB = Net Monetary Benefit; FN = False negativs identified; litigation costs represents the negative cost associated with the false negative and Costs (x) is the total cost of the strategy

 Cost-effective if: highest net monetary benefit

H.1.6 Interpreting results 3

NICE's report 'Social value judgements: principles for the development of NICE guidance' sets out the principles that GDGs should consider when judging whether an intervention offers good value for money. In general, an intervention was considered to be cost effective if either of the following criteria applied (given that the estimate was considered plausible):

- The intervention dominated other relevant strategies (that is, it was both less costly in terms of resource use and more clinically effective compared with all the other relevant alternative strategies), or
- The intervention costs less than £20,000 per quality-adjusted life-year (QALY) gained 11 12 compared with the next best strategy.
 - In the absence of data to inform a lifetime costs and QALYs associated with the strategies (i.e. data on longterm survival and medical events), the model evaluates the diagnostic startegies using three types of analyses, each referencing a different key outcome. These are:
 - a) A cost effectiveness analysis which compares the cost per false negative avoided in a given strategy.
 - b) A cost minisation analysis whereby the litigation costs accrued are evaluated against the cost of the strategy, with results expressed in net monetary benefit.
 - c) A simplistic cost utility sensitivity analysis which compares the net monetary benefit associated with each strategy given minimal QALY gains per correct diagnosis and minimal QALY loss per incorrect diagnosis.

As we have several strategies of comparison, we use Net Monetary Benefit to rank the strategies on the basis of their relative cost-effectiveness and identify dominated or extendly dominated options.

27 A note on Net Monetary Benefit Analysis using litigation costs.

28 Using information on total cost and outcome and assuming the litigation cost penalty associated with a FN was -£200,000, net monetary benefit was calculated. This statistic was 29 30 calculated as the number of False Negatives multiplied by the cost penalty (a litigation penalty of -£200,000) minus the total cost of strategy (Equation 1). Because the cost penalty 31 of a false negative was greater than the total cost of strategy, the net monetary benefit figure 32 is negative. Net Monetary Benefit Results were ranked from 1 to 7 across all strategies with 33 34 Rank 1 representing the largest Net Monetary Benefit and Rank 7 as the least Net Monetary 35 Benefit.

36 To minimise costs, the GDG would consider the strategy with the highest net monetary 37 benefit. In the sensitivity analysis where QALYs were assigned to each outcome, the 38 monetary value associated with each QALY gained was £20,000. The GDG would consider the optimal or dominant strategy from this analysis when making recommendations. 39

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1 H.1.7 Model validation

2 The model was developed in consultation with the GDG; model structure, inputs and results 3 were presented to and discussed with the GDG for clinical validation and interpretation.

The model was systematically checked by the health economist undertaking the analysis;
this included inputting null and extreme values and checking that results were plausible given
inputs. The model was peer reviewed externally and by a second experienced health
economist from the NCGC; this included systematic checking of the model calculations.

8 H.2 Results

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9 H.2.1 Base Case Results

- Each strategy is composed of diagnostic imaging, outcomes, and treatments. Thus, Table 71
 Table 74 qualify the differences in base case deterministic diagnostic imagings, outcomes
 and treatments across strategies. Understanding these differences will help the interpretation
 of base case probabilistic results in Table 75 and Table 76.
- Table 71 presents a breakdown of the total number of diagnostic imaging according to the
 strategy. The table also shows the percentage of the cohort who receives each type of
 diagnostic imaging.
- 18**Table 72** presents a breakdown of the performance of each strategy. Outcomes are19considered as the percentage of TP, FN, TN and FP. In each strategy, majority of patients20without CSI are correctly diagnosed as TN and very few are incorrectly diagnosed as FP. A21significant proportion of patients with CSI are incorrectly diagnosed as FN. The strategy with22the smallest (28%) and largest (100%) percentage of FNs are Strategy 2/Strategy 5 (CT for23all/Canadian C-spine for CT) and Strategy 1 (No Imaging) respectively.
- As **Table 73** illustrates, very few patients are treated across strategies. At the extremes, no one is treated in Strategy 1 (No Imaging) and 7 patients out of 1,000 are treated in Strategy 5 (Canadian C-spine for CT). Of those who receive treatment, the majority do not receive a procedure but are instead observed in hospitals (those who are given no procedure with or without CSI).
- 29 Table 74 presents the total cost of each strategy. Strategy 3 (X-ray all) is most costly while 30 Strategy 1 (No Imaging) is least costly. The cost of each strategy is most influenced by the 31 cost of diagnostic imaging and the cost of observation. Because of the small number of 32 patients treated across strategies, the cost of treatment assumes a relatively small proportion 33 of the total cost of strategy. By considering both the number of diagnostic imaging results as well as the differential cost across types of diagnostic imaging, the total costs of each 34 35 strategy is calculated. The strategies with the highest (£289,558) and lowest (£0) diagnostic imaging costs are Strategy 2 (CT all) and Strategy 1 (No Imaging) respectively. 36 37
- The Net Monetary Benefit analysis (**Table 75**) provides the base case deterministic results and illustrates that Strategy 5 (Canadian C-spine for CT) is the optimal strategy (highest net monetary benefit) while Strategy 1 (No Imaging) was the least optimal (lowest net monetary benefit).
- In addition, **Table 76** presents the results of the cost effectiveness analysis where
 incremental costs and false negatives avoided were calculated using Strategy 1 (No
 Imaging) as the base comparator. The lowest (£88,458) and highest (£271,310) costs per
 false negative avoided were associated with Strategy 5 (Canadian C-Spine for CT) and
 Strategy 3 (X-ray on all) respectively.
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1 H.2.2 Sensitivity Analysis Results

2 Strategy 5 remained the optimal strategy in the probabilistic analysis, and it was the most 3 cost-effective strategy in 93% of the simulations. Strategy 5 was optimal despite variation to 4 individual inputs - equal costs for no procedure with or without CSI (Table 76); GDG 5 estimated initial imaging decisions (Table 78); QALY pay-offs (Table 79); prevalence of CSI 6 between 0.5%-5% (Figure 76) in the deterministic sensitivity analysis. Assuming that 7 Strategy 1 (No Imaging) is not an ethical option, Strategy 5 was also the optimal strategy when the litigation costs associated with a missed FN was between £0 and £1,000,000 8 9 (Table 77). Strategy 5 was also the optimal strategy when the clinical decision was to not further image normal and indeterminate results or to only further image indeterminate results. 10 When the clinical decision was to further image both normal and indeterminate results, 11 12 Strategy 2 (CT all) became optimal. 13

In the sensitivity analysis that assigned a minimal QALY advantage per correct diagnosis it
 was found that no imaging ranked optimal. If no imaging was not considered an acceptable
 or ethical strategy, then strategy 5 would be the most optimal strategy.

17 H.3 Interpreting results

18 H.3.1 Summary of results

19 The probabilistic analysis identified Strategy 5 (Canadian C-spine for CT) to be dominant at a 20 threshold of £200,000 for each FP outcome meaning that Strategy 5 was less costly and avoided more FPs than all the other strategies. Strategy 5 also had the lowest cost per False 21 22 Negative avoided. This conclusion was robust to variations in the prevalence of CSI (0.5%-5%), cost of no procedure with or without CSI and GDG estimated initial imaging decisions 23 and when the decision to not further image or to further image only indeterminate results. 24 25 When the clinical decision was to further image both normal and indeterminate results, the optimal strategy changed to Strategy 2 (CT all). 26

The results were sensitive to the cost of litigation associated with a false negative, with the optimal ranking switching from no imaging to strategy 5 when litigation costs rose from £75,000 to £100,000. No imaging was also seen as an optimal strategy if only a minimal QALY advantage was associated with achieving a true positive in comparison to other diagnostic outcomes. Strategy 5 was the next optimal strategy in this analysis.

32 H.3.2 Limitations and interpretation

We acknowledge the CEA does not fully account or quantify all the trade offs involved with the diagnostic decision question, as no weighting or penalty was given to other diagnostic outcomes such as false positives (although unnecessary treatment cost is taken into account). However, the estimated negative monetary payoff of £200,000 associated with each FN outcome implicitly took into account the adverse effects of radiation and the potential of deterioration after treatment or no treatment. Nonetheless, it is necessary to interpret this analysis with caution as it has some potentially serious limitations.

40 That the 'No Imaging' strategy may be optimal in scenarios where there are limited negative consequences associated with a false negative finding and where there is little to gain with 41 42 positive findings (i.e. correct onward treatment and QALY gain) is a reflection of the low prevalence of CSI within a head injury population and the trade off involved with the decision 43 problem. A low prevalence of a condition will inevitably lower the negative predictive values 44 45 of a diagnostic intervention (in comparison to if the diagnostic intervention was placed in a 46 high prevalence setting), an in turn favour a non-imaging strategy, especially when the downstream consequences of a correct or incorrect diagnostic are marginal in relation to 47 48 each other. In this model, an extremely conservative estimate of the gains of diagnostics was 49 specified.

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4 5 The GDG felt that despite the limitations, the analysis is sufficiently robust for purposes of decision-making as it explicitly shows and attempts to quantify the parameters, assumptions and structure underpinning the decision. To interpret the results, the GDG acknowledged that the consequences of each diagnostic outcome was uncertain, and took the view that in practice a non imaging strategy was not viable to recommend.

Assuming that Strategy 1 (No Imaging) was a theoretical strategy not plausible (ethical) in
 practice, the CT according to Canadian C-spine was optimal when the false negative
 litigation costs varied form £0 - £1,000,000. The conclusion that CT using the Canadian C spine prediction rule remained gave the greatest net monetary benefit in the scenario of
 minimal QALY gain associated with each true positive and minimal QALY loss with each
 false negative under the assumption that No Imaging was not appropriate in practice.

12 With the view that a non-imaging strategy could not be recommended, the sensitivity analysis whereby an extremely conservative scenario was explored in terms of pay-off indicates that 13 despite the limitations of the CEA, the conclusions formed by the analysis appear robust. In 14 addition, that Strategy 5 (CT according to Canadian C-spine) remained robust when the 15 threshold value associated with a FN was varied from £0 to £1,000,000 (assuming the No 16 Imaging strategy was not appropriate in practice) also supports the conclusions made in this 17 analysis. In line with the NICE reference case, all parameters subject to uncertainty (i.e. unit 18 19 costs, sensitivities and specificities of the prediction rules and clinician estimates) were 20 parameterised probabilistically and probabilistic sensitivity analysis performed.

21 H.3.3 Generalisability to other populations

22 A separate subgroup analysis was not conducted for a paediatric population. The results of 23 this analysis are not applicable for children under the age of 16 with HI and suspected CSI. 24 The GDG felt this economic analysis could not be extrapolated to the paediatric population 25 as this is clinically guite different from the adult population. No evidence was identified for 26 paediatrics and so, it was not possible to determine the appropriateness of model inputs for the paediatric population (in particular, the prevalence of CSI & the clinical judgements for 27 further imaging and treatment used in the analysis for adults). For this population, the trade-28 29 off between the accuracy of diagnosis and the radiation risk associated with a CT scan (equivalent to 2 years background radiation) requires particular discussion. The GDG would 30 31 consider that a plain film X-ray has lower levels of radiation than a CT scan when writing 32 recommendations for children.

33 H.3.4 Comparisons with published studies

34 No studies that looked at the use of prediction rules for the selection of HI patients with suspected CSI for diagnostic imaging were identified. One study by Pandor et al 2011,379 35 which investigated the use of prediction rules for the management of patients with minor HI 36 found that in comparison to 9 other strategies, the Canadian CT Head Rule (CCHR) medium 37 and high-risk prediction rule was the most cost-effective. Given this conclusion, the GDG 38 39 considered that the CCHR could be used for a patient with HI and suspected CSI to rule out 40 HI. Then, according to the conclusions from this analysis, Canadian CT Spine rule could be 41 used for the same patient to rule out suspected CSI.

42 H.3.5 Conclusion

43 For patients with HI and suspected CSI, the Canadian C-spine decision rule is cost-effective 44 for selecting patients for diagnostic imaging.

45 **H.3.6 Implications for future research**

46 The time horizon of this analysis only extended to the end of treatment. Considering this 47 short time horizon and exclusion of quality-of-life health outcomes in this analysis, future

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research could explore the costs and health outcomes for a lifetime horizon. Results from this analysis were not extrapolated to the patient subgroup under the age of 16 because of a dearth of available information. Should clinical studies that look at the accuracy of prediction rules for children be available in the future, this analysis can be modified to provide information on the cost-effectiveness of C-spine injury clearance strategies for this subgroup.

H.4 Additional Tables and Figures 1

Base Case Results

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Table 32 : Base Case Deterministic Analysis— Breakdown of Diagnostic Imaging for each Strategy

Base case Breakdown of Diagnostic Imaging for each strategy (prevalence 0.5% , cohort N = 1000)								
Strategy	# of Xrays (%)	# of CTs(%)	# of MRIs(%)	# of FE X- rays(%)				
Strategy 1: No imaging	0 (0%)	0 (0%)	0 (0%)	0 (0%)				
Strategy 2: CT all	0 (0%)	1000 (100%)	751 (75%)	812 (82%)				
Strategy 3: X- ray all	1000 (100%)	792 (80%)	602 (61%)	643 (65%)				
Strategy 4: Canadian C- spine for X-ray	433(43%)	626 (63%)	473 (48%)	508 (51%)				
Strategy 5: Canadian C- spine for CT	495 (50%)	403 (40%)	307 (31%)	326 (33%)				
Strategy 6: NEXUS for X- ray	371 (37%)	608 (61%)	459 (46%)	493 (50%)				
Strategy 7: NEXUS for CT	379 (38%)	542 (54%)	410 (41%)	439 (44%)				

Table 33: Base Case Deterministic Analysis – Performance of Strategies

Base case Results: Performance of Strategy (prevalence 0.5%, cohort N = 1000)									
	patients with	out CSI	patients with	n CSI					
Strategy	% True Negative	% False Positive	%False Negative	%True Positive					
Strategy 1: No imaging	100.0%	0.0%	100%	0%					
Strategy 2: CT all	99.7%	0.3%	28%	72%					
Strategy 3: X-ray all	99.7%	0.3%	56%	44%					
Strategy 4: Canadian C- spine for X-ray	99.8%	0.2%	56%	44%					
Strategy 5: Canadian C- spine for CT	99.8%	0.1%	28%	72%					
Strategy 6: NEXUS for X- ray	99.8%	0.2%	57%	43%					
Strategy 7: NEXUS for CT	99.8%	0.2%	33%	67%					

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Base case Results: Breal 1000)	kdown of [·]	Types of T	reatment	(prevalence	0.5%, coho	rt N =
Strategy	# of T* using surgical proced ures only	# of T* using non surgical procedu res only	# of T* where surgic alor non surgic al treatm ent is possibl e	#of T* with CSCI using no procedure	# of T* without CSCI using no procedure 2	Total # Treated
Strategy 1: No imaging	-	-	-	-	-	0.0
Strategy 2: CT all	0.4	0.0	0.7	0.1	5.5	6.8
Strategy 3: X-ray all	0.3	0.0	0.5	0.1	3.9	4.8
Strategy 4: Canadian C- spine for X-ray	0.3	0.0	0.4	0.1	3.4	4.2
Strategy 5: Canadian C- spine for CT	0.3	0.0	0.5	0.1	4.0	4.9
Strategy 6: NEXUS for X-ray	0.3	0.0	0.4	0.1	3.3	4.1
Strategy 7: NEXUS for CT	0.3	0.0	0.5	0.1	4.1	5.1

Table 34: Base Case Deterministic Analysis – Breakdown of Treatment Types

T* = treatments

¹ = Number of patients with CSCI where diagnostic result indicates the need for treatment but injury characteristics indicate that no surgical or non surgical procedure is beneficial. Thus, no procedure is provided.

²= Number of patients without CSCI where diagnostic result indicates the need for treatment but injury characteristics indicate that no surgical or non surgical procedure is beneficial. Thus, no procedure is provided.

Table 35: Base Case Deterministic Analysis – Breakdown of Cost of Strategy

Base case Breakdown o	Base case Breakdown of Costs of Strategy (prevalence 0.5%, cohort N = 1000)									
Strategy	Cost of Treatment	Cost of Diagnostic Imaging	Cost of Observation	Total Cost of Strategy						
Strategy 1: No imaging	£-	£-	£1,245	£1,245						
Strategy 2: CT all	£37,930	£289,558	£1,264	£328,753						
Strategy 3: X-ray all	£26,547	£260,916	£270,549	£558,012						
Strategy 4: Canadian C- spine for X-ray	£23,496	£194,888	£117,019	£335,403						
Strategy 5: Canadian C- spine for CT	£27,151	£132,283	£135,132	£294,566						
Strategy 6: NEXUS for X-ray	£22,957	£187,678	£100,324	£310,960						
Strategy 7: NEXUS for CT	£28,313	£168,905	£103,883	£301,102						

Table 36: Base Case Deterministic Analysis Results with Probabilistic Analysis Rank

Base Case Deterministic Analysis CEA Results (prevalence 0.5%, cohort N = 1000)										
Strategy	Total Cost of Strategy	Total # of False Negatives Identified	Net Monetary Benefit	Rank	% ranke d in PSA					
Strategy 1: No imaging	£1,245	5.00	- £1,001,245	6	0%					
Strategy 2: CT all	£328,753	1.42	-£612,099	2	7%					
Strategy 3: X-ray all	£558,012	2.79	- £1,116,022	7	0%					
Strategy 4: Canadian C- spine for X-ray	£335,403	2.79	-£893,413	5	0%					
Strategy 5: Canadian C- spine for CT	£294,566	1.42	-£577,912	1	93%					
Strategy 6: NEXUS for X- ray	£310,960	2.83	-£876,751	4	0%					
Strategy 7: NEXUS for CT	£301,102	1.66	-£633,022	3	0%					

Table 37: Base Case Probablistic Analysis—Cost per False Negative Avoided

Base Case Probablistic Results (prevalence 0.5%, cohort N = 1000)

Strategy	Total Cost of Strategy	Incremen tal Cost of Strategy	Total # of FN Identified	Increme ntal # of FN Avoided	Net Benefit	Incremental Cost per False Negative Avoided
Strategy 1: No imaging (reference)	£1,214	-	5.00	-	-£1,000,947	-
Strategy 2: CT all	£328,041	£326,828	1.69	3.31	-£665,914	£98,760
Strategy 3: x ray all	£556,884	£555,670	2.95	2.05	-£1,146,996	£271,310
Strategy 4: Canadian C spine for Xray	£333,997	£332,783	2.95	2.05	-£924,109	£162,483
Strategy 5: Canadian C Spine for CT	£293,948	£292,734	1.69	3.31	-£631,821	£88,458
Strategy 6: NEXUS for Xray	£310,297	£309,083	2.99	2.01	-£907,807	£153,875
Strategy 7: NEXUS for CT	£300,537	£299,324	1.91	3.09	-£683,070	£96,994

Deterministic Sensitivity Analyses Results

Table 38: DSA with Cost for No Procedure with or without CSI Equal

Deterministic Sensitivity Analysis on Costs for no procedure with and without CSI (prevalence 0.5%, cohort N = 1000, Equal cost for no procedure with and without CSI

Strategy	Total Cost of Strategy	Total # of False Negatives identified	Net Monetary Benefit	Ran k
Strategy 1: No imaging	£1,285	5.0	-£1,001,285	6
Strategy 2: CT all	£327,933	1.4	-£611,278	2
Strategy 3: X-ray all	£566,211	2.8	-£1,124,221	7
Strategy 4: Canadian C- spine for X-ray	£338,677	2.8	-£896,687	5
Strategy 5: Canadian C- spine for CT	£298,346	1.4	-£581,692	1
Strategy 6: NEXUS for X- ray	£313,702	2.8	-£879,493	4
Strategy 7: NEXUS for CT	£303,838	1.7	-£635,759	3

Table 39 DSA with Litigation Costs (£0 - £1,000,000)

Deterministic Sensitivity Analysis on Litigation costs (£0-£1,000,000) ; Prevalence of CSI 0.5%; Cohort N =1000

Litigation Cost												
Strategy	£0	£25,000	£50,000	£75,000	£100,000	£125,000	£150,000	£175,000	£200,000	£225,000	£250,000	£1,000,000
Strategy 1: No imaging	-£1,245	- £126,245	- £251,245	- £376,245	- £501,245	- £626,245	- £751,245	-£876,245	- £1,001,245	- £1,126,245	- £1,251,245	- £5,001,245
Strategy 2: CT all	- £328,753	- £364,171	- £399,589	- £435,008	- £470,426	- £505,844	- £541,262	-£576,680	-£612,099	-£647,517	-£682,935	- £1,745,482
Strategy 3: X-ray all	- £558,012	- £627,763	- £697,514	- £767,266	- £837,017	- £906,768	- £976,520	-£1,046,271	- £1,116,022	- £1,185,774	- £1,255,525	- £3,348,064
Strategy 4: Canadian C- spine for X-ray	- £335,403	- £405,154	- £474,906	- £544,657	- £614,408	- £684,160	- £753,911	-£823,662	-£893,413	-£963,165	- £1,032,916	- £3,125,455
Strategy 5: Canadian C- spine for CT	- £294,566	- £329,984	- £365,402	- £400,821	- £436,239	- £471,657	- £507,075	-£542,494	-£577,912	-£613,330	-£648,748	- £1,711,295
Strategy 6: NEXUS for X-ray	- £310,960	- £381,684	- £452,407	- £523,131	- £593,855	- £664,579	- £735,303	-£806,027	-£876,751	-£947,475	- £1,018,198	- £3,139,915
Strategy 7: NEXUS for CT	- £301,102	- £342,592	- £384,082	- £425,572	- £467,062	- £508,552	- £550,042	-£591,532	-£633,022	-£674,512	-£716,002	- £1,960,704

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Table 40: DSA with GDG estimates for initial imaging decisions

Base case CEA Results (prevalence 0.5%, cohort N = 1000, prediction rule performance according to GDG estimates)										
Strategy	Total Cost of Strategy	Total # of False Negatives identified	Net Monetary Benefit	Ran k						
Strategy 1: No imaging	£1,245	5.0	-£1,001,245	6						
Strategy 2: CT all	£328,753	1.4	-£612,099	2						
Strategy 3: X-ray all	£558,012	2.8	-£1,116,022	7						
Strategy 4: Canadian C- spine for X-ray	£335,403	2.8	-£893,413	5						
Strategy 5: Canadian C- spine for CT	£294,566	1.4	-£577,912	1						
Strategy 6: NEXUS for X- ray	£310,960	2.8	-£876,751	4						
Strategy 7: NEXUS for CT	£301,102	1.7	-£633,022	3						

Table 41 : DSA using QALY pay-offs (per cohort of 1000 patients)

	QALYs from TP ¹	QALYs from FN ²	QALYs from TN ³	QALYs from FP ⁴	Total QALY	NMB (£20K)	Rank
Strategy 1: No imaging	0.00	5.00	1990.00	0.00	1,995.00	£39,898,755	1
Strategy 2: CT all	5.37	1.42	1983.51	6.49	1,996.79	£39,607,080	4
Strategy 3: X-ray all	3.29	2.79	1983.42	5.17	1,994.67	£39,335,356	7
Strategy 4: Canadian C-spine for X-ray	3.29	2.79	1985.32	4.07	1,995.47	£39,574,054	6
Strategy 5: Canadian C-spine for CT	5.37	1.42	1986.69	2.61	1,996.09	£39,627,239	2
Strategy 6: NEXUS for X-ray	3.23	2.83	1985.53	3.95	1,995.54	£39,599,905	5
Strategy 7: NEXUS for CT	5.01	1.66	1985.95	3.52	1,996.13	£39,621,524	3

¹ QALYs from TP = # of TP multiplied by 1. 5 QALYs ² QALYs from FP = # of FP multiplied by 1 QALY ³ QALYs from TN = # of TN multiplied by 2 QALYs ⁴ QALYs from FP = # of FP multiplied by 2 QALYs

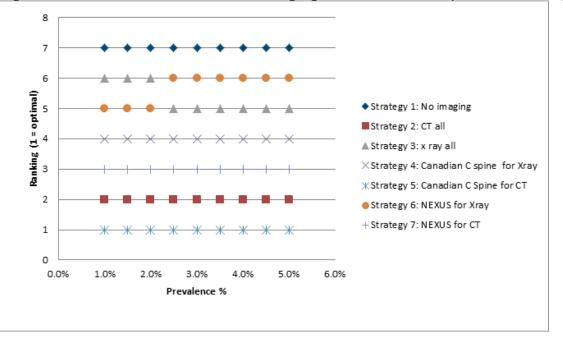
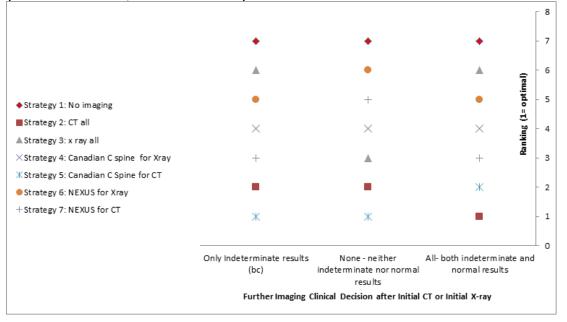


Figure 62 DSA with Prevalence of CSI ranging between 0.5%-5% (Cohort N =1000)

Figure 63 DSA with Further Imaging Clinical Decision Scenarios after Initial CT/X-ray (Prevalence 0.5%, Cohort N =1000)



5 Appendix I – Excluded studies

6 Clinical studies

7 Table 42: Studies excluded from the clinical review

Study	Code [Reason]
Albrecht RM, Kingsley D, Schermer CR et al. (2001) Evaluation of cervical spine in intensive care patients following blunt trauma. World journal of surgery 25(8): 1089-1096	- Study design - not diagnostic accuracy or test and treat
Anekstein, Y., Jeroukhimov, I., Bar-Ziv, Y. et al. (2008) The use of dynamic CT surview for cervical spine clearance in comatose trauma patients: a pilot prospective study. Injury 39(3): 339-46	- Study design - not diagnostic accuracy or test and treat
Antevil, J. L., Sise, M. J., Sack, D. I. et al. (2006) Spiral computed tomography for the initial evaluation of spine trauma: A new standard of care?. J Trauma 61(2): 382-7	- Not specifically cervical spine injury
Arbuthnot, Mary and Mooney, David P (2017) The sensitivity and negative predictive value of a pediatric cervical spine clearance algorithm that minimizes computerized tomography. Journal of pediatric surgery 52(1): 130-135	- Diagnostic test not relevant to review protocol
Awan, O., Safdar, N. M., Siddiqui, K. M. et al. (2011) Detection of cervical spine fracture on computed radiography images a monitor resolution study. Acad Radiol 18(3): 353-8	- Diagnostic test not relevant to review protocol
Bach CM, Steingruber IE, Peer S et al. (2001) Radiographic evaluation of cervical spine trauma. Plain radiography and conventional tomography versus computed tomography. Archives of orthopaedic and trauma surgery 121(7): 385-387	- Study design - only included those with injury already confirmed either on test or reference standard
Badhiwala, Jetan H, Lai, Chung K, Alhazzani, Waleed et al. (2015) Cervical spine clearance in obtunded patients after blunt traumatic injury: a systematic review. Annals of internal medicine 162(6): 429-37	- Study design - all were CT negative to be included
Berritto, Daniela, Pinto, Antonio, Michelin, Paul et al. (2017) Trauma Imaging of the Acute Cervical Spine. Seminars in musculoskeletal radiology 21(3): 184-198	- Review article but not a systematic review

Study	Code [Reason]
Bolinger, B.; Shartz, M.; Marion, D. (2004) Bedside fluoroscopic flexion and extension cervical spine radiographs for clearance of the cervical spine in comatose trauma patients. J Trauma 56(1): 132-6	- Study design - all were CT negative to be included
Brichko, Lisa, Giddey, Birinder, Tee, Jin et al. (2018) Cervical spine traumatic epidural haematomas: Incidence and characteristics. Emergency medicine Australasia : EMA 30(3): 359-365	- Study design - not diagnostic accuracy or test and treat
Brinckman, M.A.; Chau, C.; Ross, J.S. (2015) Marrow edema variability in acute spine fractures. Spine Journal 15(3): 454-460	- Diagnostic test not relevant to review protocol
	- Study design - only included those with injury already confirmed either on test or reference standard
Brown, C. V., Antevil, J. L., Sise, M. J. et al. (2005) Spiral computed tomography for the diagnosis of cervical, thoracic, and lumbar spine fractures: its time has come. J Trauma 58(5): 890-5; discussion 895	- Study design - only included those with injury already confirmed either on test or reference standard
Brown, C.V.R.; Foulkrod, K.H.; Reifsnyder, A. (2010) Computed Tomography versus Magnetic Resonance Imaging for Evaluation of the Cervical Spine: How Many Slices do you Need?. The American Surgeon 76(4): 365-368	- Insufficient information to calculate diagnostic accuracy measures
Carter, A.W., Jacups, S.P., Ackland, H.M. et al. (2017) Spinal clearance practices at a regional Australian hospital: A window to major trauma management performance outside metropolitan trauma centres. Journal of Emergency Medicine, Trauma and Acute Care 2017(1)	- Study design - not diagnostic accuracy or test and treat
Chew, Brandon G, Swartz, Christopher, Quigley, Matthew R et al. (2013) Cervical spine clearance in the traumatically injured patient: is multidetector CT scanning sufficient alone? Clinical article. Journal of neurosurgery. Spine 19(5): 576-81	 Study design - not diagnostic accuracy or test and treat Study design - all were CT negative to be included
Chilvers, G; Janjua, U; Choudhary, S (2017) Blunt cervical spine injury in adult polytrauma: incidence, injury patterns and predictors of significant ligament injury on CT. Clinical radiology 72(11): 907-914	- Study design - not diagnostic accuracy or test and treat

Study	Code [Reason]
Chiu, Ryan G, Siddiqui, Neha, Rosinski, Clayton L et al. (2020) Effect of Magnetic Resonance Imaging on Surgical Approach and Outcomes in the Management of Subaxial Cervical Fractures. World neurosurgery 138: e169-e176	- Study design - only included those with injury already confirmed either on test or reference standard
	- Study design - not diagnostic accuracy or test and treat
Como, J. J., Leukhardt, W. H., Anderson, J. S. et al. (2011) Computed tomography alone may clear the cervical spine in obtunded blunt trauma patients: a prospective evaluation of a revised protocol. J Trauma 70(2): 345-9; discussion 349	- Study design - all were CT negative to be included
Como, J. J., Thompson, M. A., Anderson, J. S. et al. (2007) Is magnetic resonance imaging essential in clearing the cervical spine in obtunded patients with blunt trauma?. J Trauma 63(3): 544-9	- Full text paper not available
Cui, Li W, Probst, Marc A, Hoffman, Jerome R et al. (2016) Sensitivity of plain radiography for pediatric cervical spine injury. Emergency radiology 23(5): 443-8	- Study design - only included those with injury already confirmed either on test or reference standard
Davies, J; Cross, S; Evanson, J (2016) Radiological assessment of paediatric cervical spine injury in blunt trauma: the potential impact of new NICE guidelines on the use of CT. Clinical radiology 71(9): 844-53	- Diagnostic test not relevant to review protocol
Davis JW, Kaups KL, Cunningham MA et al. (2001) Routine evaluation of the cervical spine in head-injured patients with dynamic fluoroscopy: a reappraisal. The Journal of trauma 50(6): 1044-1047	- Diagnostic test not relevant to review protocol
Diaz JJ, Gillman C, Morris JA et al. (2003) Are five-view plain films of the cervical spine unreliable? A prospective evaluation in blunt trauma patients with altered mental status. The Journal of trauma 55(4): 658	- Reference standard test not relevant to protocol
Diaz, J. J., Jr., Aulino, J. M., Collier, B. et al. (2005) The early work-up for isolated ligamentous injury of the cervical spine: does computed tomography scan have a role?. J Trauma 59(4): 897-903; discussion 903	- Study design - all were CT negative to be included

Study	Code [Reason]
Duane, T. M., Scarcella, N., Cross, J. et al. (2010) Do flexion extension plain films facilitate treatment after trauma?. Am Surg 76(12): 1351- 4	- Reference standard test not relevant to protocol
El Saman, Andre, Laurer, Helmut, Maier, Bernd et al. (2007) Diagnosis, Timing and Treatment of Cervical Spine Injuries in Polytraumatized Patients. European journal of trauma and emergency surgery : official publication of the	- Study design - not diagnostic accuracy or test and treat
European Trauma Society 33(5): 501-11	- Study design - only included those with injury already confirmed either on test or reference standard
Ertel, Audrey E; Robinson, Bryce R H; Eckman, Mark H (2016) Cost-effectiveness of cervical spine clearance interventions with litigation and long-term-care implications in obtunded adult patients following blunt injury. The journal of trauma and acute care surgery 81(5): 897-904	- Study design - not diagnostic accuracy or test and treat
Gamal, G.H. (2014) Evaluation of spinal trauma by multi detector computed tomography and magnetic resonance imaging. Egyptian Journal of Radiology and Nuclear Medicine 45(4): 1209- 1214	- Not specifically cervical spine injury
Garton, H. J. and Hammer, M. R. (2008) Detection of pediatric cervical spine injury. Neurosurgery 62(3): 700-8; discussion 700	- Study design - only included those with injury already confirmed either on test or reference standard
Gerrelts BD, Petersen EU, Mabry J et al. (1991) Delayed diagnosis of cervical spine injuries. The Journal of trauma 31(12): 1622-1626	- Study design - only included those with injury already confirmed either on test or reference standard
Ghasemi, A.; Haddadi, K.; Shad, A.A. (2015) Comparison of diagnostic accuracy of MRI with and without contrast in diagnosis of traumatic	- Not specifically cervical spine injury
spinal cord injuries. Medicine (United States) 94(43): e1942	- Study design - only included those with injury already confirmed either on test or reference standard
Griffith B, Bolton C, Goyal N et al. (2011) Screening cervical spine CT in a level I trauma center: overutilization?. AJR. American journal of roentgenology 197(2): 463-467	- Diagnostic test not relevant to review protocol
Haas, Brian M; Hahn, Lewis D; Oliva, Isabel (2019) What is the added sensitivity of non- lateral cervical spine radiographs in the	- Study design - not diagnostic accuracy or test and treat

Study	Code [Reason]
evaluation of acute cervical spine trauma?. Emergency radiology 26(2): 133-138	
Hale, Andrew T, Alvarado, Abraham, Bey, Amita K et al. (2017) X-ray vs. CT in identifying significant C-spine injuries in the pediatric population. Child's nervous system : ChNS : official journal of the International Society for Pediatric Neurosurgery 33(11): 1977-1983	- Study design - only included those with injury already confirmed either on test or reference standard
Halpern CH, Milby AH, Guo W et al. (2010) Clearance of the cervical spine in clinically unevaluable trauma patients. Spine 35(18): 1721-1728	- Systematic review used as source of primary studies
Hamard, A., Greffier, J., Bastide, S. et al. (2021) Ultra-low-dose CT versus radiographs for minor spine and pelvis trauma: a Bayesian analysis of accuracy. European Radiology 31(4): 2621- 2633	- Not specifically cervical spine injury
Harris, T. J., Blackmore, C. C., Mirza, S. K. et al. (2008) Clearing the cervical spine in obtunded patients. Spine (Phila Pa 1976) 33(14): 1547-53	- Study design - all were CT negative to be included
Hashem, R., Evans, C. C., Farrokhyar, F. et al. (2009) Plain radiography does not add any clinically significant advantage to multidetector row computed tomography in diagnosing cervical spine injuries in blunt trauma patients. J Trauma 66(2): 423-8	- Study design - only included those with injury already confirmed either on test or reference standard
Hennessy, D., Widder, S., Zygun, D. et al. (2010) Cervical spine clearance in obtunded blunt trauma patients: a prospective study. J Trauma 68(3): 576-82	- Study design - all were CT negative to be included
Holmes JF and Akkinepalli R (2005) Computed tomography versus plain radiography to screen for cervical spine injury: a meta-analysis. The Journal of trauma 58(5): 902-905	- Systematic review used as source of primary studies
Holmes, J. F., Mirvis, S. E., Panacek, E. A. et al. (2002) Variability in computed tomography and magnetic resonance imaging in patients with cervical spine injuries. J Trauma 53(3): 524-9; discussion 530	- Study design - only included those with injury already confirmed either on test or reference standard
Huang, Raymond, Ryu, Robert C, Kim, Terrence T et al. (2020) Is magnetic resonance imaging becoming the new computed tomography for	- Study design - not diagnostic accuracy or test and treat

Study	Code [Reason]
cervical spine clearance? Trends in magnetic resonance imaging utilization at a Level I trauma center. The journal of trauma and acute care surgery 89(2): 365-370	
Imerci, A., Canbek, U., Bozoglan, M. et al. (2013) An evaluation of the necessity of computed tomography used for the cervical spine assessment of the patients who present with trauma in the pediatric emergency department. Nobel Medicus 9(2): 91-95	- Study not reported in English
Jeong, SY., Jeon, SJ., Seol, M. et al. (2020) Diagnostic performance of dual-energy computed tomography for detection of acute spinal fractures. Skeletal Radiology 49(10): 1589-1595	- Not specifically cervical spine injury
Kanji, Hussein D, Neitzel, Andrew, Sekhon, Mypinder et al. (2014) Sixty-four-slice computed tomographic scanner to clear traumatic cervical spine injury: systematic review of the literature. Journal of critical care 29(2): 314e9-13	- Systematic review used as source of primary studies
Kanna, Rishi Mugesh, Gaike, Chandrasekar V, Mahesh, Anupama et al. (2016) Multilevel non- contiguous spinal injuries: incidence and patterns based on whole spine MRI. European spine journal : official publication of the European Spine Society, the European Spinal Deformity Society, and the European Section of the Cervical Spine Research Society 25(4): 1163-9	- Study design - not diagnostic accuracy or test and treat
Keller, S, Bieck, K, Karul, M et al. (2015) Lateralized Odontoid in Plain Film Radiography: Sign of Fractures? A Comparison Study with MDCT. RoFo : Fortschritte auf dem Gebiete der Rontgenstrahlen und der Nuklearmedizin 187(9): 801-7	- Insufficient information to calculate diagnostic accuracy measures
Klein, G. R., Vaccaro, A. R., Albert, T. J. et al. (1999) Efficacy of magnetic resonance imaging in the evaluation of posterior cervical spine fractures. Spine (Phila Pa 1976) 24(8): 771-4	- Study design - only included those with injury already confirmed either on test or reference standard
Laham JL, Cotcamp DH, Gibbons PA et al. (1994) Isolated head injuries versus multiple trauma in pediatric patients: do the same indications for cervical spine evaluation apply?. Pediatric neurosurgery 21(4): 221-6; discussion 226	- Study design - not diagnostic accuracy or test and treat

Study	Code [Reason]
Liao, Shiyao, Jung, Matthias K, Hornig, Lukas et al. (2020) Injuries of the upper cervical spine- how can instability be identified?. International orthopaedics 44(7): 1239-1253	- Review article but not a systematic review
Lindholm, Erika B, Malik, Archana, Parikh, Darshan et al. (2019) Single-lateral cervical radiograph in pediatric trauma is equivalent to multiple views. The journal of trauma and acute care surgery 87(4): 813-817	- Study design - only included those with injury already confirmed either on test or reference standard
Liu, Q, Liu, Q, Zhao, J et al. (2015) Early MRI finding in adult spinal cord injury without radiologic abnormalities does not correlate with the neurological outcome: a retrospective study. Spinal cord 53(10): 750-3	 Study design - not diagnostic accuracy or test and treat Study design - all were CT negative to be included
MacDonald, R. L., Schwartz, M. L., Mirich, D. et al. (1990) Diagnosis of cervical spine injury in motor vehicle crash victims: how many X-rays are enough?. J Trauma 30(4): 392-7	- X-ray not currently used as initial imaging test in severely injured population
Makino, Yohsuke, Yokota, Hajime, Nakatani, Eiji et al. (2017) Differences between postmortem CT and autopsy in death investigation of cervical spine injuries. Forensic science international 281: 44-51	- Study design - only included those with injury already confirmed either on test or reference standard
Malhotra, A. and Malhotra, A.K. (2019) Evaluation of Cervical Spine Injuries. Current Trauma Reports 5(1): 48-53	- Review article but not a systematic review
Malhotra, Ajay, Wu, Xiao, Kalra, Vivek B et al. (2017) Utility of MRI for cervical spine clearance after blunt traumatic injury: a meta-analysis. European radiology 27(3): 1148-1160	- Study design - all were CT negative to be included
Martin, Matthew J, Bush, Lisa D, Inaba, Kenji et al. (2017) Cervical spine evaluation and clearance in the intoxicated patient: A prospective Western Trauma Association Multi- Institutional Trial and Survey. The journal of trauma and acute care surgery 83(6): 1032- 1040	- Secondary publication of an included study that does not provide any additional relevant information
Mavros, Michael N, Kaafarani, Haytham M A, Mejaddam, Ali Y et al. (2015) Additional Imaging in Alert Trauma Patients with Cervical Spine Tenderness and a Negative Computed	- Study design - not diagnostic accuracy or test and treat

Study	Code [Reason]
Tomographic Scan: Is it Needed?. World journal of surgery 39(11): 2685-90	- Study design - all were CT negative to be included
McCallum, J., McLaughlin, P., Hameed, M. et al. (2018) 64-Slice CT compared to MRI to clear cervical spine injury in high-risk GCS < 14 blunt trauma patients admitted to the ICU. Trauma	- Study design - not diagnostic accuracy or test and treat
(United Kingdom) 20(1): 38-45	- Study design - all were CT negative to be included
McCutcheon, Lucy, Schmocker, Nicole, Blanksby, Kayla et al. (2015) Best Practice in Diagnostic Imaging after Blunt Force Trauma Injury to the Cervical Spine: A Systematic Review. Journal of medical imaging and radiation sciences 46(2): 231-240	- Systematic review used as source of primary studies
Meinig, H., Doffert, J., Linz, N. et al. (2014) Sensitivity and specificity of ultrasound in spinal trauma in 29 consecutive patients. European Spine Journal	- Diagnostic test not relevant to review protocol
Menaker, J., Philp, A., Boswell, S. et al. (2008) Computed tomography alone for cervical spine clearance in the unreliable patientare we there yet?. J Trauma 64(4): 898-903; discussion 903	- Study design - all were CT negative to be included
Menaker, J., Stein, D. M., Philp, A. S. et al. (2010) 40-slice multidetector CT: is MRI still necessary for cervical spine clearance after blunt trauma?. Am Surg 76(2): 157-63	- Study design - all were CT negative to be included
Merza, Fadia Abdul-Ameer and Lafta, Ghazwan Alwan (2022) The role of computed tomography and Glasgow Coma Scale in detecting spinal injury associated with traumatic brain injuries. Medicine and pharmacy reports 95(2): 158-164	- Reference standard test not relevant to protocol
Moeri, Michael, Rothenfluh, Dominique A, Laux, Christoph J et al. (2020) Cervical spine clearance after blunt trauma: current state of the art. EFORT open reviews 5(4): 253-259	- Review article but not a systematic review
Mohamed, Mohamed A, Majeske, Karl D, Sachwani-Daswani, Gul et al. (2016) Impact of MRI on changing management of the cervical spine in blunt trauma patients with a 'negative'	- Study design - not diagnostic accuracy or test and treat
CT scan. Trauma surgery & acute care open 1(1): e000016	- Study design - all were CT negative to be included

Study	Code [Reason]
Moore, Justin M, Hall, Jonathan, Ditchfield, Michael et al. (2017) Utility of plain radiographs and MRI in cervical spine clearance in symptomatic non-obtunded pediatric patients without high-impact trauma. Child's nervous system : ChNS : official journal of the International Society for Pediatric Neurosurgery 33(2): 249-258	- Population not relevant to this review protocol
Morais, D.F., De Melo Neto, J.S., Meguins, L.C. et al. (2014) Clinical applicability of magnetic resonance imaging in acute spinal cord trauma. European Spine Journal 23(7): 1457-1463	 Study design - not diagnostic accuracy or test and treat Not specifically cervical spine injury
Mower, W. R., Hoffman, J. R., Pollack, C. V., Jr. et al. (2001) Use of plain radiography to screen for cervical spine injuries. Ann Emerg Med 38(1): 1-7	- Study design - only included those with injury already confirmed either on test or reference standard
Murphy, Joshua M; Park, Paul; Patel, Rakesh D (2014) Cost-effectiveness of MRI to assess for posttraumatic ligamentous cervical spine injury. Orthopedics 37(2): e148-52	- Study design - not diagnostic accuracy or test and treat
Nazir, Muhammad, Khan, Shahbaz Ali, Raja, Riaz A et al. (2012) Cervical spinal injuries in moderate to severe head injuries. Journal of Ayub Medical College, Abbottabad : JAMC 24(34): 100-2	- Study design - not diagnostic accuracy or test and treat
Nigrovic, L. E., Rogers, A. J., Adelgais, K. M. et al. (2012) Utility of plain radiographs in detecting traumatic injuries of the cervical spine in children. Pediatr Emerg Care 28(5): 426-32	- Study design - only included those with injury already confirmed either on test or reference standard
Nunn, C., Negus, S., Lawrence, T. et al. (2020) Have changes in computerised tomography guidance positively impacted detection of cervical spine injury in children? A review of the Trauma Audit and Research Network data. Trauma (United Kingdom)	- Study design - not diagnostic accuracy or test and treat
Nuñez DB, Zuluaga A, Fuentes-Bernardo DA et al. (1996) Cervical spine trauma: how much more do we learn by routinely using helical CT?. Radiographics : a review publication of the Radiological Society of North America, Inc 16(6): 1307	- Study design - only included those with injury already confirmed either on test or reference standard

Study	Code [Reason]
O'Boynick, C.P.; Lonergan, T.M.; Place, H.M. (2015) Clearing the C-spine in obtunded trauma patients based on admission CT: A prospective randomized trial. Spine 2015(supplement2): 416	- Full text paper not available
Oh, Jason Jaeseong; Asha, Stephen Edward; Curtis, Kate (2016) Diagnostic accuracy of flexion-extension radiography for the detection of ligamentous cervical spine injury following a normal cervical spine computed tomography. Emergency medicine Australasia : EMA 28(4): 450-5	- Study design - all were CT negative to be included
Ojaghihaghighi, S., Vahdati, S.S., Tarzamani, M.K. et al. (2019) Diagnostic value of bedside ultrasound for detecting cervical spine injuries in patients with severe multiple trauma. Trauma Monthly 24(5): e85199	- Diagnostic test not relevant to review protocol
Onoue, Keita, Farris, Chad, Burley, Hannah et al. (2019) Role of cervical spine MRI in the setting of negative cervical spine CT in blunt trauma: Critical additional information in the setting of clinical findings suggestive of occult injury. Journal of neuroradiology = Journal de neuroradiologie	 Study design - not diagnostic accuracy or test and treat Study design - all were CT negative to be included
Overmann, Kevin M; Robinson, Bryce R H; Eckman, Mark H (2020) Cervical spine evaluation in pediatric trauma: A cost- effectiveness analysis. The American journal of emergency medicine 38(11): 2347-2355	- Study design - not diagnostic accuracy or test and treat
Padayachee, L., Cooper, D. J., Irons, S. et al. (2006) Cervical spine clearance in unconscious traumatic brain injury patients: dynamic flexion- extension fluoroscopy versus computed tomography with three-dimensional reconstruction. J Trauma 60(2): 341-5	- Study design - all were CT negative to be included
Panczykowski, D. M.; Tomycz, N. D.; Okonkwo, D. O. (2011) Comparative effectiveness of using computed tomography alone to exclude cervical spine injuries in obtunded or intubated patients: meta-analysis of 14,327 patients with blunt trauma. J Neurosurg 115(3): 541-9	- Systematic review used as source of primary studies
Patel, M S, Grannum, S, Tariq, A et al. (2013) Are soft tissue measurements on lateral cervical spine X-rays reliable in the assessment of traumatic injuries?. European journal of trauma	- Full text paper not available

Study	Code [Reason]
and emergency surgery : official publication of the European Trauma Society 39(6): 613-8	
Patel, Mayur B, Humble, Stephen S, Cullinane, Daniel C et al. (2015) Cervical spine collar clearance in the obtunded adult blunt trauma patient: a systematic review and practice management guideline from the Eastern Association for the Surgery of Trauma. The journal of trauma and acute care surgery 78(2): 430-41	- Systematic review used as source of primary studies
Plackett, Timothy P, Wright, Franklin, Baldea, Anthony J et al. (2016) Cervical spine clearance when unable to be cleared clinically: a pooled analysis of combined computed tomography	- Systematic review used as source of primary studies
and magnetic resonance imaging. American journal of surgery 211(1): 115-21	- Study design - all were CT negative to be included
Platzer, P., Jaindl, M., Thalhammer, G. et al. (2006) Clearing the cervical spine in critically injured patients: a comprehensive C-spine protocol to avoid unnecessary delays in diagnosis. Eur Spine J 15(12): 1801-10	- Study design - only included those with injury already confirmed either on test or reference standard
Plumb, J. O. and Morris, C. G. (2012) Clinical review: Spinal imaging for the adult obtunded blunt trauma patient: update from 2004. Intensive Care Med 38(5): 752-71	- Systematic review used as source of primary studies
Raja, R.; Arooj, S.; Mahmood, H. (2018) Role of magnetic resonance imaging in acute spinal cord trauma. Pakistan Journal of Medical and Health Sciences 12(3): 925-929	- Study design - not diagnostic accuracy or test and treat
Russin, Jonathan J, Attenello, Frank J, Amar, Arun P et al. (2013) Computed tomography for clearance of cervical spine injury in the unevaluable patient. World neurosurgery 80(34): 405-13	- Systematic review used as source of primary studies
Sanchez, B., Waxman, K., Jones, T. et al. (2005) Cervical spine clearance in blunt trauma: evaluation of a computed tomography-based protocol. J Trauma 59(1): 179-83	- Diagnostic test not relevant to review protocol
Sarani, B., Waring, S., Sonnad, S. et al. (2007) Magnetic resonance imaging is a useful adjunct in the evaluation of the cervical spine of injured patients. J Trauma 63(3): 637-40	- Full text paper not available

Study	Code [Reason]
Satahoo, Shevonne S, Davis, James S, Garcia, George D et al. (2014) Sticking our neck out: is magnetic resonance imaging needed to clear an obtunded patient's cervical spine?. The Journal of surgical research 187(1): 225-9	- Study design - not diagnostic accuracy or test and treat
	- Study design - all were CT negative to be included
Savakus, Jonathan C, Weinberg, Douglas S, Moore, Timothy A et al. (2020) Prevertebral Soft-Tissue Swelling at C7 Is Highly Sensitive for Cervical Spine Ligamentous Injury Study Type: Retrospective Cohort Study. Journal of the American Academy of Orthopaedic Surgeons. Global research & reviews 4(4)	- Diagnostic test not relevant to review protocol
Schenarts PJ, Diaz J, Kaiser C et al. (2001) Prospective comparison of admission computed tomographic scan and plain films of the upper cervical spine in trauma patients with altered mental status. The Journal of trauma 51(4): 663	- Study design - only included those with injury already confirmed either on test or reference standard
Schoenwaelder, M.; Maclaurin, W.; Varma, D. (2009) Assessing potential spinal injury in the intubated multitrauma patient: does MRI add value?. Emerg Radiol 16(2): 129-32	- Study design - all were CT negative to be included
Schoneberg, C, Schweiger, B, Hussmann, B et al. (2013) Diagnosis of cervical spine injuries in children: a systematic review. European journal of trauma and emergency surgery : official publication of the European Trauma Society 39(6): 653-65	- Systematic review used as source of primary studies
Schuster, R., Waxman, K., Sanchez, B. et al. (2005) Magnetic resonance imaging is not needed to clear cervical spines in blunt trauma patients with normal computed tomographic results and no motor deficits. Arch Surg 140(8): 762-6	- Diagnostic test not relevant to review protocol
Shyu, J.Y., Khurana, B., Soto, J.A. et al. (2020) ACR Appropriateness Criteria Major Blunt Trauma. Journal of the American College of Radiology 17(5supplement): 160-s174	- Review article but not a systematic review
Sierink, J C, van Lieshout, W A M, Beenen, L F M et al. (2013) Systematic review of flexion/extension radiography of the cervical spine in trauma patients. European journal of radiology 82(6): 974-81	- Systematic review used as source of primary studies

Study	Code [Reason]
Singh, S., Garg, R., Singh, P. et al. (2015) Diagnosing Cervical Spine Injury in Severe Head Injury: A Case for Replacing Plain Radiography With Computed Tomographic Scan of the Cervical Spine. Indian Journal of Neurotrauma 12(1): 35-40	- Reference standard test not relevant to protocol
Smith, Jackie S (2014) A synthesis of research examining timely removal of cervical collars in the obtunded trauma patient with negative computed tomography: an evidence-based review. Journal of trauma nursing : the official journal of the Society of Trauma Nurses 21(2): 63-7	- Study design - all were CT negative to be included
Soult, M. C., Weireter, L. J., Britt, R. C. et al. (2012) MRI as an adjunct to cervical spine clearance: a utility analysis. Am Surg 78(7): 741- 4	- Study design - all were CT negative to be included
Stassen, N. A., Williams, V. A., Gestring, M. L. et al. (2006) Magnetic resonance imaging in combination with helical computed tomography provides a safe and efficient method of cervical spine clearance in the obtunded trauma patient. J Trauma 60(1): 171-7	- Study design - excluded those positive on initial imaging (X-ray)
Sutherland, M., Bourne, M., McKenney, M. et al. (2021) Utilization of computerized tomography and magnetic resonance imaging for diagnosis of traumatic C-Spine injuries at a level 1 trauma center: A retrospective Cohort analysis. Annals of Medicine & Surgery 68: 102566	- Reference standard test not relevant to protocol (MRI, and was not able to calculate MRI+CT as reference standard)
Tomycz, N. D., Chew, B. G., Chang, Y. F. et al. (2008) MRI is unnecessary to clear the cervical spine in obtunded/comatose trauma patients: the four-year experience of a level I trauma center. J Trauma 64(5): 1258-63	- Study design - all were CT negative to be included
Veiga, Joana Raquel Santos and Mitchell, Kay (2019) Cervical spine clearance in the adult obtunded blunt trauma patient: A systematic review. Intensive & critical care nursing 51: 57- 63	- Systematic review used as source of primary studies
Wu, Xiao, Malhotra, Ajay, Geng, Bertie et al. (2018) Cost-effectiveness of Magnetic Resonance Imaging in Cervical Clearance of Obtunded Blunt Trauma After a Normal Computed Tomographic Finding. JAMA surgery 153(7): 625-632	- Study design - not diagnostic accuracy or test and treat

Study	Code [Reason]
Wu, Xiao, Malhotra, Ajay, Geng, Bertie et al. (2018) Cost-effectiveness of Magnetic Resonance Imaging in Cervical Spine Clearance of Neurologically Intact Patients With Blunt Trauma. Annals of emergency medicine 71(1): 64-73	- Study design - not diagnostic accuracy or test and treat
Yasin, A.; Saeed, U.; Munir, M. (2017) Magnetic Resonance Imaging (MRI) diagnostic accuracy in acute spinal column injuries. Pakistan Journal of Medical and Health Sciences 11(3): 971-972	- Not specifically cervical spine injury
Zhuge, W., Ben-Galim, P., Hipp, J.A. et al. (2015) Efficacy of MRI for Assessment of Spinal Trauma. Journal of Spinal Disorders and Techniques 28(4): 147-151	- Not specifically cervical spine injury

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10 Health Economic studies

- 11 Published health economic studies that met the inclusion criteria (relevant population,
- 12 comparators, economic study design, published 2006 or later and not from non-OECD
- 13 country or USA) but that were excluded following appraisal of applicability and
- 14 methodological quality are listed below. See the health economic protocol for more details.
- 15 None.