This local pathway is an example used in the NICE adoption resource on adoption of high sensitivity troponin for early rule out of NSTEMI. It was not produced for or commissioned by NICE.

Liverpool Acute Chest	Pain Proforma	Aintre	ee University Hospital NHS Foundation Trust						
Date: Time: If using CP proforma simple note 'see CP proforma ' on acute medical proforma Completed by: Lead Authors: Dr Aleem Khand, Consultant cardiologist, Dr Freddy Frost, research Fellow									
PATIENT DETAILS	Contact: Aleem.khan	d@aintree.nhs.uk (office ho	ours) Evidence Base: B.						
Name:	DOB:	OOB: Unit number:							
TIMING & DURATION									
Time onset: Duration: Activity at onset									
CHEST LOW SUSPICION	PAIN CHARACTERISTI MODERATE SUSPICE		HIGH SUSPICION						
Non-central (epigastric, L or R sided)		Central (abo	ove epigastric region)						
Localised chest pain (patient points to		Radiation to	Radiation to neck/jaw or arms						
discreet position in chest)		Worse on e	xertion or emotion						
No radiation	Elements of both	Relieved by	rest or GTN						
Pleuritic element	'High' & 'Low' suspicion		autonomic symptoms:						
'Sharp'/ 'Pins and Needles'		e.g. nausea	, sweating, vomiting						
<5 minutes in duration, non-recurring		'Dull'/ 'Sque	eezing'/ 'Heavy'/ 'tight'						
Other:		≥ 5 minutes	, recurrent						
≥2 in absence of high suspicion chest pain feature- consider differential diagnosis and investigate/ reassure appropriately. Only consider use of CP pathway/ troponin sampling if overall clinical suspicion of ACS remains	Consider initiating Rx for ACS Hstn , ECG and follow CP patl	nway ≥2 in absen ECG monito ACS. Check	Other: ≥2 in absence of low suspicion features: ECG monitoring, Initiate treatment for ACS. Check HsTn T, ECG and follow CP pathway.						
	CV Risk /PMH (please circ								
Diabetes: Type 1, type 2, diet controlled Smoking: Current: Pack years: Ex: > 1year, <1 year Dyslipidaemia: Hypertension: PVD: FH:	PMH: Free Text:								
Obesity: Y/N (wt= kg ,ht= m)									
Previous CABG Year									
MI Year									
PCI Year									
	MEDICATION	l	Allergies						
	OTHER RELEVANT H	ISTORY							
(Social history/Systems Review/P	revious relevant investiga	ntions)							
Chest Pain Proforma (v 1) Valid from: Jan 17 to Jan 20	Uncontrolled copy whe	en printed	Page 1 of 2						

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		0	DSERVATIO	JNS					
HR:	BP:	RR:	Sats:	Tem	р	MEWS:			
		FXAMI	NATION F	NDINGS					
			NATIONT	NDINGS					
INVESTIGATIONS									
ECG:			CXR:						
Presentation troponin				BLOODS: (If indicated) Na: K ⁺ : Urea Creat					
trop	oonin		Na:	K :	Urea	Creat			
			Hb:	WC	2	Platelets			
Time:		ng/l							
	troponin								
	POOL CHEST ATHWAY								
Time:		ng/l							
GRAC	E score			http://www.outcomes- umassmed.org/grace/acs risk/acs risk content.html					
Grace (6 month		uillassi	neu.org/grace/	acs_HSK/a	ics_risk_content.ntim			
mor	tality								
DIAGNOSIS									
1.						nlikely ACS nsider non-ACS			
2.						agnoses (see below)			
					Lik	cely/Probable ACS			
3.						nsider ACS treatment			
MANAGEMENT									
NON-ACS CHEST PAIN DIFFERENTIAL (for guidance)									
	TROPONIN RAISED TROPONIN NORMAL								
Cai	rdiac	Non-cardiac		Cardiac		Non-cardiac			
Myono	ricarditis	Aortic dissection		Pericarditis		Anxiety			
	ry embolus	Chronically raised		nonary emboli	ıs	Musculoskeletal			
	ythmia	troponin eg CKD		y ciliboli		Pneumonia			
	tsubo's	Pneumonia				Pancreatitis			
	Irome					GORD			
•						Cholecystitis			
						Dnoumothoray			

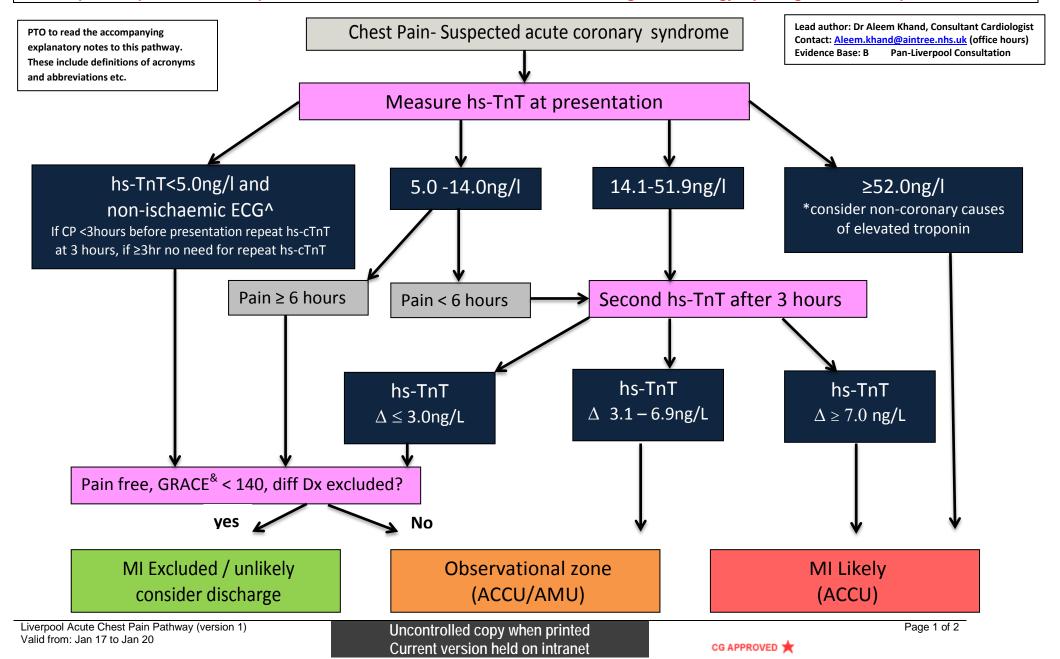
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Liverpool Acute Chest Pain Pathway

All Troponin results must be interpreted in conjunction with clinical presentation and electrocardiograms.

Repeated episodes of chest pain of an ischaemic nature should be referred for urgent cardiology input regardless of Troponin results



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Accompanying Notes for Liverpool Acute Chest Pain Pathway

All patients with STEMI on their ECG should be managed according the primary PCI pathway (with emergency transfer to LHCH if criteria fulfilled http://www.cmcsn.nhs.uk/fileuploads/PPCI Protocol Final April 20131.pdf

This pathway is not a substitute for careful history taking, clinical examination and scrutiny of serial electrocardiograms. In cases where clinician/nurse practitioner judgement differ from pathway directed care then clinician/ nurse practitioner judgement should take precedence.

Any patient with ongoing or recurrent ischaemic sounding chest pain should be referred for urgent cardiology input

Any patient with dynamic or baseline ischaemic changes on their ECG should be referred for urgent cardiology input, irrespective of the initial troponin result.

The pathway should be used for patients presenting with chest pain/discomfort/ possible 'ischaemic' symptoms who are suspected to have acute coronary syndrome. Non-cardiac chest pain should be managed accordingly without an hs-TnT check.

The pink boxes refer to time of arrival at accident and emergency. Therefore, blood samples for hs.TnT should be drawn at **presentation**, **irrespective of time of chest pain**, and (for those mandated by the pathway) at **3 hours after presentation**.

For patients with suspected acute coronary syndrome who present early (<3 hours from the time of [peak] chest pain) a 2nd troponin at 3 hours should be undertaken even if presentation troponin is <5ng/l with a non-ischaemic ECG

This algorithm is **conditional upon the use of a high sensitivity troponin** and is specific to the ROCHE (elecsys) high sensitivity troponin T (hs-cTnT) biomarker analysed with the standard 18minute assay in the COBAS e601/602 analysers.

Renal function: moderate-severe renal dysfunction is associated with troponin elevation by a number of mechanisms. These patients are more likely to be classified in the 'observation zone'. For patients with moderate-severe renal dysfunction in whom an acute coronary syndrome is suspected, using a $\Delta 20\%$ rise or fall maybe more appropriate, rather than an absolute value, in terms of diagnosing myocardial infarction.

Referrals to Rapid Access Chest Pain Clinics/ HOT clinics: In each trust there will be policies for patient referrals for those discharged for further assessment/ cardiac imaging. This service should be utilised only for patients with suspected coronary disease who are deemed safe to be discharged but require further investigation. It should not be a default position for all acute chest pain patients discharged with the aid of this pathway (a large proportion of whom can be reassured with no follow-up or investigations)

Abbreviations/acronyms:

- Hs-TnT= high sensitive (elecsys- ROCHE) troponin T
- ACCU: Acute Cardiovascular Care Unit, AMU: Acute Medical Unit
- Diff Dx: differential diagnosis
- ^ nonischaemic ECG has the following definition: sinus rhythm or atrial fibrillation/ flutter with VR <110, absence of LBBB, absence of ST segment depression or elevation, absence of T wave inversion or t wave flattening in 2 contiguous leads, absence of paced rhythm.
- *For potential non-coronary causes of troponin elevation please see table http://circ.ahajournals.org/content/124/21/2350.full
- pain ≥ or <6 hours refers to the time between chest pain onset/peak and the time of the first Hs-TnT sample.
- Δ refers to the absolute change in troponin values between 1st and 2nd sample (at 3 hours) and can be a **rise or fall**
- All patients in the 'observation zone' should be cared for in the ACCU or the cardiology ward. If they are in A&E or in AMU when the 2nd troponin results indicate that they be categorised in the 'Observation Zone' triage (and thereby require further investigation /clinician input) then they should be transferred to ACCU or the cardiology ward. Care should continue in AMU (or A&E) only in the absence of ACCU or cardiology beds.
- GRACE refers to the GRACE score. (http://www.gracescore.org/website/WebVersion.aspx)

