National Institute for Health and Care Excellence

Draft for consultation

Workplace health: longterm sickness absence and capability to work

Evidence reviews for workplace health: cost effectiveness outcomes

NICE guideline <number>

Evidence reviews

May 2019

Draft for Consultation

These evidence reviews were developed by York Health Economics Consortium





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Contents

Preventing recurring short-term sickness absence (RQ 1a)	6
Review question	6
Introduction	6
PICO table	6
Methods and process	7
Economic evidence	8
Evidence statements	9
The committee's discussion of the evidence	9
Reducing movement from short-term to long-term sickness (RQ 2a)	10
Review question	10
Introduction	10
PICO table	10
Methods and process	11
Economic evidence	11
Summary of studies included in the economic evidence review	13
Evidence statements	16
The committee's discussion of the evidence	16
Helping employees return to work and reducing long-term sickness absence	(RQ
3a)	
Review question	
Introduction	
PICO table	
Methods and process	
Economic evidence	
Summary of studies included in the economic evidence review	
Evidence statements	
The committee's discussion of the evidence	
References	
Appendices	
Appendix A – Review protocols	
Review protocols for review questions 1, 2 and 3	
Appendix B – Literature search strategies	
Appendix G – Economic evidence study selection	54
Figure 1: Flow chart of economic evidence study selection for the guideline	54
54	
Appendix H – Economic evidence tables	
Appendix I – Health economic evidence profiles	83
Appendix J – Health economic analysis	116

DRAFT FOR CONSULTATION

Appendix K – Excluded studies	117
Economic studies	117
Appendix M – Health economic quality assessment	124

Preventing recurring short-term sickness absence (RQ 1a)

3 Review question

- 4 Review question 1a What interventions, programmes, policies or strategies are cost-
- 5 effective in preventing or reducing recurrence of short-term sickness absence among
- 6 employees?

7 Introduction

- 8 Frequent absence may indicate general ill health which requires medical investigation and, if
- 9 continued, may indicate work stress or lack of capability to do the job. Repeated absence for
- short periods is likely both to undermine the individual employee's own performance and
- 11 cause disruption for colleagues and the wider organisation, including:
- the need to find temporary replacement cover (sometimes for quite specialist tasks);
- increasing the workload of others;
- general disruption of the remaining workforce and workflow;
- other employees feeling resentful if they think an individual's repeated absences are not
 being addressed;
- reduction in employee morale;
- the risk that a culture of frequent absenteeism may develop across the wider workforce.

19 PICO table

Table 1: PICO inclusion criteria for interventions to prevent or reduce recurrent shortterm sickness absence

term sickness absence							
Population	 Adult employees (≥16 years; full- or part-time; paid or unpaid) who: have experienced 4 or more episodes of short-term sickness absence in a 12 month period (each episode lasting less than 4 weeks) or are currently absent from work for less than 4 weeks due to sickness (with a minimum study follow-up of 12 months to enable patterns of recurrent absence to be identified) 						
	Organisational level All employers in the public, private and 'not-for-profit' sectors						
Interventions	Any intervention to prevent or reduce recurring short-term sickness absence (4 or more episodes in a 12-month period, each episode lasting <4 weeks). Where interventions are not delivered in a workplace or primary care setting, there should be some element of employer or primary care involvement in the design, content, implementation or funding of the intervention.						
Comparator	 No work-related intervention (includes 'usual care' or usual sickness absence practice / guidance) Any other active comparator for managing sickness absence or return to work Other active workplace comparator (intervention, programme, policy or strategy) Time 						

Outcomes	Effectiveness studies (review question 1a)
	Primary outcome
	Short-term sickness absence, as measured and reported by the authors
	Secondary outcomes
	Health-related quality of life - using validated patient-report measures, for
	example EQ-5DPsychological and/or social functioning - using any patient-report measure
	Adverse / unintended effects:
	- Self-reported presenteeism or work performance (individual-level
	studies); - Job satisfaction (individual or organisational-level)
	- Rate of staff turnover (organisational-level studies)
	- Number of grievances (organisational-level studies)
	Qualitative studies (review question 1b)
	Participant views on:
	Intervention acceptability (including preferences for content, frequency,
	location, etc.)Barriers and facilitators to successful intervention delivery
	- Same and Identatore to Successful Intervention delivery

1 Methods and process

- 2
- This evidence review was developed using the methods and process described in Developing NICE guidelines: the manual. Methods specific to this review question are 3
- described in the review protocol in appendix A. 4
- 5 Declarations of interest were recorded according to NICE's 2014 conflicts of interest policy.

1 Economic evidence

2 Included studies

- 3 8,040 records were assessed against the eligibility criteria.
- 4 7,974 records were excluded based on information in the title and abstract. One reviewer
- 5 assessed all of the records and a second reviewer blind-screened 10% of the records. The
- 6 level of agreement between the two reviewers was 100%.
- 7 The full-text papers of 66 documents were retrieved and assessed and 0 studies were
- 8 assessed as meeting the eligibility criteria for research question 1a. One reviewer assessed
- 9 all of the full texts and a second reviewer blind-screened 10% of the records. The level of
- agreement between the two reviewers was 100%. For review question 1a, no studies were
- 11 included.

12 Excluded studies

- 13 66 full text documents were excluded for this question. The documents and the reasons for
- 14 their exclusion are listed in Appendix K Excluded studies. Documents were excluded for
- the following reasons: ineligible patient population (n=27), ineligible outcomes (n=15),
- ineligible study design (n=12), ineligible intervention (n=5), ineligible outcomes (n=4) and
- ineligible setting (n=3). The selection process is shown in Appendix G.

1 Evidence statements

2 No eligible studies were identified.

3 The committee's discussion of the evidence

4 Cost effectiveness and resource use

No cost-effectiveness studies were identified that met the inclusion criteria for review question 1 – interventions to prevent or reduce recurrent short-term sickness absence.

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A health economic model was developed to determine how cost-effective an intervention will be in helping employees on sickness absence to return to work. Because the interventions and size and type of organisation vary greatly and a myriad of factors can impact sickness absence and return to work, the model adopted a generalised approach and multiple sensitivity analyses were carried out which showed the results varied greatly by key model inputs such as the cost and effectiveness of the intervention, reduction in absenteeism and baseline rate of absenteeism. The committed noted that in general a company with high turnover costs or costs of absenteeism will likely benefit from an intervention to reduce sickness absence, particularly if the intervention is effective and less expensive than the overall costs of absenteeism or replacing a worker. The reverse is also true. For example, an organisation with low baseline turnover costs or low levels of absenteeism will find it more difficult to realise cost savings by implementing an intervention aimed at reducing sickness absence, though this does not mean that other factors could not also benefit the organisation. The committee appreciated employers may be interested in factors other than pure cost savings. The overall willingness to pay for an intervention by an organisation is important: there is no requirement for the intervention to be cost saving if the organisation is willing to pay for an intervention that will benefit the workers and the organisation itself.

Reducing movement from short-term to long-term sickness (RQ 2a)

3 Review question

- 4 Review question 2a What interventions, programmes, policies or strategies are cost
- 5 effective in reducing the number of employees who move from short- to long-term sickness
- 6 absence?

7 Introduction

- 8 There is substantial evidence that work is beneficial for physical and mental health, whereas
- 9 unemployment and long-term sickness absence often have a harmful impact (Marmot and
- Bell 2012). Data have shown that those who had been unemployed for more than six months
- 11 had lower wellbeing than those who had been unemployed for less time (DH 2008).
- 12 Reducing the extent of sickness absence in the UK, and in particular long-term sickness
- 13 absence (defined as a period of four weeks or more) has therefore been a policy priority for
- 14 at least the last ten years.

15 PICO table

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Table 2: PICO inclusion criteria for interventions to reduce movement from short- to long-term sickness absence

Population	Individual level Adult employees (≥16 years; full- or part-time; paid or unpaid) who are currently absent from work for less than 4 consecutive weeks due to sickness. Organisation level All employers in the public, private and 'not-for-profit' sectors
Interventions	Any intervention that aims to reduce the risk of employees progressing from short-term to long-term absence (that is, lasting ≥4 consecutive weeks).
Comparator	 No work-related intervention (includes 'usual care' or usual sickness absence practice / guidance) Any other active comparator for managing sickness absence or return to work Other active workplace comparator (intervention, programme, policy or strategy) Time
Outcomes	Effectiveness studies (review question 2a) Primary outcome Return to work. Measured as any of: - Proportion returning to work within 4 weeks of start of absence - Time taken to return to work or

- Sickness absence, as reported by the authors, including:
 - Proportion with any long-term sickness absence (≥4 consecutive weeks duration)
 - Total number of sickness absence days

Secondary outcomes

- Health-related quality of life using validated patient-report measures, for example EQ-5D
- Psychological and/or social functioning using any patient-report measure
- Adverse / unintended effects:
 - Self-reported 'presenteeism' or work performance (individual-level studies)
 - Job satisfaction (individual or organisational-level)
 - Rate of staff turnover (organisational-level studies)
 - Number of grievances (organisational-level studies)

Qualitative studies (review question 2b)

Participant views on:

- Sickness absence recurrence following RTW (individual-level studies)
- Intervention acceptability (including preferences for content, frequency, location, etc.)
- Barriers and facilitators to successful intervention delivery

1 Methods and process

- 2 This evidence review was developed using the methods and process described in
- 3 Developing NICE guidelines: the manual. Methods specific to this review question are
- 4 described in the review protocol in appendix A.
- 5 Declarations of interest were recorded according to NICE's 2014 conflicts of interest policy.

6 Economic evidence

7 Included studies

- 8 8,040 records were assessed against the eligibility criteria.
- 9 7,974 records were excluded based on information in the title and abstract. One reviewer
- 10 assessed all of the records and a second reviewer blind-screened 10% of the records. The
- 11 level of agreement between the two reviewers was 100%.
- 12 The full-text papers of 66 documents were retrieved and assessed and 1 study was
- 13 assessed as meeting the eligibility criteria for research question 2a. One reviewer assessed
- all of the full texts and a second reviewer blind-screened 10% of the records. The level of
- agreement between the two reviewers was 100%.
- One economic study was eligible for inclusion for review question 2a [1]. This study is
- summarised in the health economic evidence profile in appendix I and the health economic
- evidence tables below in Table 3 and in appendix H.

19 Excluded studies

- 20 65 full text documents were excluded for this question. The documents and the reasons for
- 21 their exclusion are listed in Appendix K Excluded studies. Documents were excluded for
- 22 the following reasons: ineligible patient population (n=27), ineligible outcomes (n=14),

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Reducing movement from short-term to long-term sickness (RQ 2a)

- 1 ineligible study design (n=12), ineligible intervention (n=5), ineligible outcomes (n=4) and
- 2 ineligible setting (n=3). The selection process is shown in Appendix G.

1 Summary of studies included in the economic evidence review

Table 3: Summary of the study included in the economic evidence review for reducing movement from short-term to long-term sickness absence –

3 RQ 2a

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
van Oostrom 2009 (Netherlands) Population: Workers on sick leave for 2 to 8 weeks due to distress Interventions: Workplace intervention (WI): a stepwise communication process to identify and solve obstacles to return to work (RTW).a Comparators: Usual care	Potentially serious limitations °	Partially applicable ^d	None	Mean total costs, over 12 months Societal perspective WI: €3,201 UC: €2,758 Employer perspective WI: €1,386 UC: €802	Mean duration of sick leave, over 12 months Cost Effectivene ss Analysis (CEA) WI: 133 days UC: 134 days Cost utility analysis (CUA) WI: 0.77 UC: 0.78	Mean cost difference, over 12 months Societal perspective WI vs UC: €443 more costly (not statistically significant) Employer perspective WI vs UC: €583 more costly (not statistically significant)	Mean duration of sick leave, over 12 months WI vs UC: 1 day fewer (not statistically significant) QALYs WI vs UC: 0.01 less (not statistically significant)	CEA WI vs UC ICER: €627 per sick day avoided Neither change in costs or change in sick days were statistically different between WI and UC CBA Net monetary benefit with WI was -€1,987 with human capital approach (HCA) and -	The base case was a bootstrapped analysis to account for stochastic uncertainty. Cos effectiveness planes showed substantial uncertainty in results which reflects the statistical uncertainty in the point estimates o cost differences and effectiveness measures between WI and UC. Subgroup analysis

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
(UC): treatment by an occupational physician, according to the Dutch Guidelines. b								€1,700 with friction cost approach (FCA) WI was statistically significantly more costly than UC and changes in costs of productivity loss favoured UC but were not statistically significant regardless of productivity measure. CUA WI vs UC, incremental cost-utility ratio (ICUR) (HCA): -€184,562 per QALY gained (HCA)	suggested WI may be most cost-effective for patients with an intention to return to work but findings were still limited in statistical significance.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
								WI vs UC, ICER (FCA): - €155,850 UC dominates WI but neither cost differences with WI or QALY gains were statistically significant	

CEA: cost-effectiveness analysis; CUA: cost-utility analysis; FCA: frictional cost approach; HCA: human capital approach; ICER: incremental cost-effectiveness ratio; ICUR: incremental cost-utility ratio; RtW: return to work; QALY: quality-adjusted life year; UC: usual care; VAS: visual analogue scale; WI: workplace intervention

- (a) Three meetings were planned to take place within 2 weeks. The purpose of the first meeting between the sick-listed employee and the RTW coordinator was to identify obstacles for RTW from the perspective of the employee. The second meeting was between the supervisor and the RTW coordinator, where obstacles to the employee's RTW were identified from the supervisor's perspective. In the third meeting, which was generally the longest, the employee, supervisor and RTW coordinator discussed solutions and formulated a consensus-based plan for their implementation.
- (b) According to the evidence-based guideline of the Dutch Association of Occupational Physicians (NVAB) published in 2000 and updated in 2007. This guideline aims to facilitate the optimal functioning of employees with mental health problems and to prevent long-term sick leave and frequent recurrences. An early start to the treatment by occupational physicians is recommended. Occupational physicians act as motivating counsellors using cognitive behavioural elements to enhance the problem-solving capacity of employees. In addition, the Improved Gatekeeper Act requires that both the employer and employee take responsibility for a RTW plan.
- (c) Only a 12 month time horizon, so insufficient to capture sustainability of outcomes.
- (d) This study included people with 2 to 4 weeks of sick leave. The study was in the Netherlands where the occupational support offered is differently organised to the UK. EQ-5D VAS, rather than 5 level health state instrument, was used for utility values.

Evidence statements

2 One cost-effectiveness, cost-benefit and cost-utility analysis (van Oostrom, 2009) found that a workplace intervention consisting of a stepwise communication process to identify 3 and solve obstacles to return to work for people absent with distress found that it did not 4 5 improve outcomes and had a higher cost compared to usual care. The workplace intervention for all people was unlikely to be more or less cost-effective than usual care 6 but the workplace intervention was more likely to be cost-effective than usual care in 7 people with an intention to return to work. The analysis was assessed as partially 8 applicable to the review question, with potentially serious limitations. 9

10 The committee's discussion of the evidence

11 Cost effectiveness and resource use

- The cost effectiveness review identified one study which found that a workplace intervention 12 consisting of a stepwise communication process to identify and solve obstacles to return to 13
- work for people absent with distress did not improve outcomes and had a higher cost 14
- compared to usual care. Although the intervention for all people was unlikely to be more or 15
- less cost-effective than usual care the committee were mindful that it was more likely to be 16
- cost-effective than usual care in people with an intention to return to work. However, given 17
- the limitations of the study and the lack of evidence from effectiveness studies the committee 18 19
 - did not consider there to be sufficient evidence to determine the value for money of these
- types of interventions. 20

Helping employees return to work and reducing long-term sickness absence (RQ 3a)

4 Review question

- 5 Review question 3a What interventions, programmes, policies or strategies are cost
- 6 effective in:

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- helping employees on long-term sickness absence to return to work?
 - reducing the recurrence of long-term sickness absence following a return to work?

9 Introduction

- There is substantial evidence that work is beneficial for physical and mental health, whereas
- 11 unemployment and long-term sickness absence often have a harmful impact (Marmot and
- Bell 2012). Data have shown that those who had been unemployed for more than six months
- had lower wellbeing than those who had been unemployed for less time (DH 2008).
- 14 Reducing the extent of sickness absence in the UK, and in particular long-term sickness
- absence (defined as a period of four weeks or more) has therefore been a policy priority for
- 16 at least the last ten years.

17 PICO table

Table 4: PICO inclusion criteria for interventions to help employees on long-term sickness absence return to work and prevent recurrence

Population	Adult employees (≥16 years; full- or part-time; paid or unpaid) who
	 are currently absent from work for 4 or more consecutive weeks due to sickness
	or
	 have returned to work in the past 6 months after an episode of long-term sickness absence (lasting 4 or more consecutive weeks)
	Organisation level
	All employers in the public, private and 'not-for-profit' sectors
Interventions	Any interventions, programmes, policies or strategies that aim to increase the return to work of employees who experience an episode of long-term sickness absence (≥4 consecutive weeks) and / or prevent the recurrence of long-term absence
	Where interventions are not delivered in a workplace or primary care setting, there should be some element of employer or primary care involvement in the design, content, implementation or funding of the intervention.
Comparator	 No work-related intervention (includes 'usual care' or usual sickness absence practice / guidance)
	 Any other active comparator for managing sickness absence or return to work
Outcomes	Effectiveness studies (review question 3a)

Primary outcomes

- Return to work (full / partial). Measured as any of:
 - Proportion returning to work
 - Proportion assessed as capable of returning to work physical or functional assessments using validated or self-report measure, clinical indicators or clinical opinion
 - Time taken to return to work
 - Hours worked per week / month
 - Proportion who take ill-health retirement
- Long-term sickness absence (following the return to work, for those on long-term sickness at baseline) as reported by the authors, including:
 - Proportion with any long-term sickness absence (4 or more weeks duration)
 - Number of episodes of long-term sickness absence (per participant)
 - Number of days sick leave per episode
 - Total number of days sickness absence

Secondary outcomes

- Health-related quality of life using validated patient-report measures, for example EQ-5D
- Psychological and/or social functioning using any patient-report measure
- Adverse / unintended effects:
 - Self-reported 'presenteeism' or work performance (individual-level studies);
 - Job satisfaction (individual or organisational-level)
 - Rate of staff turnover (organisational-level studies)
 - Number of grievances (organisational-level studies)

Qualitative studies (review question 3b)

Participant views on:

- Intervention acceptability (including preferences for content, frequency, location, etc.)
- · Barriers and facilitators to successful intervention delivery

1 Methods and process

- 2 This evidence review was developed using the methods and process described in
- 3 Developing NICE guidelines: the manual. Methods specific to this review question are
- 4 described in the review protocol in appendix A.
- 5 Declarations of interest were recorded according to NICE's 2014 conflicts of interest policy.

6 Economic evidence

7 Included studies

- 8 8,040 records were assessed against the eligibility criteria.
- 9 7,974 records were excluded based on information in the title and abstract. One reviewer
- 10 assessed all of the records and a second reviewer blind-screened 10% of the records. The
- level of agreement between the two reviewers was 100%.
- 12 The full-text papers of 66 documents were retrieved and assessed and 14 studies were
- assessed as meeting the eligibility criteria for research question 3a. One reviewer assessed

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Helping employees return to work and reducing long-term sickness absence (RQ 3a)

- all of the full texts and a second reviewer blind-screened 10% of the records. The level of
- 2 agreement between the two reviewers was 100%.
- 3 14 economic studies met the review inclusion criteria for review question 3a [2-15]. These
- 4 are summarised in the health economic evidence profile in appendix I and the health
- 5 economic evidence table below in Table 4 and in appendix H.

6 Excluded studies

- 7 52 of the 66 full text papers retrieved were excluded on full text review. The documents and
- 8 the reasons for their exclusion are listed in Appendix K Excluded studies. Documents were
- 9 excluded for the following reasons: ineligible patient population (n=27), ineligible study
- design (n=12), ineligible intervention (n=5), ineligible outcomes (n=4), ineligible setting (n=3)
- and ineligible outcomes (n=1). The selection process is shown in Appendix G.

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2 Summary of studies included in the economic evidence review

Table 4: Summary of studies included in the economic evidence review for helping employees return to work and reducing recurrent long-term

4 sickness absence – RQ 3a

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Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Arends 2013 (Netherlands) Population: Workers (aged 18 to 63 years) who were diagnosed at start of sickness absence with a common mental disorder (CMD) and were now partially or fully ready to return to work.	Potentially serious limitations ^c	Partially applicable d	None	Mean cost per person SHARP Cost-effectivene ss analysis (CEA): €4,167 Cost-benefit analysis (CBA): Between €29,337 (human capital approach (HCA) to productivity loss) and €37,215	Recurrent sickness absence over 12 months SHARP: 39% CAU: 62%	SHARP vs CAU ^e CEA: €1,764 more CBA: €4,730 (HCA) CBA: €5,530 (FCA)	Recurrent sickness absence over 12 months, SHARP vs CAU (bootstrappe d estimate): 24% lower	CEA, incremental cost- effectiveness ratio (ICER), SHARP vs CAU, per 1% of recurrent sickness absence prevented: €10,605 CBA Employer occupational health costs only, SHARP was €800 greater than with CAU	CEA Excluding an outlier, which was attributed to high costs due to hospitalisation in a psychiatric ward, an ICER of -€533 was calculated for the incidence of recurrent sickness absence, indicating SHARP could be cost-effective. Reduced SHARP costs did not change the direction of the primary analyses.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
SHARP-at work. A five steps intervention: return to work (RtW) was started with the occupational physician (OP) monitoring and supporting the person through the steps. ^a Comparator: Care as usual (CAU) ^b				(friction cost approach (FCA) to productivity loss) CAU CEA: €2,403 CBA: Between €24,607 (HCA) and €31,685 (FCA)				Productivity loss (HCA) SHARP vs CAU: €6,046 Productivity loss (FCA) SHARP vs CAU: €3,995	CBA Reduced SHARP costs did not change these results
Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Brouwers 2007 (Netherlands) Population: People (aged 18-60 years) absent from work on sick leave for no more than 3 months due to	Potentially serious limitations ^h	Partially applicable ⁱ	None	Mean cost per person IG: €14,493 (exclusive of intervention costs of €13,305 total for all people)	Mean quality- adjusted life years (QALYs) per person not reported Sick leave duration	CBA IG vs CAU (excluding costs of intervention): €11 more expensive (not statistically different)	Mean QALYs per person IG vs CAU (Dutch EQ- 5D): 0.056 higher IG vs CAU (UK EQ-5D): 0.044 higher	CUA, ICER Probabilistic ICERs reported (ICERs are negative). IG vs CAU (Dutch EQ-5D): - €4,179	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA. This demonstrated an even split between the north- east and south-east quadrants of the cost-effectiveness

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
a minor mental disorder Intervention group (IG): 5 individual sessions (50 minutes each), covering 3 stages, with a social worker: cause, coping and implementation . f Comparator: Care as usual (CAU): routine general practitioner (GP) care ^g				CAU: €14,482 CAU CEA: €2,403 CBA: €24,607 (HCA) to €31,685 (FCA)	until full return to work IG: 152.7 days CAU: 156.5 days No statistically significantly difference	Cost-utility analysis (CUA), based on 2,000 bootstrap pairs IG vs CAU: €234 cheaper	Sick leave duration until full return to work IG vs CAU: 3.8 days sooner	IG vs CAU (UK EQ-5D): - €5,306 IG was less expensive and more effective than CAU CBA IG vs CAU: €11 more expensive (not statistically significant but did not include intervention costs)	plane indicating that any difference in costs between IG and CAU were likely small. 52% of bootstrap estimates were in the south east quadrant where IG dominates CAU.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Bultmann 2009 (Denmark) Population: Workers on	Potentially serious limitations ^k	Partially applicable ¹	None	Mean cost per person, over 12 months	Sickness absence, per person, over 12 months	Over 12 months, per person	Mean absence days averted	CEA CTWR vs CCM, per absence day	One way deterministic sensitivity analysis of a doubling of intervention costs

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
sick leave for 4 to 12 weeks due to lower back pain (LBP) or musculoskelet al disorders (MSD) Intervention: Coordinated and tailored work rehabilitation (CTWR), screening followed by a tailored rehabilitation plan developed by an interdisciplinar y team j		Approcuanty		CTWR: \$31,144 (\$3,321 without productivity loss) CCM: \$41,812 (\$1,773 without productivity loss)	CTWR: 656.6 hours CCM: 997.3 hours	CTWR vs CCM: \$10,668 (\$1,548 more over 12 months without productivity loss)	CTWR vs CCM: 46.0	avoided (without productivity loss): \$33.7	and 25% reduction in wages still resulted in cost savings under the CBA.
Comparator: Conventional case management (CCM) provided by the									

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
municipality. No further information given.									
Finnes 2017 (Sweden) Population: Workers (at least 50% whole time equivalent (WTE)) with sickness absence due to anxiety, depression, stress or exhaustion Interventions: Acceptance and Commitment Therapy (ACT): a psychologica I intervention consisting of 6 X 60-	Potentially serious limitations q	Partially applicable ^r	None	12 month costs Healthcare perspective ACT: \$5,507 WDI: \$6,465 ACT+WDI: \$6,141 TAU: \$6,207 Societal perspective ACT: \$14,452 WDI: \$15,649 ACT+WDI: \$17,066 TAU: \$15,593	QALY gains over 12 months ACT: 0.164 WDI: 0.122 ACT+WDI: 0.168 TAU: 0.155	12 months, vs ACT Healthcare perspective WDI: \$958 ACT+WDI: \$634 TAU: \$700 Societal perspective WDI: \$1,197 ACT+WDI: \$2,614 TAU: \$1,141	QALY gains over 12 months vs ACT WDI: -0.042 ACT+WDI: 0.046 TAU: -0.009	Healthcare perspective Both TAU and WDI were dominated by ACT ACT vs baseline: \$33,579 per QALY gained ACT+WDI vs ACT: \$158,500 per QALY gained Societal perspective Both TAU and WDI were dominated by ACT alone.	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA. This showed that for ACT compared to ACT+WDI from both the healthcare and societal perspectives, the percentage of bootstrap iterations were spread roughly equally across all four quadrants, although approximately 60% of iterations in both perspectives had ACT+WDI more costly than ACT and 50% of iterations of ACT+WDI were more effective.

Chudu	Limitations	Annlinghilitu	Other	04-	Effects	Incrementa	Incremental	Cost- effectiveness	Unacostalisti
minute sessions m Workplace dialogue intervention (WDI): three meetings involving the participant plus work supervisor n ACT + WDI: conducted by two different therapists o	Limitations	Applicability	comments	Costs	Enects	I cost	effects	Compared to ACT, ACT+WDI had an ICER of \$30,804 per QALY gained	Uncertainty Scenario analysis showed that using Swedish utility weights (rather than English in the base case) resulted in ACT being the dominant strategy. A second scenario explored the impact of using costs as if the intervention was delivered in a 'regular' setting in which case ACT would no longer dominate TAU but
Comparator: Treatment as usual (TAU) was treatment as planned in a primary care centre or other care facility. ^p									have an ICER of \$71 per QALY gained (healthcare perspective) and ACT+WDI had an ICER of \$286,000 per QALY gained compared to ACT.
Jensen 2005 (Sweden) Population: Blue-collar and service/care	Potentially serious limitations ^u	Partially applicable ^v	None	Mean cost per person, at 3 years: female; male	Working days lost per year: female; male	Incremental cost per person, at 3 years, vs TAU:	Change in working days lost per year pre and post	For women, BM, PT and CBT were all less expensive over 3 years	Not undertaken

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
vorkers (aged						female;	intervention:	vs TAU with	
18 to 60 years)				BM:	Pre	male	female; male	BM having the	
vith non-				€107,703;	intervention			lowest cost per	
pecific back				€130,015	BM: 67; 72	BM: -	BM: 22; 51	person vs CBT	
pain resulting n sick leave					PT: 76; 92	€ 137,509;	PT: 19; 18		
for 1-6 months				PT:	CBT:115;	-€63,224	CBT:-6; -8	BM vs CBT:	
ioi 1-0 months				€189,760;	109		TAU: 17; 6	€50,097 less	
nton (ontions:				€220,268	TAU: 65;	PT:	,	expensive	
nterventions:					80	-€55,452;		BM vs PT:	
Behaviour-				CBT:		€27,029		\$82,057 less	
oriented physiotherapy				€157,800;	Post			expensive	
(PT) s				€199,824	intervention	CBT:		BM vs TAU: €137,509 less	
Cognitive					BM: 99;	-€87,412;		expensive	
behavioural				TAU:	123	€6,585		охропото	
therapy				€245,212;	PT: 95; 110			For men, CBT	
(CBT): to				€193,239	CBT:109;			and PT were	
improve pain					101			both more	
management t					TAU: 82;			expensive	
Behavioural					86			over three	
medicine								years than	
programme (BM):								TAU with BM	
PT+CBT								being less	
								expensive than TAU by	
Comparator:								€63,224	
reatment as								C00,224	
ısual (TAU).									
lo additional									
nterventions									
outside the									

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
normal routines in health care. This was not otherwise described. Lambeek 2010	Potentially	Partially	None	At 12	Days until	At 12	Days until	CEA	Stochastic
(Netherlands) Population: Employed or self-employed workers (aged 18 to 65 years) on full or partial sick leave for 12 weeks to 2 years due to non-specific low back pain (LBP). Intervention: Integrated care (IC), a graded activity protocol at the workplace. w	serious limitations ×	applicable y		months IC: £13,165 (£1,479 direct costs, £11,686 indirect costs) UC: £18,475 (£1,262 direct costs, £17,213 indirect costs)	sustainable return to work IC: 129 UC: 197 QALYs IC: 0.74 UC: 0.65	months IC vs UC: - £5,310 (£217 direct costs, - £5,527 indirect costs)	sustainable return to work, per person IC vs UC: -68	£3 extra cost for every day earlier return to work with IC. Direct costs only considered. CUA IC vs UC: dominant IC cost saving over UC, per person: £5,310 QALY gain, per person: 0.09 (direct and indirect costs	uncertainty in the data were dealt with using nonparametric bootstraps for CUA and CEA. This showed that in 98% of iterations IC dominated UC, for the CUA and for the CEA that if there was a willingness to pay of £10 for one day earlier return to work there was a 95% chance that IC was cost-effective. Scenario analysis showed that if only complete cases were considered then there was no
usual care								considered)	statistical difference

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
(UC): referred to occupational physician and GP with a letter containing the advice to treat them according to the Dutch guidelines for patients with LBP.								CBA IC, return on investment for every £1 spent: £26 (direct and indirect costs considered)	in costs between IC and UC. A further scenario explored the impact of the intervention only for people aged under 55 years which resulted in a doubling in the costs of IC. A final scenario analysis showed that varying productivity levels did not impact on results.
Loisel 2002 (Canada) Population: Workers absent for more than 4 weeks with occupational back pain	Minor limitations ^{aa}	Partially applicable bb	The study had fewer than 30 people in each study arm and no statistical significance testing of results was performed	At 12 months OI: \$9,569 CRI: \$12,038 Sherbrooke model: \$12,137 UC: \$9,789	Mean number of days on full benefits (DFB) At 12 months OI: 116.1 CI: 114.9	Incremental costs vs UC At 12 months OI: -\$220 CI: \$2,250 Sherbrooke model:	Mean number of DFB, incremental difference vs UC At 12 months OI: -10.8 CI: -12	CEA (cost per DFB saved vs UC) At 12 months OI: -\$20.40 (dominated UC) CI: \$187.40	Sensitivity analyses were performed by varying the total healthcare costs by 60% to 190% and income per capita by 85% to 125%. Over a 6.4 year time period all interventions remained dominant

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
Interventions:				At 6.4	Sherbrooke	-\$2,348	Sherbrooke	Sherbrooke	vs UC over the cost
 Occupationa 				years	model:		model: -11	model:	ranges considered.
I intervention					115.9	At 6.4		\$213.50	
(OI): visits to				OI: \$16,252	UC: 126.9	years	At 6.4 years		
the study				CI: \$16,902				At 6.4 years	
occupational				Sherbrooke	At 6.4	OI:	OI: -190.3		
medicine				model:	years	-\$16,827	CI: -239.6	OI: -\$88.40	
physician and a				\$14,494		CI:	Sherbrooke	(dominated	
participatory				UC:	OI: 228.0	-\$16,176	model:	UC)	
ergonomics				\$33,079	CI: 178.7	Sherbrooke	-293.7	CI: -\$67.50	
intervention					Sherbrooke	model:	UC: 418.3	(dominated	
with the					model:	-\$18,585		UC)	
study					125.6	ψ.ο,σοσ		Sherbrooke	
ergonomist,					UC: 418.3			model: -\$63.50 (dominated	
the injured								UC)	
worker, their supervisor,								33)	
and								СВА	
managemen								Cost	
t and union								differential vs	
representativ								UC	
es.									
Clinical								At 12 months	
rehabilitation								OI: \$220	
intervention (CRI), with a								CI: -\$2,250	
back pain								Sherbrooke	
specialist								model: -\$2,348	
and								ποασι. ψ2,040	
potentially a								At 6.4 years	
multidisciplin								At 6.4 years	

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
ary work rehabilitation intervention at 12 weeks of absence ^z • OI+CRI ("Sherbrook e Model")		Дериодоні		000.0				OI: \$16,827 CI: \$16,176 Sherbrooke model: \$18,585	Checkumy
Comparator: Usual care (UC): with worker's physician receiving no advice about return to work.									
Meijer 2006 (Netherlands) Population: Bank and university workers on at least 50% contracts with 50% sick leave in the last 4 to 20 weeks due to non-specific upper	Potentially serious limitations ^{dd}	Partially applicable ee	None	Total costs per week At 2 months MDT: €1,335 UC: €448 At 6 months	Percentage of regular hours worked Baseline MDT: 29% UC: 29% 6 months MDT: 82%	Incremental costs of MDT vs UC per week At 2 months: €887 At 6 months: €305	Difference in percentage of regular hours worked Baseline: 0% 6 months: 10% 12 months: 13% There was no statistical difference	MDT was more expensive compared to UC and did not increase the proportion of days worked. MDT was not cost-effective.	Not undertaken

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
extremity musculoskelet al disorders (MSDs) Intervention: Multidisciplinar y treatment (MDT): an outpatient training programme carried out at Dutch rehabilitation centres. cc		, фризавису		MDT: €664 UC: €359 At 12 months MDT: €430 UC: €315	UC: 72% 12 months MDT: 86% UC: 73% There was no statistical difference between MDT and UC at any time point	At 12 months: €115	between MDT and UC at any time point		
Comparator: Usual care (UC): supervision by occupational health services. UC could include treatment at the workplace and in the regular health care system, initiated by a									

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
GP, or medical specialist.									
Radford 2012 (UK) Population: Patients in paid or voluntary work or in full time education, hospitalised for at least 48 hours due to traumatic brain injury (TBI) Intervention: TBI Vocational Rehabilitation (TBI-VR), provided by an occupational therapist ff Comparators: Usual care (UC): participants in hospitals	Potentially serious limitations hh	Partially applicable ii	None	At 12 months Health and social care perspective TBI-VR: £2,106.94 UC: £2,031.71 Societal perspective TBI-VR: £8,786 UC: £10,648	QALYs (at 12 months) TBI-VR: 0.1938 UC: 0.1763 Return to work or education (at 12 months) TBI-VR: 75% UC: 60%	At 12 months Health and social care perspective TBI-VR vs UC: £76.24 more expensive Societal perspective TBI-VR vs UC: £1,867 less expensive	QALYs (at 12 months) TBI-VR vs UC: 0.0175 more QALYs Return to work or education (at 12 months) TBI-VR vs UC: 15% more people returned to work	CEA, per person returned to work, TBI-VR vs UC Health and social care perspective: £501.33 Societal perspective: TBI-VR was more effective and saved money CBA Health and social care perspective: TBI-VR was £75.23 more costly vs UC	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA. The boot strapped ICER was £2,567 lower than the deterministic ICER. Sensitivity analysis using imputed data for missing values more than doubled the cost per person returned to work in the CEA and increased the ICER per QALY gained in the CUA to £35,873 with TBI-VR.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Study without TBI- VR. ⁹⁹	Limitations	Applicability	comments	Costs	Effects	I cost	effects	Societal perspective: TBI-VR was £1,863 less expensive than UC. Neither difference was statistically significant CUA, TBI-VR vs UC From a health and social care perspective: £4,299 per QALY gained Neither the QALY gain nor	Uncertainty
								cost difference was statistically significant	
Rebergen 2009 (Netherlands)	Potentially serious limitations kk	Partially applicable ^{II}	None	At 12 months	Days of sick leave	At 12 months	Days of sick leave, at 12 months	CEA GBC vs UC, ICER per sick	Stochastic uncertainty in the data were dealt with using

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
Population: Police workers on sick leave due to common mental disorders (CMDs) Intervention: Guideline based care (GBC), treatment by occupational physicians (OPs) according to the Dutch guideline for workers with mental health problems. Comparator: Usual care (UC): minimal involvement of the OP and easy access to				Health care perspective GBC: €2,145 UC: €2,664 Societal perspective (HCA for productivity loss) TBI-VR: €14,114 UC: €14,202 No costs were statistically significantly different between groups	GBC: 113 UC: 114 These were not statistically significantly different	Health care perspective GBC vs UC: €520 less expensive	GBC vs UC: 1 fewer The difference was not statistically significant	day avoided: - €736 CBA Estimated net monitory benefit of GBC, per person: €3,582 Outcomes are similar between GBC and UC, but direct costs were lower with GBC. The authors concluded that GBC could be cost-effective.	nonparametric bootstraps for CEA. The iterations showed there was never more than a 50% chance of GBC being cost- effective per day of sick leave avoided regardless of the value of the day of work lost. Different approaches to measuring productivity loss were analysed but did not affect the main findings.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
counselling by a psychologist.	Limitations	Applicability	comments	Costs	Ellects	TCOST	enects	enectiveness	Uncertainty
Schene 2007 (Netherlands) Population: Workers (aged 18 years or over) with at least 50% absence over 10 weeks to 2 years due to work related major depressive disorder (WRMDD)	Potentially serious limitations ⁿⁿ	Partially applicable ^{oo}	None	At 12 months (cost of intervention only) TAU+OT: \$3,149 TAU: \$1,891	No health or employmen t outcomes were reported beyond earnings over 12 month period	TAU+OT vs TAU at 12 months (cost of intervention only): \$1,258 more expensive (Not statistically significant)	Not applicable	Difference in total earnings minus costs of intervention at 12 months TAU+OT vs TAU: \$3,952 higher (Not statistically significant)	The base case was a bootstrapped analysis to account for stochastic uncertainty. The only sensitivity analysis performed was on the value of an hour's work. As the value reduces the probability that TAU+OT is more cost-effective than TAU falls. In the base case it is 75.5% at \$36.88
Intervention: Treatment as usual (TAU) + occupational therapy (OT): TAU was as described below (comparator). OT was the addition of OT, which had									per hour and falls to 52.5% at \$10 per hour

0. 1		A P 1 704	Other	0.11	Ecc. 4	Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
diagnostic and									
therapeutic									
phases. mm									
0									
Comparator:									
Treatment as									
usual (TAU):									
out-patient psychiatric									
treatment for									
depression									
according to									
American									
Psychiatric									
Association									
(APA)									
guideline and									
antidepressant									
s and/or									
cognitive									
behavioural									
therapy (CBT) with senior									
psychiatric									
residents.									
Visits lasted 30									
minutes every									
2–3 weeks.									

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Squires 2012 (UK) Population: Workers on sick leave for 1 week to 6 months due to musculoskelet al disorders (MSDs) Interventions (synthesized evidence identified through a review): • Workplace interventions (WI): a workplace assessment and work modification s based on participative ergonomics involving all relevant	Potentially serious limitations ss	Partially applicable tt	Costs and effects data were very poorly reported. Results were presented on a cost-effectiveness plane and not in a detailed table or text.	Not reported	Increased likelihood of return to work (i.e. relative risk) within the first 6 months of sickness absence (obtained from a literature review) WI: 1.12 PAE: 1.06 PAEW: 1.43	Not reported	Not reported	CUA (from societal perspective so includes costs to NHS and from lost wages) WI and PAEW are both cheaper than UC and more effective. PAE is more costly but more effective than UC. PAEW dominates all interventions. CEA, cost per sick day avoided PAEW is the dominant strategy	Sensitivity and scenario analyses were undertaken. PAEW was not dominant if only the employer perspective was taken and the probability of sick leave recurring was doubled. In a threshold analysis, if the intervention costs were less than an additional £3,000 and returns at least an additional 3% of people to work (32/1,000) in comparison to UC, then it is likely to result in a cost per QALY gained below £20,000.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
stakeholders		- Applications						111111111111111111111111111111111111111	, , , , , , , , , , , , , , , , , , , ,
pp									
 Physical activity and 									
education									
intervention									
(PAE): any									
form of physical									
activity and									
education									
around how to deal with									
pain and									
body									
mechanics									
 Physical activity, 									
education									
and									
workplace visit									
(PAEW):									
WI+PAE									
plus a									
workplace visit by the									
employee									
and the									
physical									
therapist to inform									

			Other			Incrementa	Incremental	Cost-	
rehabilitation . qq Comparator: Usual care (UC) treatment of MSDs in the UK "	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
Steenstra 2006 (Netherlands) Population: Workers on sick leave for 2 to 6 weeks due to low back pain (LBP) The study had a 2-stage design. Between 2 to 8 weeks of sick-leave, patients received either the workplace intervention (WI) or usual care (UC). After 8 weeks, approximately	Potentially serious limitations **	Partially applicable yy	None	Mean total costs WI: €8,993 UC: €9,109 UC+CI: €10,537 UC+UC: €10,885 WI+CI: €12,391 WI+WI: €11,096	Actual QALY values not reported Sick leave (calendar days) WI: 108.5 UC: 135.2 UC+CI: 172.9 UC+UC: 155.9 WI+CI: 181.7 WI+WI: 115.3	Vs UC over 12 months WI: €16 UC+CI: €1,428 WI+CI: €3,282 None of the differences were statistically significant	Incremental QALY values were not reported Sick leave vs UC, calendar days WI: -26.7 days UC+UC: 20.7 days WI+CI: 46.5 days	CEA (per 1 day less of sick leave) WI vs UC: €19 WI+CI vs UC: €11 UC+CI vs UC: €29 CUA (cost per QALY) WI vs UC: -€1,483 WI+CI vs UC: €24,416 UC+CI vs UC: €5,447	The base case was a bootstrapped analysis to account for stochastic uncertainty. The cost-effective planes (and confidence intervals of point estimates) suggested that WI and UC were likely similar in cost but that WI was more effective. CI is likely less effective and more costly than both WI and UC. Scenario analyses suggested using a fixed sum per day

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
half of each									of production lost,
group also									net rather than
received a									calendar sick days
clinical intervention									and using a HCA approach to
(CI).									productivity loss did
(01).									not significantly
Interventions:									influence results.
 WI: usual 									
care, a									
workplace									
assessment									
and									
modification									
and communicati									
on between									
OP and GP									
in order to									
discuss how									
to counsel									
the worker to									
RtW ^{uu} • Clinical									
intervention									
(CI): a									
graded									
activity									
programme									
of 26 x 1-									
hour _.									
sessions,									

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
with a frequency of 2 sessions per week ** Comparator: Usual Care (UC): Dutch OP guidelines for LBP delivered by a GP. ***									
Uegaki 2010 (Netherlands) Population: Workers with partial sick leave over 3 months due to distress Intervention: Minimal intervention for stress-related mental disorders with sick leave (MISS): a GP	Potentially serious limitations ^{aaa}	Partially applicable bbb	None	At 12 months MISS: €12,538 UC: €12,722	QALYs over 12 months MISS: 0.78 UC: 0.76	MISS vs UC, incremental cost at 12 months: A saving of €184 This was not statistically significant	QALY gain, over 12 months MISS vs UC: 0.02 QALYs This was not statistically significant	ICER MISS vs UC: -€7,356 per QALY gained Neither change in costs nor change in QALYs were statistically significantly different between MISS and UC	The base case was a bootstrapped analysis to account for stochastic uncertainty. Cost-effectiveness planes showed that in the base case 77% of bootstrapped pairs would be considered cost-effective at a willingness to pay (WTP) threshold of €25,600 per QALY.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
customized version of an activating approach zz Comparator: usual care (UC) managed by a GP. No									Sensitivity analysis explored different approaches to costing lost productivity but did not significantly influence the results.
further information given.									Subgroup analysis suggested MISS may be most costeffective for patients with stress related mental disorders, which was the only analysis which had statistically significant improvement in QALYs vs UC: -€28,278

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty

ACT: acceptance and commitment therapy; APA: American Psychiatric Association; BM: behavioural medicine; CAU: care as usual; CBA: cost-benefit analysis; CBT: cognitive behavioural therapy; CCM: conventional case management; CEA: cost-effectiveness analysis; CG: control group; CI: clinical intervention; CMD: common mental disorder; CRI: clinical rehabilitation intervention; CTWR: coordinated and tailored work rehabilitation; CUA: cost-utility analysis; DFB: days on full benefits; FCA: frictional cost approach; GBC: guideline based care; GP: general practitioner; HCA: human capital approach; IC: Integrated care; ICER: incremental cost-effectiveness ratio; IG: intervention group; LBP: lower back pain; MDT: multidisciplinary treatment; MISS: minimal Intervention for stress-related mental disorders with sick leave; MSD: musculoskeletal disorders; MSK: musculoskeletal disorders; NA: not applicable; OI: occupational intervention; OP: occupational physician; OT: occupational therapy; PAE: physical activity and education intervention; PAEW: physical activity, education and workplace visit; PSA: probabilistic sensitivity analysis; PT: physiotherapy; QALY: quality-adjusted life year; RtW: return to work; TAU: treatment as usual; TBI: traumatic brain injury; TBI-VR: traumatic brain injury vocational rehabilitation; UC: usual care; VAS: visual analogue scale; VR: vocational rehabilitation; WDI: workplace dialogue intervention; WI: workplace intervention; WRMDD: work related major depressive disorder; WTE: whole time equivalent; WTP: willingness to pay

- (a) The five steps comprised: (1) making an inventory of problems and/or opportunities encountered at work after RtW; (2) brainstorming about solutions/realisations; (3) writing down solutions/ realisations and the support needed and assessing the applicability of these solutions; (4) discussing solutions/ realisations and making an action plan with the supervisor; (5) evaluating the action plan/implementation of solutions.
- (b) Occupational physicians (OPs) enacted the guideline of the Netherlands Society of Occupational Medicine "The treatment of workers with mental health problems by the OP". It is primarily aimed at structuring OPs' treatment to help sick-listed workers with mental health issues to RtW. Limited focus is given to follow-up after RtW has been achieved: only one consultation, to address relapse.
- (c) A 12 month time horizon was used, which is insufficient to capture whether outcomes are sustained. Impact on QALYs was not considered.
- (d) It was unclear how long people in the study had been absent from work. The study was in the Netherlands where the occupational support offered is differently organised than in the UK.
- (e) Calculated from the total costs reported in the paper.
- (f) Treatment was over 10 weeks and entailed 3 stages: (i) understanding the cause of loss of control (ii) the development of problem-solving strategies; and (iii) their implementation.
- (g) This could include medication, counselling or referral.
- (h) An 18 month time horizon was used, which is insufficient to capture whether outcomes were sustained. Deterministic sensitivity analysis was not performed.
- (i) People in the study could have been absent for less than four weeks. The study was in the Netherlands where the occupational support offered is differently organised than the UK. It is unclear how utility values were derived.
- (j) After 4 to 12 weeks of sick leave: (1) work disability screening was conducted: a multidisciplinary assessment of disability and functioning and identification of barriers for RtW and (2) a coordinated, tailored and action-oriented work rehabilitation plan was developed by an interdisciplinary team with continuous feedback on the plan from the sick listed worker, the interdisciplinary team, the workplace, and major stakeholders. The interdisciplinary team consisted of an occupational physician, an occupational physician, an occupational physician, a chiropractor, a psychologist, and a social worker who had the role of case worker.

			Other			Incrementa	Incremental	Cost-		
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty	

- (k) A 12 month time horizon was used which is insufficient to capture whether outcomes are sustained. Impact on QALYs was not considered. Probabilistic sensitivity analysis (PSA) was not undertaken.
- (I) The study was in Denmark where the organisation of the sickness benefits system is similar to the UK, but may still be different enough to limit the generalisability of findings.
- (m) The first part (sessions 1 to 3) emphasized mindfulness, cognitive defusion, and acceptance. During sessions 4 to 6, the focus was on exploring and clarifying personal values and committing to pursuing valued life activities.
- (n) The first step was an individual interview with the participant at the clinic followed by an interview with the participant's supervisor at the workplace. These meetings, lasting up to 60 minutes, aimed to investigate the participants' and the supervisors' views upon causes of the sickness absence, and what might facilitate RTW.
- (o) There was no integration or coordination of the two interventions, and no interaction between therapists.
- (p) This typically included psychotherapy, cognitive behavioural therapy and/or pharmacological treatments, physical therapy and counselling.
- (q) A 12 month time horizon was used which is insufficient to capture sustainability of outcomes. Medication costs were not considered.
- (r) It was unclear how much time people in the study had been absent from work and the study was in Sweden where the occupational support offered is differently organised than in the UK. Drug costs were not considered despite a societal perspective.
- (s) This consisted of approximately 20 scheduled hours per week aimed at enhancing the physical functioning and facilitating lasting behaviour change. Each participant was assigned to an individually tailored training programme.
- (t) This was an average of 13 to 14 scheduled hours per week aimed at improving participants' ability to manage their pain and resume a normal level of activity.
- (u) A time horizon of 3 years was used which is only partly sufficient to assess whether outcomes are sustainable over the long-term. No deterministic sensitivity analysis or probabilistic sensitivity analysis was performed. Cost sources were poorly reported.
- (v) The study was conducted in Sweden where the organisation of sickness benefits system is similar to the UK, but may still be different enough to limit the generalisability of the findings.
- (w) The workplace intervention protocol formulated a consensus-based plan for adaptations at work to facilitate return to work. The integrated care team consisted of a medical specialist, occupational therapist, physiotherapist, and clinical occupational physician.
- (x) Only a 12 month time horizon was used which is insufficient to capture whether outcomes are sustained.
- (y) The study includes people on partial sick leave so may not be considered 'continuous absence'. The study was set in the Netherlands where the organisation of healthcare and sickness may be different enough from the UK to limit the generalisability of findings.
- (z) Clinical examination by a back pain medical specialist, participation in a back school after eight weeks of absence from regular work and, if necessary, a multidisciplinary work rehabilitation intervention after 12 weeks of absence from work.
- (aa) Impact on QALYs was not considered and no probabilistic sensitivity analysis was reported.

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty

- (bb) The study was conducted in Canada where the organisation of the sickness benefits system is similar to the UK, but where it may still be different enough to limit the generalisability of findings. The perspective included the employment insurer.
- (cc) The main part of the intervention took 13 full days, 5 return-to-work sessions and 1 feedback session, all of which took place within 2 months. Each day's schedule consisted of four (1.5 hours) sessions: two physical sessions and two psychological sessions, twice a week supplemented with a fifth session consisting of 30 minutes of relaxation exercises.
- (dd) The study only had a 12 month time horizon which is insufficient to assess whether outcomes are sustained. Impact on QALYs was not considered. No deterministic or PSA was performed.
- (ee) The population includes people with 50% to 100% sick leave. The study was set in the Netherlands where the occupational support offered is differently organised to the UK.
- (ff) Vocational rehabilitation involved: assessing the impact of TBI on the participant, family and their roles; community reintegration training; pre-work training; liaison with employers, tutors or employment advisors.
- (gg) Local differences in service provision meant that this varied widely between participants, but potentially involved support from Headway (a voluntary organization providing advice and support to TBI people and their families), community occupational therapy (OT) or physiotherapy and routine GP follow-up.
- (hh) A 12 month time horizon was used which was insufficient to capture whether outcomes were sustained. QALY data were estimated using VAS. Effectiveness data were not derived from a RCT.
- (ii) Although this was a UK study, the population included students and those in unpaid employment. EQ-5D VAS data were used for utilities rather than health states valued by a population.
- (jj) The course focused on an early start of the intervention by OPs, in which they operated as an activating counsellor using CBT to enhance the problem-solving capacity of workers, especially in relation to their work environment. This consisted of clinical management according to the APA Guideline (2000) and antidepressants and/or CBT with senior psychiatric residents. Visits lasted 30 minutes every 2 to 3 weeks.
- (kk) A 12 month time horizon was used which was insufficient to capture whether outcomes were sustained. QALYs were not considered and neither were all healthcare costs.
- (II) The length of unemployment was unclear for study participants. The study was conducted in the Netherlands where occupational support is differently organised to the UK.
- (mm) TAU included antidepressants, if indicated and accepted by patients. They were treated by senior psychiatric residents with visits lasting 30 minutes every 2 to 3 weeks. OT consisted of two skilled occupational therapists providing the intervention over three manual-based phases: diagnostic phase (4 weeks) five contacts with an occupational physician from the patient's employer and a plan for work reintegration; therapeutic phase (24 weeks) 24 weekly group sessions (8 to 10 patients) and 12 individual sessions; follow-up phase (20 weeks) three individual visits.
- (nn) A 12 month time horizon was used which is insufficient to capture whether outcomes were sustained. QALYs were not considered. The source of costs was unclear.

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty

- (oo) The population includes people with 50% to 100% sick leave. The study was set in the Netherlands where the occupational support offered is differently organised to the UK.
- (pp) Work modifications were defined as those based on participative ergonomics involving all relevant stakeholders.
- (qq) This enabled the employer to become actively involved in the rehabilitation process. The PAEW intervention did not include a workplace assessment and work modifications, as was part of the WI.
- (rr) UC included 4.5 GP visits, 4.5 prescriptions, 3 packs of pain relief medication, 4 half hour sessions of physiotherapy (in 7% of cases), 2.5 sessions of osteopathy (in 5% of cases), 2.5 sessions of chiropractic treatment (in 2% of cases) and a hospital outpatient visit (in 10% of cases).
- (ss) Whilst a lifetime horizon was used in the model it was based on only 12 months of effectiveness data with outcomes not influenced by the intervention after 12 months. Costs, QALYs and incremental analysis were not reported and PSA was not performed.
- (tt) Although this is a UK study, it is based upon effectiveness studies that were conducted outside the UK and the authors stated that this fact may limit study generalisability. Costs and outcomes data were also not well reported.
- (uu) WI started at baseline, at least 8 weeks before sick-leave. The intervention consisted of: Dutch OP guidelines for LBP; A workplace assessment and work modifications based on participative ergonomics, which involved all important stakeholders: the occupational health service's ergonomist or occupational health nurse, the worker on sick-leave, the worker's supervisor and other communication between the OP and the GP, to reach consensus on counselling the worker in RTW.
- (vv) A graded activity programme based on operant behavioural therapy principles based on the findings from patient history, physical examination, functional capacity evaluation, the demands from the patients' work and the patients' expectations on time to RTW. The entire programme consisted of a maximum of 26 one-hour sessions, with a frequency of two sessions a week. The first session took half an hour more since taking the patients' history and a physical examination were part of this session. The programme ended as soon as a full RTW had been established, according to an earlier agreed upon individual schedule. During the programme the worker had an active role in RTW and the physiotherapist acted as a coach and supervisor, using a hands-off approach
- (ww)This included resuming daily activities, working within two weeks was encouraged and a clinical intervention recommended after 12 weeks.
- (xx) A 12 month time horizon was used which is insufficient to capture whether outcomes were sustained. QALYs were estimated using VAS.
- (yy) This study included people with 2 to 4 weeks of sick leave. The study was in the Netherlands where the occupational support offered is differently organised than the UK. EQ-5D VAS rather than 5 level health state was used for utility values.
- (zz) This was developed on the basis of three consultations over a time span of four weeks, and encompassed the following five key tasks: 1 diagnosing stress-related mental disorders; 2 providing education about the problem and importance of taking an active role in one's functional recovery; 3 advising patients on how to reflect, cope and problem-solve; 4 monitoring progress; 5 referring to specialists.
- (aaa) A 12 month time horizon was used which is insufficient to capture the sustainability of outcomes.
- (bbb) Participants were those with partial sick leave over six months. The study was in the Netherlands where the occupational support offered is differently organised than in the UK.

Evidence statements

- One cost-effectiveness and cost-benefit analysis (Arends 2013), in a sensitivity analysis
 that excluded an outlier, found that a five-step return to work programme for people with
 common mental disorders could be cost-effective compared to care as usual. The
 analysis was assessed as partially applicable to the review question, with potentially
 serious limitations.
- One cost-utility and cost-benefit analysis (Brouwers, 2007) found that an activating
 intervention by social workers with people absent from work with distress or minor mental
 disorders reduced sick leave duration and increased QALYs with a reduction in costs with
 a negative deterministic ICER indicating that the intervention dominated usual care. The
 analysis was assessed as partially applicable to the review question, with potentially
 serious limitations.
- One cost-effectiveness and cost-benefit analysis (Bultmann, 2009) found that work disability screening using a multidisciplinary assessment of disability and functioning and identification of barriers for return to work followed by a coordinated, tailored and action-oriented work rehabilitation plan developed for workers on sick leave for 4–12 weeks due to muskuloskeletal disorders had fewer sickness absence hours than controls receiving usual care. The economic evaluation showed that coordinated and tailored work rehabilitation seems to be cost saving for society. The analysis was assessed as partially applicable to the review question, with potentially serious limitations.
- One cost-utility analysis (Finnes, 2017) found that a psychological intervention either
 alone or in combination with a work place intervention for people on sickness absence
 due to mental disorders was likely to be cost-effective compared to treatment as usual.
 However, neither improvements in quality-adjusted life-years (QALYs) or differences in
 cost with the interventions were statistically significant. The analysis was assessed as
 partially applicable to the review question, with potentially serious limitations.
- One cost-benefit analysis (Jensen, 2005) found that a physiotherapy intervention with a CBT component had the lowest overall costs (healthcare costs, lost days of work and disability pension costs were included) compared to physiotherapy or cognitive behavioural therapy (CBT) alone or usual care for workers with neck and back pain. The analysis was assessed as partially applicable to the review question, with potentially serious limitations.
- One cost-effectiveness, cost-benefit and cost-utility analysis (Lambeek, 2010) concluded that an integrated work based CBT programme for people on sick-leave with lower back pain was cost-effective compared to usual care, saving money and generating QALYs. The analysis was assessed as partially applicable to the review question, with potentially serious limitations.
- One cost-effectiveness and cost-benefit analysis (Loisel, 2002) found evidence that an
 occupational intervention, clinical intervention or combination of the two (the Sherbrook
 Model) for people absent with back pain may be cost-effective in pairwise comparisons
 with usual care. The study had fewer than 30 people in each study arm and no statistical
 significance testing of results was performed. This analysis was assessed as partially
 applicable to the review question, with minor study limitations.
- One cost-effectiveness analysis (Meijer, 2006) found no evidence that a residential multi-disciplinary team programme for people on sick-leave with upper musculoskeletal disorders was cost-effective compared to usual care. The analysis was assessed as partially applicable to the review question, with potentially serious limitations.
- One cost-effectiveness, cost-benefit and cost-utility analysis (Radford, 2012) found that a vocational rehabilitation programme for people with traumatic brain injury returned more people to work and was potentially cost-effective. The analysis was assessed as not being applicable to the review question because the population included students and those in unpaid employment with potentially serious limitations.

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- One cost-effectiveness and cost-benefit analysis (Rebergen, 2009) found that care
 following a national guideline delivered by occupational physicians for people on sick
 leave due to mental disorders resulted in no difference in work outcomes but lower
 healthcare costs than usual care and so was possibly cost-effective. The analysis was
 assessed as partially applicable to the review question, with potentially serious limitations.
- One cost-benefit analysis (Schene, 2007) found that an occupational therapist led CBT intervention for people absent from work with depression reduced work days lost and was likely to be cost-effective compared to treatment as usual. The analysis was assessed as partially applicable to the review question, with potentially serious limitations.
- One cost-utility analysis (Squires, 2011) using modelling based upon previously published studies found that for people on sickness absence with musculoskeletal disorders, a workplace intervention and a combination workplace intervention with physical activity and education intervention generate more QALYs compared to usual care at a lower cost and so therefore dominating usual care. The combination intervention had lower costs and generated more QALYs than a workplace intervention or physical activity intervention alone. The analysis was assessed as partially applicable to the review question, with potentially serious limitations.
- One cost-effectiveness and cost-utility analysis (Steenstra, 2006) found that for workers sick-listed due to low back pain, a work place intervention based upon ergonomics returned people to work faster than usual care at a slightly higher cost. A clinical intervention involving a graded activity programme with behavioural therapy was found to be less effective and more costly than usual care. The analysis was assessed as partially applicable to the review question, with potentially serious limitations.
- One cost-utility analysis (Uegaki, 2010) found that a general practitioner (GP) customised
 activation programme for people on sick leave due to stress related sick leave was not
 statistically superior in costs or QALYs compared to usual care managed by a GP
 although there was as high a likelihood of the intervention being cost-effective for people
 on sick leave due to stress-related mental disorders. The analysis was assessed as
 partially applicable to the review question, with potentially serious limitations.

30 The committee's discussion of the evidence

31 Cost effectiveness and resource use

- The committee noted the lack of health economic literature directly applicable to the UK. And even though it was mixed, they were mindful that overall it suggested interventions for people
- on sick leave due to musculoskeletal disorders including back pain or common mental
- disorders to support them to return to work could be cost effective. Therefore a new health
- 36 economic model was developed to determine how cost-effective an intervention will be in
- 37 helping employees on sickness absence to return to work.
- 38 Because the committee were concerned that interventions and size and type of organisation
- 39 vary greatly and a myriad of factors can impact sickness absence and return to work the
- 40 model adopted a generalised approach. Multiple sensitivity analyses were carried out which
- showed the results varied greatly by key model inputs such as the cost and effectiveness of
- the intervention, reduction in absenteeism and baseline rate of absenteeism.
- The committee noted that the results of the model reinforced the findings of the cost
- 44 effectiveness review that interventions for people on sick leave due to musculoskeletal
- disorders or common mental disorders could be cost effective. However, they were mindful
- 46 that these results are influenced by multiple factors some of which are specific to the local
- 47 conditions and that these may explain the mixed findings reported earlier.
- The committed also noted that the analysis showed in general a company with high turnover
- 49 costs or costs of absenteeism will likely benefit from an intervention to reduce sickness

DRAFT FOR CONSULTATION

Helping employees return to work and reducing long-term sickness absence (RQ 3a)

- absence, particularly if the intervention is effective and less expensive than the overall costs
- of absenteeism or replacing a worker. The committee were aware that the reverse is also
- 3 true. For example, an organisation with low baseline turnover costs or low levels of
- 4 absenteeism will find it more difficult to realise cost savings by implementing an intervention
- 5 aimed at reducing sickness absence, though this does not mean that other factors could not
- also benefit the organisation. The committee appreciated employers may be interested in
- 7 factors other than pure cost savings for example if the organisation is willing to pay for an
- 8 intervention that will benefit the workers and the organisation itself.
- 9 The committee noted that the results were influenced by multiple factors that are highly
- dependent on factors specific to each organisation as well as external factors such as the
- individual's personal life, labour market and culture of the workplace. They also noted that
- some identified benefits could not be quantified suggesting that the overall benefits might be
- greater than those reported by the model. So the committee concluded that such
- interventions could offer good value for money dependent on local circumstances.

References

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- 2 1. van Oostrom SH, Heymans MW, de Vet HCW, van Tulder MW, van Mechelen W, Anema
- 3 JR. Economic evaluation of a workplace intervention for sick-listed employees with distress.
- 4 Occup Environ Med. 2010;67(9):603-10.
- 5 2. Arends I, Bultmann U, van Rhenen W, H G, van der Klink JJL. Economic evaluation of a
- 6 problem solving intervention to prevent recurrent sickness absence in workers with common
- 7 mental disorders. PLoS ONE. 2013;8(8):e71937.
- 8 3. Brouwers E, de Bruijne M, Terluin B, Tiemens BG, Verhaak PFM. Cost-effectiveness of an
- 9 activating intervention by social workers for patients with minor mental disorders on sick
- leave: a randomized controlled trial. Eur J Public Health. 2007;17(2):214-20.
- 4. Bultmann U, Sherson D, Olsen J, Hansen CL, Lund T, Kilsgaard J. Coordinated and
- 12 tailored work rehabilitation: a randomized controlled trial with economic evaluation
- undertaken with workers on sick leave due to musculoskeletal disorders. J Occup Rehabil.
- 14 2009;19(1):81-93.
- 15 5. Finnes A, Enebrink P, Sampaio F, Sorjonen K, Dahl J, Ghaderi A, et al. Cost-Effectiveness
- of Acceptance and Commitment Therapy and a Workplace Intervention for Employees on
- 17 Sickness Absence due to Mental Disorders. J Occup Environ Med. 2017;59(12):1211-20.
- 18 6. Jensen IB, Bergstrom G, Ljungquist T, Bodin L. A 3-year follow-up of a multidisciplinary
- rehabilitation programme for back and neck pain. Pain. 2005;115(3):273-83.
- 7. Lambeek LC, Bosmans JE, Van Royen BJ, Van Tulder MW, Van Mechelen W, Anema JR.
- 21 Effect of integrated care for sick listed patients with chronic low back pain: economic
- evaluation alongside a randomised controlled trial. BMJ. 2010;341:c6414.
- 8. Loisel P, Lemaire J, Poitras S, Durand MJ, Champagne F, Stock S, et al. Cost-benefit and
- cost-effectiveness analysis of a disability prevention model for back pain management: a six
- 25 year follow up study. Occup Environ Med. 2002;59(12):807-15.
- 9. Meijer EM, Sluiter JK, Heyma A, Sadiraj K, Frings-Dresen MH. Cost-effectiveness of
- 27 multidisciplinary treatment in sick-listed patients with upper extremity musculoskeletal
- disorders: a randomized, controlled trial with one-year follow-up. Int Arch Occup Environ
- 29 Health. 2006;79(8):654-64.
- 30 10. Radford K, Phillips J, Drummond A, Sach T, Walker M, Tyerman A, et al. Return to work
- 31 after traumatic brain injury: cohort comparison and economic evaluation. Brain Inj.
- 32 2013;27(5):507-20.
- 11. Rebergen DS, DJ B, van Tulder MW, van der Beek AJ, van Mechelen W. Cost-
- 34 effectiveness of guideline-based care for workers with mental health problems. J Occup
- 35 Environ Med. 2009;51(3):313-22.
- 36 12. Schene AH, Koeter MW, Kikkert MJ, Swinkels JA, McCrone P. Adjuvant occupational
- 37 therapy for work-related major depression works: randomized trial including economic
- 38 evaluation. Psychol Med. 2007;37(3):351-62.
- 39 13. Squires H, Rick J, Carroll C, Hillage J. Cost-effectiveness of interventions to return
- 40 employees to work following long-term sickness absence due to musculoskeletal disorders. J
- 41 Public Health. 2012;34(1):115-24.
- 42 14. Steenstra IA, Anema JR, van Tulder MW, Bongers PM, de Vet HC, van Mechelen W.
- Economic evaluation of a multi-stage return to work program for workers on sick-leave due to
- 44 low back pain. J Occup Rehabil. 2006;16(4):557-78.

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Helping employees return to work and reducing long-term sickness absence (RQ 3a)

- 1 15. Uegaki K, Bakker I, de Bruijne M, van der Beek A, Terluin B, van Marwijk H, et al. Cost-
- 2 effectiveness of a minimal intervention for stress-related sick leave in general practice:
- 3 results of an economic evaluation alongside a pragmatic randomised control trial. J Affect
- 4 Disord. 2010;120(1-3):177-87.

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Appendices

2 Appendix A – Review protocols

- 3 Review protocols for review questions 1, 2 and 3
- 4 Review question 1a What interventions, programmes, policies or strategies are cost-effective in preventing or reducing recurrence of short-
- 5 term sickness absence among employees?
- Review question 2a What interventions, programmes, policies or strategies are cost effective in reducing the number of employees who move
- 7 from short- to long-term sickness absence?
- 8 Review question 3a What interventions, programmes, policies or strategies are cost effective in:
- helping employees on long-term sickness absence to return to work?
- reducing the recurrence of long-term sickness absence following a return to work?
- 11 The same protocols were used for both the effectiveness and cost-effectiveness reviews. See effectiveness reviews for full details.

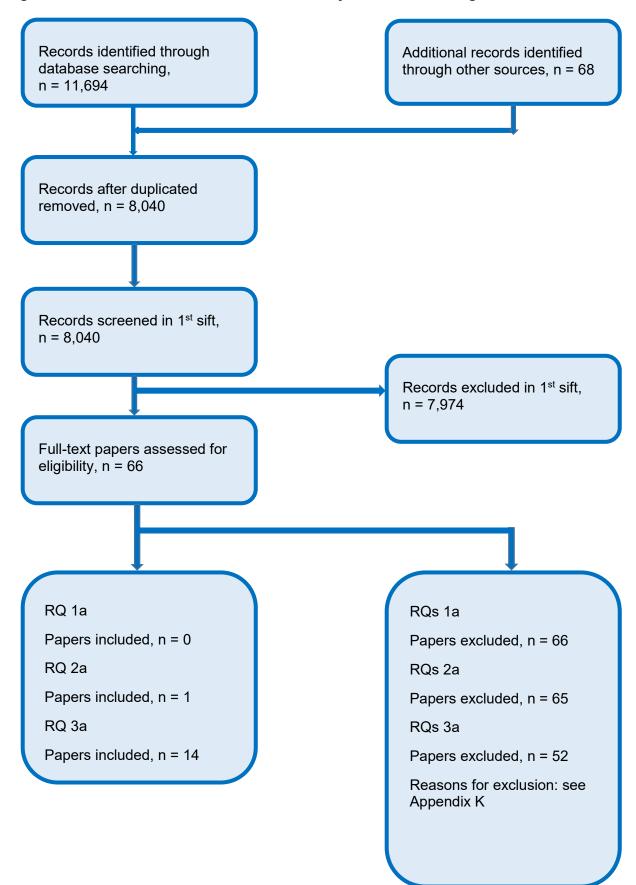
Appendix B – Literature search strategies

Guideline-wide search strategies were undertaken based on the review protocols provided for all review questions. See effectiveness reviews for full details.

Appendix G – Economic evidence study selection

The following flowchart shows the record selection process for all three review questions.

Figure 1: Flow chart of economic evidence study selection for the guideline



Appendix H – Economic evidence tables

Table 5: Summary of studies included in the economic evidence review for workplace health interventions for RQ 1a, 2a and 3a

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
Arends 2013 (Netherlands) Population: Workers (aged 18 to 63 years) who were diagnosed at start of sickness absence with a common mental disorder (CMD) and were now partially or fully ready to return to work. Intervention: SHARP-at work. A five steps intervention: return to work	Potentially serious limitations ^c	Partially applicable ^d	None	Mean cost per person SHARP Cost- effectivene ss analysis (CEA): €4,167 Cost- benefit analysis (CBA): Between €29,337 (human capital approach (HCA) to productivity loss) and €37,215 (friction cost approach (FCA) to	Recurrent sickness absence over 12 months SHARP: 39% CAU: 62%	SHARP vs CAU ^e CEA: €1,764 more CBA: €4,730 (HCA) CBA: €5,530 (FCA)	Recurrent sickness absence over 12 months, SHARP vs CAU, bootstrapped estimate: 24% lower	CEA, incremental cost- effectiveness ratio (ICER), SHARP vs CAU, per 1% of recurrent sickness absence prevented: €10,605 CBA Employer occupational health costs only, SHARP was €800 greater than with CAU Productivity loss (HCA)	Sensitivity analyses CEA Excluding an outlier, which was attributed to high costs due to hospitalisation in a psychiatric ward, an ICER of €-533 was calculated for the incidence of recurrent sickness absence, indicating SHARP could be cost-effective. Reduced SHARP costs did not change the direction of the primary analyses. CBA Reduced SHARP costs did not

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
(RtW) was started with the occupational physician (OP) monitoring and supporting the person through the steps. ^a Comparator: Care as usual (CAU) ^b				productivity loss) CAU CEA: €2,403 CBA: Between €24,607 (HCA) and €31,685 (FCA)				SHARP vs CAU: €6,046 Productivity loss (FCA) SHARP vs CAU: €3,995	change these results
Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Brouwers 2007 (Netherlands) Population: People (aged 18-60 years) absent from work on sick leave for no more than 3 months due to a minor mental disorder Intervention group (IG): 5 individual sessions (50	Potentially serious limitations ^h	Partially applicable ⁱ	None	Mean cost per person IG: €14,493 (exclusive of intervention costs of €13,305 total for all people) CAU: €14,482 CAU	Mean quality- adjusted life years (QALYs) per person not reported Sick leave duration until full return to work IG: 152.7 days	CBA IG vs CAU (excluding costs of intervention): €11 more expensive (not statistically different) Cost-utility analysis (CUA), based on 2,000	Mean QALYs per person IG vs CAU (Dutch EQ-5D): 0.056 higher IG vs CAU (UK EQ-5D): 0.044 higher Sick leave duration until full return to work IG vs CAU: 3.8 days sooner	CUA, ICER Probabilistic ICERs reported (ICERs are negative). IG vs CAU (Dutch EQ-5D): - €4,179 IG vs CAU (UK EQ-5D): - €5,306 IG was less expensive and more effective than CAU	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA. This demonstrated an even split between the north east and south east quadrants of the cost-effectiveness plane indicating that any difference in costs between IG and CAU were likely small. 52% of bootstrap estimates were in the south

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
minutes each), covering 3 stages, with a social worker: cause, coping and implementation . f				CEA: €2,403 CBA: €24,607 (HCA) to €31,685 (FCA)	CAU: 156.5 days No statistically significantly difference	bootstrap pairs IG vs CAU: €234 cheaper		CBA IG vs CAU: €11 more expensive (not statistically significant but did not include intervention	east quadrant where IG dominates CAU.
Comparator: Care as usual								costs)	
(CAU): routine general practitioner (GP) care ^g									

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Bultmann 2009 (Denmark) Population: Workers on sick leave for 4 to 12 weeks due to lower back pain (LBP) or musculoskelet al disorders (MSD)	Potentially serious limitations ^k	Partially applicable ¹	None	Mean cost per person, over 12 months CTWR: \$31,144 (\$3,321 without productivity loss) CCM: \$41,812	Sickness absence, per person, over 12 months CTWR: 656.6 hours CCM: 997.3 hours	Over 12 months, per person CTWR vs CCM: \$10,668 (\$1,548 more over 12 months without productivity loss)	Mean absence days averted CTWR vs CCM: 46.0	CEA CTWR vs CCM, per absence day avoided (without productivity loss): \$33.7	One way deterministic sensitivity analysis of a doubling of intervention costs and 25% reduction in wages still resulted in cost savings under the CBA.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Intervention: Coordinated and tailored work rehabilitation (CTWR), screening followed by a tailored rehabilitation plan developed by an interdisciplinar y team j	Limitations	Аррисавину	Comments	(\$1,773 without productivity loss)	EHEGIS	1 GOST	enecis	enectiveness	Oncertainty
Comparator: Conventional case management (CCM) provided by the municipality. No further information given.									
Finnes 2017 (Sweden) Population: Workers (at least 50% whole time	Potentially serious limitations ^q	Partially applicable ^r	None	12 month costs Healthcare perspective	QALY gains over 12 months ACT: 0.164 WDI: 0.122	12 months, vs ACT Healthcare perspective WDI: \$958	QALY gains over 12 months vs ACT WDI: -0.042	ICER Healthcare perspective	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA. This showed that

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
equivalent (WTE)) with sickness absence due to anxiety, depression, stress or exhaustion Interventions: • Acceptance and Commitment Therapy (ACT): a psychologica I intervention consisting of 6 X 60- minute sessions m • Workplace dialogue intervention (WDI): three meetings involving the participant plus work supervisor n • ACT + WDI: conducted by two	Limitations	Applicability	comments	ACT: \$5,507 WDI: \$6,465 ACT+WDI: \$6,141 TAU: \$6,207 Societal perspective ACT: \$14,452 WDI: \$15,649 ACT+WDI: \$17,066 TAU: \$15,593	ACT+WDI: 0.168 TAU: 0.155	ACT+WDI: \$634 TAU: \$700 Societal perspective WDI: \$1,197 ACT+WDI: \$2,614 TAU: \$1,141	effects ACT+WDI: 0.046 TAU: -0.009	effectiveness Both TAU and WDI were dominated by ACT ACT vs baseline: \$33,579 per QALY gained ACT+WDI vs ACT: \$158,500 per QALY gained Societal perspective Both TAU and WDI were dominated by ACT alone. Compared to ACT, ACT+WDI had an ICER of \$30,804 per QALY gained	for ACT compared to ACT+WDI from both the healthcare and societal perspectives, the percentage of bootstrap iterations were spread roughly equally across all four quadrants, although approximately 60% of iterations in both perspectives had ACT+WDI more costly than ACT and 50% of iterations of ACT+WDI were more effective. Scenario analysis showed that using Swedish utility weights (rather than English in the base case) resulted in ACT being the dominant strategy. A second scenario explored the impact of using costs as if the intervention

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
different therapists of the comparator: Treatment as usual (TAU) was treatment as planned in a primary care centre or other care facility.	Limitations	Аррисавину	Comments	COSIS	LITECTS	T COST	enects	enectiveness	'regular' setting in which case ACT would no longer dominate TAU but have an ICER of \$71 per QALY gained (healthcare perspective) and ACT+WDI had an ICER of \$286,000 per QALY gained compared to ACT.
Jensen 2005 (Sweden) Population: Blue-collar and service/care workers (aged 18 to 60 years) with nonspecific back pain resulting in sick leave for 1-6 months Interventions: Behaviour-oriented physiotherapy (PT) s Cognitive behavioural	Potentially serious limitations ^u	Partially applicable v	None	Mean cost per person, at 3 years: female; male BM: €107,703; €130,015 PT: €189,760; €220,268 CBT: €157,800; €199,824	Working days lost per year: female; male Pre intervention BM: 67; 72 PT: 76; 92 CBT:115; 109 TAU: 65; 80 Post intervention BM: 99; 123	Incremental cost per person, at 3 years, vs TAU: female; male BM: - €137,509; -€63,224 PT: -€55,452; €27,029 CBT: -€87,412; €6,585	Change in working days lost per year pre and post intervention: female; male BM: 22; 51 PT: 19; 18 CBT:-6; -8 TAU: 17; 6	For women, BM, PT and CBT were all less expensive over 3 years vs TAU with BM having the lowest cost per person vs CBT BM vs CBT: €50,097 less expensive BM vs PT: \$82,057 less expensive BM vs TAU: €137,509 less expensive	Not undertaken

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
therapy (CBT): to improve pain management t Behavioural medicine programme (BM): PT+CBT Comparator: Treatment as usual (TAU). No additional interventions outside the normal routines in health care. This was not otherwise described.				TAU: €245,212; €193,239	PT: 95; 110 CBT:109; 101 TAU: 82; 86			For men, CBT and PT were both more expensive over three years than TAU with BM being less expensive than TAU by €63,224	
Lambeek 2010 (Netherlands) Population: Employed or self-employed workers (aged 18 to 65 years) on full or partial sick leave for 12 weeks to 2	Potentially serious limitations ^x	Partially applicable ^y	None	At 12 months IC: £13,165 (£1,479 direct costs, £11,686 indirect costs)	Days until sustainable return to work IC: 129 UC: 197 QALYs	At 12 months IC vs UC: -£5,310 (£217 direct costs, -£5,527 indirect costs)	Days until sustainable return to work, per person IC vs UC: -68	£3 extra cost for every day earlier return to work with IC. Direct costs only considered.	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA and CEA. This showed that in 98% of iterations IC dominated UC, for the CUA and for the

			Other			Incrementa	Incremental	Cost-	
Study years due to non-specific low back pain (LBP). Intervention: Integrated care (IC), a graded activity	Limitations	Applicability	Other comments	Costs UC: £18,475 (£1,262 direct costs, £17,213 indirect costs)	Effects IC: 0.74 UC: 0.65	Incrementa I cost	Incremental effects	Cost- effectiveness IC vs UC: dominant IC cost saving over UC, per person: £5,310 QALY gain, per person:	Uncertainty CEA that if there was a willingness to pay of £10 for one day earlier return to work there was a 95% chance that IC was cost-effective. Scenario analysis
protocol at the workplace. w Comparators: usual care (UC): referred to occupational physician and GP with a letter containing the advice to treat them according to the Dutch guidelines for patients with LBP.								0.09 (direct and indirect costs considered) CBA IC, return on investment for every £1 spent: £26 (direct and indirect costs considered)	showed that if only complete cases were considered then there was no statistical difference in costs between IC and UC. A further scenario explored the impact of the intervention only for people aged under 55 years which resulted in a doubling in the costs of IC.
									A final scenario analysis showed that varying productivity levels did not impact on results.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Loisel 2002 (Canada) Population: Workers absent for more than 4 weeks with occupational back pain Interventions: Occupationa I intervention (OI): visits to the study occupational medicine physician and a participatory ergonomics intervention with the study ergonomist, the injured worker, their supervisor, and managemen t and union	Minor limitations aa	Partially applicable bb	The study had fewer than 30 people in each study arm and no statistical significance testing of results was performed	At 12 months OI: \$9,569 CRI: \$12,038 Sherbrooke model: \$12,137 UC: \$9,789 At 6.4 years OI: \$16,252 CI: \$16,902 Sherbrooke model: \$14,494 UC: \$33,079	Mean number of days on full benefits (DFB) At 12 months OI: 116.1 CI: 114.9 Sherbrooke model: 115.9 UC: 126.9 At 6.4 years OI: 228.0 CI: 178.7 Sherbrooke model: 125.6 UC: 418.3	Incremental costs vs UC At 12 months OI: -\$220 CI: \$2,250 Sherbrooke model: -\$2,348 At 6.4 years OI: -\$16,827 CI: -\$16,176 Sherbrooke model: -\$18,585	Mean number of DFB, incremental difference vs UC At 12 months OI: -10.8 CI: -12 Sherbrooke model: -11 At 6.4 years OI: -190.3 CI: -239.6 Sherbrooke model: -293.7 UC: 418.3	CEA (cost per DFB saved vs UC) At 12 months OI: -\$20.40 (dominated UC) CI: \$187.40 Sherbrooke model: \$213.50 At 6.4 years OI: -\$88.40 (dominated UC) CI: -\$67.50 (dominated UC) Sherbrooke model: -\$63.50 (dominated UC) CBA Cost differential vs UC	Sensitivity analyses were performed by varying the total healthcare costs by 60% to 190% and income per capita by 85% to 125%. Over a 6.4 year time period all interventions remained dominant vs UC over the cost ranges considered.

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
representativ es. • Clinical								At 12 months	
rehabilitation								OI: \$220	
intervention								CI: -\$2,250	
(CRI), with a								Sherbrooke	
back pain								model: -\$2,348	
specialist and								At 6.4 years	
potentially a								OI: \$16,827	
multidisciplin								CI: \$16,176	
ary work rehabilitation								Sherbrooke	
intervention								model:	
at 12 weeks								\$18,585	
of absence z									
OI+CRI ("Sherbrook									
e Model")									
,									
Comparator:									
Usual care									
(UC): with worker's									
physician									
receiving no									
advice about return to work.									
	Detentially	Dortiolly	None	Total costs	Doroontogo	Ingramantal	Difference in	MDT was	Not undertaken
Meijer 2006 (Netherlands)	Potentially serious	Partially applicable ee	None	Total costs per week	Percentage of regular	Incremental costs of	percentage	more	NOT UTILIETTAKETI
(limitations dd	, ,			hours	MDT vs UC	of regular	expensive	
Population:				At 2	worked	per week	hours worked	compared to	
				months				UC and did not	

			Other			Incrementa	Incremental	Cost-	
Bank and university workers on at least 50% contracts with 50% sick leave in the last 4 to 20 weeks due to non-specific upper extremity musculoskelet al disorders (MSDs) Intervention: Multidisciplinar y treatment (MDT): an outpatient training programme carried out at Dutch rehabilitation centres. © Comparator: Usual care (UC): supervision by occupational health	Limitations	Applicability	comments	MDT: €1,335 UC: €448 At 6 months MDT: €664 UC: €359 At 12 months MDT: €430 UC: €315	Effects Baseline MDT: 29% UC: 29% 6 months MDT: 82% UC: 72% 12 months MDT: 86% UC: 73% There was no statistical difference between MDT and UC at any time point	At 2 months: €887 At 6 months: €305 At 12 months: €115	effects Baseline: 0% 6 months: 10% 12 months: 13% There was no statistical difference between MDT and UC at any time point	effectiveness increase the proportion of days worked. MDT was not cost-effective.	Uncertainty

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
services. UC could include treatment at the workplace and in the regular health care system, initiated by a GP, or medical specialist.									
Radford 2012 (UK) Population: Patients in paid or voluntary work or in full time education, hospitalised for at least 48 hours due to traumatic brain injury (TBI) Intervention: TBI Vocational Rehabilitation (TBI-VR), provided by an occupational therapist ff	Potentially serious limitations hh	Partially applicable ⁱⁱ	None	At 12 months Health and social care perspective TBI-VR: £2,106.94 UC: £2,031.71 Societal perspective TBI-VR: £8,786 UC: £10,648	QALYs (at 12 months) TBI-VR: 0.1938 UC: 0.1763 Return to work or education (at 12 months) TBI-VR: 75% UC: 60%	At 12 months Health and social care perspective TBI-VR vs UC: £76.24 more expensive Societal perspective TBI-VR vs UC: £1,867 less expensive	QALYs (at 12 months) TBI-VR vs UC: 0.0175 more QALYs Return to work or education (at 12 months) TBI-VR vs UC: 15% more people returned to work	CEA, per person returned to work, TBI-VR vs UC Health and social care perspective: £501.33 Societal perspective: TBI-VR was more effective and saved money CBA Health and social care	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA. The boot strapped ICER was £2,567 lower than the deterministic ICER. Sensitivity analysis using imputed data for missing values more than doubled the cost per person returned to work in the CEA and increased the ICER per QALY gained in the CUA to £35,873 with TBI-VR.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Study Comparators: Usual care (UC): participants in hospitals without TBI- VR. 99	Limitations	Applicability	comments	Costs	Effects	I cost	effects	perspective: TBI-VR was £75.23 more costly vs UC Societal perspective: TBI-VR was £1,863 less expensive than UC. Neither difference was statistically significant CUA, TBI-VR vs UC From a health and social care perspective: £4,299 per QALY gained Neither the QALY gain nor cost difference was statistically significant	Uncertainty

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
Rebergen 2009 (Netherlands) Population: Police workers on sick leave due to common mental disorders (CMDs) Intervention: Guideline based care (GBC), treatment by occupational physicians (OPs) according to the Dutch guideline for workers with mental health problems. Comparator: Usual care (UC): minimal involvement of	Potentially serious limitations kk	Partially applicable	None	At 12 months Health care perspective GBC: €2,145 UC: €2,664 Societal perspective (HCA for productivity loss) TBI-VR: €14,114 UC: €14,202 No costs were statistically significantly different between groups	Days of sick leave GBC: 113 UC: 114 These were not statistically significantly different	At 12 months Health care perspective GBC vs UC: €520 less expensive	Days of sick leave, at 12 months GBC vs UC: 1 fewer The difference was not statistically significant	CEA GBC vs UC, ICER per sick day avoided: - €736 CBA Estimated net monitory benefit of GBC, per person: €3,582 Outcomes are similar between GBC and UC, but direct costs were lower with GBC. The authors concluded that GBC could be cost-effective.	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CEA. The iterations showed there was never more than a 50% chance of GBC being cost- effective per day of sick leave avoided regardless of the value of the day of work lost. Different approaches to measuring productivity loss were analysed but did not affect the main findings.

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
the OP and easy access to counselling by a psychologist.									
Schene 2007 (Netherlands) Population: Workers (aged 18 years or over) with at least 50% absence over 10 weeks to 2 years due to work related major depressive disorder (WRMDD) Intervention: Treatment as usual (TAU) + occupational therapy (OT): TAU was as described below (comparator). OT was the addition of OT, which had	Potentially serious limitations nn	Partially applicable °°	None	At 12 months (cost of intervention only) TAU+OT: \$3,149 TAU: \$1,891	No health or employmen t outcomes were reported beyond earnings over 12 month period	TAU+OT vs TAU at 12 months (cost of intervention only): \$1,258 more expensive (Not statistically significant)	Not applicable	Difference in total earnings minus costs of intervention at 12 months TAU+OT vs TAU: \$3,952 higher (Not statistically significant)	The base case was a bootstrapped analysis to account for stochastic uncertainty. The only sensitivity analysis performed was on the value of an hour's work. As the value reduces the probability that TAU+OT is more cost-effective than TAU falls. In the base case it is 75.5% at \$36.88 per hour and falls to 52.5% at \$10 per hour

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
diagnostic and									
therapeutic									
phases. mm									
Comparator:									
Treatment as									
usual (TAU):									
out-patient									
psychiatric									
treatment for									
depression									
according to American									
Psychiatric									
Association									
(APA)									
guideline and									
antidepressant									
s and/or									
cognitive									
behavioural									
therapy (CBT)									
with senior									
psychiatric residents.									
Visits lasted 30									
minutes every									
2–3 weeks.									
2 0 WOOKS.									
Squires 2012	Potentially	Partially	Costs and	Not	Increased	Not	Not reported	CUA (from	Sensitivity and
(UK)	serious	applicable ^{tt}	effects data	reported	likelihood of	reported		societal	scenario analyses
	limitations ss		were very		return to			perspective so	were undertaken.
Population:					work (i.e.			includes costs	PAEW was not

Workers on sick leave for 1 reported. week to 6 months due to musculoskelet al disorders (MSDs) Interventions poorly relative risk) within from lost employer risk) within from lost employer misk) within from lost employer meand the months of sickness absence (obtained from a literature effective. Interventions to NHS and dominant if or misk) within from lost employer misk) within from lost employer wages) Perspective to taken and the probability of leave recurring doubled. WI and PAEW are both cheaper than UC and more effective. In a threshold the provious of t				Other			Incrementa	Incremental	Cost-	
sick leave for 1 week to 6 months due to musculoskelet al disorders (MSDs) Results were presented on a cost- effectiveness plane and not Interventions reported. risk) within the first 6 months of months of sickness absence (obtained from a literature literature roviow) reported. risk) within months of sickness absence (obtained from a literature roviow) risk) within months wages) perspective or taken and the probability of leave recurring doubled. In a threshold	Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
evidence identified WI: 1.12 PAE is more costly but more effective than UC. • Workplace interventions (WI): a workplace assessment and work • Workplace interventions.	Workers on sick leave for 1 week to 6 months due to musculoskelet al disorders (MSDs) Interventions (synthesized evidence identified through a review): • Workplace interventions (WI): a workplace assessment and work modification s based on participative ergonomics involving all relevant stakeholders pp • Physical activity and education	Limitations	Applicability	comments poorly reported. Results were presented on a cost- effectiveness plane and not	Costs	relative risk) within the first 6 months of sickness absence (obtained from a literature review) WI: 1.12 PAE: 1.06 PAEW:			effectiveness to NHS and from lost wages) WI and PAEW are both cheaper than UC and more effective. PAE is more costly but more effective than UC. PAEW dominates all interventions. CEA, cost per sick day avoided PAEW is the dominant	dominant if only the employer perspective was taken and the probability of sick leave recurring was doubled. In a threshold analysis, if the intervention costs were less than an additional £3,000 and returns at least an additional 3% of people to work (32/1,000) in comparison to UC, then it is likely to result in a cost per QALY gained below

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
form of									
physical									
activity and									
education									
around how									
to deal with									
pain and									
body									
mechanics									
 Physical 									
activity,									
education									
and									
workplace									
visit									
(PAEW):									
WI+PAE									
plus a									
workplace									
visit by the									
employee									
and the									
physical									
therapist to									
inform									
rehabilitation									
. qq									
Comparator:									
Usual care									
(UC) treatment									
of MSDs in the									
UK ^{rr}									

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
Steenstra 2006 (Netherlands) Population: Workers on sick leave for 2 to 6 weeks due to low back pain (LBP) The study had a 2-stage design. Between 2 to 8 weeks of sick-leave, patients received either the workplace intervention (WI) or usual care (UC). After 8 weeks, approximately half of each group also received a clinical intervention (CI).	Potentially serious limitations **	Partially applicable yy	None	Mean total costs WI: €8,993 UC: €9,109 UC+CI: €10,537 UC+UC: €10,885 WI+CI: €12,391 WI+WI: €11,096	Actual QALY values not reported Sick leave (calendar days) WI: 108.5 UC: 135.2 UC+CI: 172.9 UC+UC: 155.9 WI+CI: 181.7 WI+WI: 115.3	Vs UC over 12 months WI: €16 UC+CI: €1,428 WI+CI: €3,282 None of the differences were statistically significant	Incremental QALY values were not reported Sick leave vs UC, calendar days WI: -26.7 days UC+UC: 20.7 days WI+CI: 46.5 days	CEA (per 1 day less of sick leave) WI vs UC: €19 WI+CI vs UC: €11 UC+CI vs UC: €29 CUA (cost per QALY) WI vs UC: -€1,483 WI+CI vs UC: €24,416 UC+CI vs UC: €5,447	The base case was a bootstrapped analysis to account for stochastic uncertainty. The cost-effective planes (and confidence intervals of point estimates) suggested that WI and UC were likely similar in cost but that WI was more effective. CI is likely less effective and more costly than both WI and UC. Scenario analyses suggested using a fixed sum per day of production lost, net rather than calendar sick days and using a HCA approach to productivity loss did not significantly influence results.

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
 WI: usual care, a workplace assessment and modification and communicati on between OP and GP in order to discuss how to counsel the worker to RtW ^{uu} Clinical intervention (CI): a graded activity programme of 26 x 1-hour sessions, with a frequency of 2 sessions per week ^w 									
Comparator: Usual Care (UC): Dutch OP guidelines									

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty
for LBP delivered by a GP. ^{ww}									
Uegaki 2010 (Netherlands) Population: Workers with partial sick leave over 3 months due to distress Intervention: Minimal intervention for stress-related mental disorders with sick leave (MISS): a GP customized version of an activating approach zz Comparator: usual care (UC) managed by a GP. No further	Potentially serious limitations aaa	Partially applicable bbb	None	At 12 months MISS: €12,538 UC: €12,722	QALYs over 12 months MISS: 0.78 UC: 0.76	MISS vs UC, incremental cost at 12 months: A saving of €184 This was not statistically significant	QALY gain, over 12 months MISS vs UC: 0.02 QALYs This was not statistically significant	ICER MISS vs UC: -€7,356 per QALY gained Neither change in costs nor change in QALYs were statistically significantly different between MISS and UC	The base case was a bootstrapped analysis to account for stochastic uncertainty. Cost-effectiveness planes showed that in the base case 77% of bootstrapped pairs would be considered cost-effective at a willingness to pay (WTP) threshold of €25,600 per QALY. Sensitivity analysis explored different approaches to costing lost productivity but did not significantly influence the results. Subgroup analysis suggested MISS may be most cost-

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
information given.									effective for patients with stress related mental disorders, which was the only analysis which had statistically significant improvement in QALYs vs UC: -€28,278
van Oostrom 2009 (Netherlands) Population: Workers on sick leave for 2 to 8 weeks due to distress Interventions: Workplace intervention (WI): a stepwise communication process to identify and solve obstacles to return to work (RTW).ccc	Potentially serious limitations eee	Partially applicable fff	None	Mean total costs, over 12 months Societal perspective WI: €3,201 UC: €2,758 Employer perspective WI: €1,386 UC: €802	Mean duration of sick leave, over 12 months Cost Effectivene ss Analysis (CEA) WI: 133 days UC: 134 days Cost utility analysis (CUA) WI: 0.77 UC: 0.78	Mean cost difference, over 12 months Societal perspective WI vs UC: €443 more costly (not statistically significant) Employer perspective WI vs UC: €583 more costly (not statistically significant)	Mean duration of sick leave, over 12 months WI vs UC: 1 day fewer (not statistically significant) QALYs WI vs UC: 0.01 less (not statistically significant)	CEA WI vs UC ICER: €627 per sick day avoided Neither change in costs or change in sick days were statistically different between WI and UC CBA Net monetary benefit with WI was -€1,987 with human capital	The base case was a bootstrapped analysis to account for stochastic uncertainty. Cost effectiveness planes showed substantial uncertainty in results which reflects the statistical uncertainty in the point estimates of cost differences and effectiveness measures between WI and UC. Subgroup analysis suggested WI may be most cost-

Comparators: Usual care (UC): treatment by approach (HCA) and - patients with an intention to return friction cost to work but findir	Study	Limitations	Applicability	Other	Coete	Effects	Incrementa	Incremental	Cost-	Uncertainty
CUA WI vs UC, incremental cost-utility ratio (ICUR) (HCA): -€184,562 per	Usual care (UC): treatment by an occupational physician, according to the Dutch	Limitations	Applicability	comments	Costs	Effects	I cost	effects	approach (HCA) and - €1,700 with friction cost approach (FCA) WI was statistically significantly more costly than UC and changes in costs of productivity loss favoured UC but were not statistically significant regardless of productivity measure. CUA WI vs UC, incremental cost-utility ratio (ICUR) (HCA):	patients with an intention to return to work but findings were still limited in statistical

Study	Limitations	Applicability	Other comments	Costs	Effects	Incrementa I cost	Incremental effects	Cost- effectiveness	Uncertainty
								WI vs UC, ICER (FCA): - €155,850 WI dominates UC. Neither cost differences with WI or QALY gains were statistically significant	

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty

ACT: acceptance and commitment therapy; APA: American Psychiatric Association; BM: behavioural medicine; CAU: care as usual; CBA: cost-benefit analysis; CBT: cognitive behavioural therapy; CCM: conventional case management; CEA: cost-effectiveness analysis; CG: control group; CI: clinical intervention; CMD: common mental disorder; CRI: clinical rehabilitation intervention; CTWR: coordinated and tailored work rehabilitation; CUA: cost-utility analysis; DFB: days on full benefits; FCA: frictional cost approach; GBC: guideline based care; GP: general practitioner; HCA: human capital approach; IC: Integrated care; ICER: incremental cost-effectiveness ratio; ICUR: incremental cost-utility ratio; IG: intervention group; LBP: lower back pain; MDT: multidisciplinary treatment; MSD: musculoskeletal disorders; MISS: minimal Intervention for stress-related mental disorders with sick leave; MSK: musculoskeletal disorders; NA: not applicable; OI: occupational intervention; OP: occupational physician; OT: occupational therapy; PAE: physical activity and education intervention; PAEW: physical activity, education and workplace visit; PSA: probabilistic sensitivity analysis; PT: physiotherapy; QALY: quality-adjusted life year; RtW: return to work; TAU: treatment as usual; TBI: traumatic brain injury; TBI-VR: traumatic brain injury vocational rehabilitation; UC: usual care; VAS: visual analogue scale; VR: vocational rehabilitation; WDI: workplace intervention; WRMDD: work related major depressive disorder; WTE: whole time equivalent; WTP: willingness to pay

- (a) The five steps comprised: (1) making an inventory of problems and/or opportunities encountered at work after RtW; (2) brainstorming about solutions/realisations; (3) writing down solutions/ realisations and the support needed and assessing the applicability of these solutions; (4) discussing solutions/ realisations and making an action plan with the supervisor; (5) evaluating the action plan/implementation of solutions.
- (b) Occupational physicians (OPs) enacted the guideline of the Netherlands Society of Occupational Medicine "The treatment of workers with mental health problems by the OP". It is primarily aimed at structuring OPs' treatment to help sick-listed workers with mental health issues to RtW. Limited focus is given to follow-up after RtW has been achieved: only one consultation, to address relapse.
- (c) A 12 month time horizon was used, which is insufficient to capture whether outcomes are sustained. Impact on QALYs was not considered.
- (d) It was unclear how long people in the study had been absent from work. The study was in the Netherlands where the occupational support offered is differently organised than in the UK.
- (e) Calculated from the total costs reported in the paper.
- (f) Treatment was over 10 weeks and entailed 3 stages: (i) understanding the cause of loss of control (ii) the development of problem-solving strategies; and (iii) their implementation.
- (g) This could include medication, counselling or referral.
- (h) An 18 month time horizon was used, which is insufficient to capture whether outcomes were sustained. Deterministic sensitivity analysis was not performed.
- (i) People in the study could have been absent for less than four weeks. The study was in the Netherlands where the occupational support offered is differently organised than the UK. It is unclear how utility values were derived.
- (j) After 4 to 12 weeks of sick leave: (1) work disability screening was conducted: a multidisciplinary assessment of disability and functioning and identification of barriers for RtW and (2) a coordinated, tailored and action-oriented work rehabilitation plan was developed by an interdisciplinary team with continuous feedback on the plan from the sick listed worker, the interdisciplinary team, the workplace, and major stakeholders. The interdisciplinary team consisted of an occupational physician, an occupational physician, an occupational physician, a chiropractor, a psychologist, and a social worker who had the role of case worker.
- (k) A 12 month time horizon was used which is insufficient to capture whether outcomes are sustained. Impact on QALYs was not considered. Probabilistic sensitivity analysis (PSA) was not undertaken.

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty

- (I) The study was in Denmark where the organisation of the sickness benefits system is similar to the UK, but may still be different enough to limit the generalisability of findings.
- (m) The first part (sessions 1 to 3) emphasized mindfulness, cognitive defusion, and acceptance. During sessions 4 to 6, the focus was on exploring and clarifying personal values and committing to pursuing valued life activities.
- (n) The first step was an individual interview with the participant at the clinic followed by an interview with the participant's supervisor at the workplace. These meetings, lasting up to 60 minutes, aimed to investigate the participants' and the supervisors' views upon causes of the sickness absence, and what might facilitate RTW.
- (o) There was no integration or coordination of the two interventions, and no interaction between therapists.
- (p) This typically included psychotherapy, cognitive behavioural therapy and/or pharmacological treatments, physical therapy and counselling.
- (q) A 12 month time horizon was used which is insufficient to capture sustainability of outcomes. Medication costs were not considered.
- (r) It was unclear how much time people in the study had been absent from work and the study was in Sweden where the occupational support offered is differently organised than in the UK. Drug costs were not considered despite a societal perspective.
- (s) This consisted of approximately 20 scheduled hours per week aimed at enhancing the physical functioning and facilitating lasting behaviour change. Each participant was assigned to an individually tailored training programme.
- (t) This was an average of 13 to 14 scheduled hours per week aimed at improving participants' ability to manage their pain and resume a normal level of activity.
- (u) A time horizon of 3 years was used which is only partly sufficient to assess whether outcomes are sustainable over the long-term. No deterministic sensitivity analysis or probabilistic sensitivity analysis was performed. Cost sources were poorly reported.
- (v) The study was conducted in Sweden where the organisation of sickness benefits system is similar to the UK, but may still be different enough to limit the generalisability of the findings.
- (w) The workplace intervention protocol formulated a consensus-based plan for adaptations at work to facilitate return to work. The integrated care team consisted of a medical specialist, occupational therapist, physiotherapist, and clinical occupational physician.
- (x) Only a 12 month time horizon was used which is insufficient to capture whether outcomes are sustained.
- (y) The study includes people on partial sick leave so may not be considered 'continuous absence'. The study was set in the Netherlands where the organisation of healthcare and sickness may be different enough from the UK to limit the generalisability of findings.
- (z) Clinical examination by a back pain medical specialist, participation in a back school after eight weeks of absence from regular work and, if necessary, a multidisciplinary work rehabilitation intervention after 12 weeks of absence from work.
- (aa) Impact on QALYs was not considered and no probabilistic sensitivity analysis was reported.
- (bb) The study was conducted in Canada where the organisation of the sickness benefits system is similar to the UK, but where it may still be different enough to limit the generalisability of findings. The perspective included the employment insurer.

				Other			Incrementa	Incremental	Cost-	
5	Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty

- (cc) The main part of the intervention took 13 full days, 5 return-to-work sessions and 1 feedback session, all of which took place within 2 months. Each day's schedule consisted of four (1.5 hours) sessions: two physical sessions and two psychological sessions, twice a week supplemented with a fifth session consisting of 30 minutes of relaxation exercises.
- (dd) The study only had a 12 month time horizon which is insufficient to assess whether outcomes are sustained. Impact on QALYs was not considered. No deterministic or PSA was performed.
- (ee) The population includes people with 50% to 100% sick leave. The study was set in the Netherlands where the occupational support offered is differently organised to the UK.
- (ff) Vocational rehabilitation involved: assessing the impact of TBI on the participant, family and their roles; community reintegration training; pre-work training; liaison with employers, tutors or employment advisors.
- (gg) Local differences in service provision meant that this varied widely between participants, but potentially involved support from Headway (a voluntary organization providing advice and support to TBI people and their families), community occupational therapy (OT) or physiotherapy and routine GP follow-up.
- (hh) A 12 month time horizon was used which was insufficient to capture whether outcomes were sustained. QALY data were estimated using VAS. Effectiveness data were not derived from a RCT.
- (ii) Although this was a UK study, the population included students and those in unpaid employment. EQ-5D VAS data were used for utilities rather than health states valued by a population.
- (jj) The course focused on an early start of the intervention by OPs, in which they operated as an activating counsellor using CBT to enhance the problem-solving capacity of workers, especially in relation to their work environment. This consisted of clinical management according to the APA Guideline (2000) and antidepressants and/or CBT with senior psychiatric residents. Visits lasted 30 minutes every 2 to 3 weeks.
- (kk) A 12 month time horizon was used which was insufficient to capture whether outcomes were sustained. QALYs were not considered and neither were all healthcare costs.
- (II) The length of unemployment was unclear for study participants. The study was conducted in the Netherlands where occupational support is differently organised to the UK.
- (mm) TAU included antidepressants, if indicated and accepted by patients. They were treated by senior psychiatric residents with visits lasting 30 minutes every 2 to 3 weeks. OT consisted of two skilled occupational therapists providing the intervention over three manual-based phases: diagnostic phase (4 weeks) five contacts with an occupational physician from the patient's employer and a plan for work reintegration; therapeutic phase (24 weeks) 24 weekly group sessions (8 to 10 patients) and 12 individual sessions; follow-up phase (20 weeks) three individual visits.
- (nn) A 12 month time horizon was used which is insufficient to capture whether outcomes were sustained. QALYs were not considered. The source of costs was unclear.
- (oo) The population includes people with 50% to 100% sick leave. The study was set in the Netherlands where the occupational support offered is differently organised to the UK.
- (pp) Work modifications were defined as those based on participative ergonomics involving all relevant stakeholders.

			Other			Incrementa	Incremental	Cost-	
Study	Limitations	Applicability	comments	Costs	Effects	I cost	effects	effectiveness	Uncertainty

- (qq) This enabled the employer to become actively involved in the rehabilitation process. The PAEW intervention did not include a workplace assessment and work modifications, as was part of the WI.
- (rr) UC included 4.5 GP visits, 4.5 prescriptions, 3 packs of pain relief medication, 4 half hour sessions of physiotherapy (in 7% of cases), 2.5 sessions of osteopathy (in 5% of cases), 2.5 sessions of chiropractic treatment (in 2% of cases) and a hospital outpatient visit (in 10% of cases).
- (ss) Whilst a lifetime horizon was used in the model it was based on only 12 months of effectiveness data with outcomes not influenced by the intervention after 12 months. Costs, QALYs and incremental analysis were not reported and PSA was not performed.
- (tt) Although this is a UK study, it is based upon effectiveness studies that were conducted outside the UK and the authors stated that this fact may limit study generalisability. Costs and outcomes data were also not well reported.
- (uu) WI started at baseline, at least 8 weeks before sick-leave. The intervention consisted of: Dutch OP guidelines for LBP; A workplace assessment and work modifications based on participative ergonomics, which involved all important stakeholders: the occupational health service's ergonomist or occupational health nurse, the worker on sick-leave, the worker's supervisor and other communication between the OP and the GP, to reach consensus on counselling the worker in RTW.
- (vv) A graded activity programme based on operant behavioural therapy principles based on the findings from patient history, physical examination, functional capacity evaluation, the demands from the patients' work and the patients' expectations on time to RTW. The entire programme consisted of a maximum of 26 one-hour sessions, with a frequency of two sessions a week. The first session took half an hour more since taking the patients' history and a physical examination were part of this session. The programme ended as soon as a full RTW had been established, according to an earlier agreed upon individual schedule. During the programme the worker had an active role in RTW and the physiotherapist acted as a coach and supervisor, using a hands-off approach
- (ww)This included resuming daily activities, working within two weeks was encouraged and a clinical intervention recommended after 12 weeks.
- (xx) A 12 month time horizon was used which is insufficient to capture whether outcomes were sustained. QALYs were estimated using VAS.
- (yy) This study included people with 2 to 4 weeks of sick leave. The study was in the Netherlands where the occupational support offered is differently organised than the UK. EQ-5D VAS rather than 5 level health state was used for utility values.
- (zz) This was developed on the basis of three consultations over a time span of four weeks, and encompassed the following five key tasks: 1 diagnosing stress-related mental disorders; 2 providing education about the problem and importance of taking an active role in one's functional recovery; 3 advising patients on how to reflect, cope and problem-solve; 4 monitoring progress; 5 referring to specialists.
- (aaa) A 12 month time horizon was used which is insufficient to capture the sustainability of outcomes.
- (bbb) Participants were those with partial sick leave over six months. The study was in the Netherlands where the occupational support offered is differently organised than in the UK.
- (ccc) Three meetings were planned to take place within 2 weeks. The purpose of the first meeting between the sick-listed employee and the RTW coordinator was to identify obstacles for RTW from the perspective of the employee. The second meeting was between the supervisor and the RTW coordinator, where obstacles to the employee's RTW were identified from the supervisor's perspective. In the third meeting, which was generally the longest, the employee, supervisor and RTW coordinator discussed solutions and formulated a consensus-based plan for their implementation.

Appendix I – Health economic evidence profiles

Table 6: Health economic evidence profiles of studies included in the economic evidence review for workplace health interventions for RQ 1a, 2a and 3a

Study	Arends 2013			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Arends 2013 (Netherlands) Economic analysis: Cost-benefit analysis (CBA) and cost- effectiveness analysis (CEA) Study design: An economic evaluation using data directly from a trial. No economic model was constructed. Approach to analysis: An economic evaluation was conducted alongside a cluster- randomised controlled trial (RCT). Different costs and outcomes were used to undertake CEA (with prevention of an episode of recurrent sickness as the effectiveness measure) and CBA. Perspective: CEA: societal CBA: employer	Population: Workers (aged 18 to 63 years) who were diagnosed at start of sickness absence with a common mental disorder (CMD) and were now partially or fully ready to return to work. Cohort settings: Not applicable (NA) Intervention 1: SHARP-at work (n=80, mean age 41.3 years, 66.2% female). A five steps intervention return to work (RtW) was started with the occupational physician (OP) monitoring and supporting the person through the steps. a Intervention 2: Care as usual (CAU) (n=78, Mean age 42.3, 51.3% female): OP	Mean cost per person: SHARP CEA: €4,167 CBA: Between €29,337 (human capital approach (HCA) to productivity loss) and €37,215 (friction cost approach (FCA) to productivity loss) CAU CEA: €2,403 CBA: Between €24,607 (HCA) and €31,685 (FCA) Currency & cost year: Cost year not stated, Euros (€) Cost components incorporated: CEA: Health care costs for society and individual (e.g. GP, specialist, prescriptions and over the counter medications). CBA: Occupational health services for employer and	Recurrent sickness absence over 12 months SHARP: 39% CAU: 62%	Full incremental analysis CEA, incremental cost-effectiveness ratio (ICER), SHARP vs CAU, per 1% of recurrent sickness absence prevented: €10,605 CBA Employer occupational health costs only, SHARP was €800 greater than with CAU Productivity loss (HCA) SHARP vs CAU: €6,046 Productivity loss (FCA) SHARP vs CAU: €3,995 Analysis of uncertainty Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CEA. This showed that if the ICER is €20,000 per 1% reduction in recurrence of sickness absence there is an 84% chance that CAU would be cost-effective. Deterministic sensitivity analysis looked at plausibly cheaper costs of SHARP but found this made no significant difference to results. Sensitivity analyses CEA – Excluding an outlier, which was attributed to high costs due to hospitalisation in a psychiatric ward, an ICER of €-533 was calculated for the incidence of recurrent

Study	Arends 2013			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Time horizon: 12 months Treatment effect duration: Not relevant Discounting: Not conducted	followed an evidence based national guideline on helping people on sick leave with mental health problems return to work. ^b	productivity loss (sickness days adjusted by productivity loss from hours of absence measured using HCA and FCA).		sickness absence, indicating SHARP could be cost-effective. Reduced SHARP costs did not change the direction of the primary analyses. CBA – Reduced SHARP costs did not change these results.

Health outcomes: Within trial analysis using retrospective questionnaires. **Quality-of-life weights:** Not applicable. **Cost sources:** Underlying trial for resource use and published sources for costs.

Comments

Source of funding: A grant from Stichting Instituut GAK, a Dutch funding agency. **Limitations:** Author-recognised limitations: the data that was collected was self-reported and the accuracy could not be checked, 38% of patients not having completed cost data meaning the CEA and CBA could be underpowered. There was evidence that some people could not understand the productivity question. **Other:** None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: CAU: care as usual; CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; CMD: common mental disorder; FCA: friction cost approach; HCA: human capital approach; ICER: incremental cost-effectiveness ratio; OP: occupational physician; RCT: randomised controlled trial; RTW: return to work

- (a) The five steps comprised: (1) making an inventory of problems and/or opportunities encountered at work after RtW; (2) brainstorming about solutions/realisations; (3) writing down solutions/ realisations and the support needed and assessing the applicability of these solutions; (4) discussing solutions/ realisations and making an action plan with the supervisor; (5) evaluating the action plan/implementation of solutions.
- (b) OPs enacted the guideline of the Netherlands Society of Occupational Medicine "The treatment of workers with mental health problems by the OP". It is primarily aimed at structuring OPs' treatment to help sick-listed workers with mental health issues to RtW. Limited focus is given to follow-up after RtW has been achieved: only one consultation, to address relapse.

Study	Brouwers 2007				
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness	
Brouwers 2007 (Netherlands) Economic analysis: Cost-utility analysis (CUA) and cost-benefit analysis (CBA) Study design: An economic evaluation using data directly from a trial. No economic model was constructed. Approach to analysis: An economic evaluation was conducted alongside a randomised controlled trial (RCT) where healthcare and sick leave costs and utility values data were captured. Different costs and outcomes were used to undertake CUA and CBA. Perspective: CUA: societal CBA: public health insurer Time horizon: 18 months	Population: People (aged 18 to 60 years) absent from work on sick leave for no more than 3 months due to a minor mental disorder. Cohort settings: Total trial cohort Mean age: 40; female: 60% Intervention 1: Intervention group (IG) (n=95): 5 individual sessions (50 minutes each), covering 3 stages, with a social worker: cause, coping and implementation. a Intervention 2: Care As Usual (CAU)) (n=90): routine general practitioner (GP) care, which could include medication or counselling or referral.	Mean cost per person: IG: €14,493 (exclusive of intervention costs of €13,305 total for all people) CG: €14,482 CAU CEA: €2,403 CBA: €24,607 human capital approach (HCA) to €31,685 frictional cost analysis (FCA) Currency & cost year: Cost year not stated, Euros (€) Cost components incorporated: Health care costs for society (e.g. GP, specialist, prescriptions), productivity loss (sickness days adjusted by productivity loss from hours of absence measured using HCA and FCA. Intervention costs.	Mean quality-adjusted life-years (QALYs) per person IG vs CAU (Dutch EQ-5D): 0.056 higher IG vs CAU (UK EQ-5D): 0.044 higher Sick leave duration until full return to work IG: 152.7 days CAU: 156.5 days These were not statistically significantly different	Full incremental analysis CUA (based on 2,000 bootstrapped pairs) Probabilistic ICERs reported (ICERs are negative). IG vs CAU: €234 less expensive ICER, IG vs CAU (Dutch EQ-5D): -€,4179 ICER, IG vs CAU (UK EQ-5D): -€5,306 IG was less expensive and more effective than CAU CBA IG vs CAU: €11 more expensive (not statistically significant but did not include intervention costs) Analysis of uncertainty Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA. This demonstrated an even split between the north-east and south-east quadrants of the cost-effectiveness plane indicating that any difference in costs between IG and CAU were likely small. 52% of bootstrap estimates were in the south east quadrant where IG dominates CAU.	

Study	Brouwers 2007			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Treatment effect duration: Not relevant				
Discounting: Not conducted				

Health outcomes: The underlying trial. **Quality-of-life weights:** Health states derived using SF-36. Sources of utility values for SF-36 health states were not provided. **Cost sources:** The underlying trial for resource use and published sources for costs.

Comments

Source of funding: The Netherlands Organisation for Health Research and Development. **Limitations:** Author recognised limitations. Only sick leave duration was considered and not sick leave episodes after return to work. Indirect costs were not considered. Cost data were skewed by a few high cost individuals. All participants came from one part of Amsterdam. GPs in the CG may have heard of the intervention through their patients resulting in contamination of CG. **Other:** None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: CAU: care as usual; CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; CUA: cost-utility analysis; FCA: frictional cost approach; GP: general practitioner; HCA: human capital approach; ICER: incremental cost-effectiveness ratio; IG: intervention group; QALY: quality-adjusted life year; RCT: randomised controlled trial

(a) Treatment was over 10 weeks and entailed 3 stages: (i) understanding the cause of loss of control (ii) the development of problem-solving strategies; and (iii) their implementation.

Study	Bultmann 2009			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Bultmann 2009 (Denmark)	Population: Workers on sick leave for 4-12 weeks due to	Mean cost: CTWR (per person): \$31,144 over 12 months	Sickness absence hours, per person, over 12 months	Full incremental analysis CTWR vs CCM, over 12 months, per person: \$10,668 (\$1,548 more over 12 months without
Economic analysis: Cost-benefit analysis (CBA) and cost-	lower back pain (LBP) or musculoskeletal disorders (MSK).	(\$3,321 without productivity loss)	CTWR: 656.6 hours CCM: 997.3 hours	productivity loss) CEA

Study	Bultmann 2009				
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness	
Study design: An economic evaluation using data directly from a trial. No economic model was constructed. Approach to analysis: An economic evaluation was conducted alongside a randomised controlled trial (RCT) where healthcare costs and lost days of work were captured. CEA used averted absence days as the effectiveness measure. Perspective: Societal Time horizon: 12 months Treatment effect duration: Not relevant Discounting: Not conducted	Cohort settings: Not applicable (NA) Intervention 1: Coordinated and tailored work rehabilitation (CTWR) (n=60, mean age 44.2, 48.5% female) screening followed by a tailored rehabilitation plan developed by an interdisciplinary team. Intervention 2: Conventional case management (CCM) (n=47, mean age 42.9, 63.8% female): Conventional case management provided by the municipality. No further information given.	CCM (per person): \$41,812 over 12 months (\$1,773 without productivity loss) Currency & cost year: 2007 US\$ Cost components incorporated: CBA: Primary care and specialist treatment costs, productivity loss (sickness days adjusted by productivity loss from hours of absence measured using human capital approach (HCA)). Intervention costs. CEA: As for CBA but without productivity loss.		CTWR vs CCM, per absence day avoided (without productivity loss): \$33.7 Analysis of uncertainty CBA: One way deterministic sensitivity analysis of a doubling of intervention costs and 25% reduction in wages still resulted in cost savings	

Health outcomes: Within trial analysis using retrospective questionnaires. **Quality-of-life weights:** NA. **Cost sources:** Resource and cost data collected from the national Danish registries.

Comments

Study	Bultmann 2009			
	Population &	Costs	Health outcomes	Cost-effectiveness
Study details	interventions			

Source of funding: Danish National Labour Market Authority, Vejle County, and the Danish Chiropractic Research Fund. **Limitations:** Authors recognised limitations: the required sample size was not obtained; gender, education levels and prevalence of neck pain differed between intervention and control groups which may have influenced findings. There was significant loss to follow up (45% at 12 months in the control group). The trial was not blinded. **Other:** None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: CAU: care as usual; CBA: cost-benefit analysis; CCM: conventional case management; CEA: cost-effectiveness analysis; CG: control group; CTWR: coordinated and tailored work rehabilitation; FCA: frictional cost approach;; HCA: human capital approach; ICER: incremental cost-effectiveness ratio; IG: intervention group; LBP: low back pain; MSK: musculoskeletal disorder; NA: not applicable; RCT: randomised controlled trial; RtW: return to work

(a) After 4 to 12 weeks of sick leave: (1) work disability screening was conducted: a multidisciplinary assessment of disability and functioning and identification of barriers for RtW and (2) a coordinated, tailored and action-oriented work rehabilitation plan was developed by an interdisciplinary team with continuous feedback on the plan from the sick listed worker, the interdisciplinary team, the workplace, and major stakeholders. The interdisciplinary team consisted of an occupational physician, an occupational physiotherapist, a chiropractor, a psychologist, and a social worker who had the role of case worker.

Study	Finnes 2017				
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness	
Economic analysis: Cost-utilisation analysis (CUA) Study design: An economic evaluation using data directly from a trial. No economic model was constructed.	Population: Workers (at least 50% whole time equivalent (WTE)) with sickness absence due to anxiety, depression, stress or exhaustion Cohort settings: Age and gender split of trial not provided Intervention 1:	Total costs: Healthcare perspective (12 months) ACT: \$5,507 WDI: \$6,465 ACT+WDI: \$6,141 TAU: \$6,207 Societal perspective (12 months) ACT: \$14,452 WDI: \$15,649	Quality-adjusted life years (QALY) gains over 12 months ACT: 0.164 WDI: 0.122 ACT+WDI: 0.168 TAU: 0.155	Full incremental analysis Healthcare perspective, incremental costeffectiveness ratio (ICER) Both TAU and WDI were dominated by ACT ACT vs baseline: \$33,579 per QALY gained ACT+WDI vs ACT: \$158,500 per QALY gained Societal perspective Both TAU and WDI were dominated by ACT alone. Compared to ACT, ACT+WDI had an ICER of \$30804 per QALY gained	

Study	Finnes 2017			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Approach to analysis: An economic evaluation conducted alongside a randomised controlled trial (RCT) where healthcare costs, lost days of work and utility values were captured. Perspective: Healthcare and separate societal analysis including sickness benefit costs Time horizon: 12 months Treatment effect duration: Not relevant Discounting: Not conducted	Acceptance and Commitment Therapy (ACT) (n=89): a psychological intervention consisting of 6 x 60-minute sessions. a Intervention 2: Workplace dialogue intervention (WDI) (n=87): three meetings involving the participant plus work supervisor. b Intervention 3: ACT+WDI (n=88): conducted by two different therapists. c Intervention 4: Treatment as usual (TAU (n=88)): treatment as it was planned in a primary care centre or other care facility. d	ACT+WDI: \$17,066 TAU: \$15,593 Currency & cost year: 2015 US\$ Cost components incorporated: For healthcare perspective, intervention and costs of visits to health professionals (drug costs were excluded). For societal perspective, sickness benefit payments were added.		Analysis of uncertainty Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA. This showed that for ACT compared to ACT+WDI from both the healthcare and societal perspectives, the percentage of bootstrap iterations were spread roughly equally across all four quadrants, although approximately 60% of iterations in both perspectives had ACT+WDI more costly than ACT and 50% of iterations of ACT+WDI were more effective. Scenario analysis showed that using Swedish utility weights (rather than English in the base case) resulted in ACT being the dominant strategy. A second scenario explored the impact of using costs as if the intervention was delivered in a 'regular' setting in which case ACT would no longer dominate TAU but have an ICER of \$71 per QALY gained (healthcare perspective) and ACT+WDI had an ICER of \$286,000 per QALY gained compared to ACT.

Study	Finnes 2017			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness

Health outcomes: Within trial analysis using retrospective questionnaires. **Quality-of-life weights:** Health states were derived using EQ-5D and valued using UK valuation set (Swedish in the scenario analysis). **Cost sources:** The volumes of each cost category were obtained from study records, and unit costs were obtained from national public databases and websites.

Comments

Source of funding: REHSAM research fund (2011/12) and from the County Council in Stockholm, Sweden. **Limitations:** Author recognised limitations: it excluded the unemployed and self-employed people; data on healthcare resource use was retrospective and self-completed and so may have recall bias; the societal perspective did not include impacts on employers and drug costs were not included in either perspective; cost data were skewed towards a few individuals which has an impact where loss to follow up was not insignificant (although actual follow up rates were not reported). **Other:** None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: ACT: acceptance and commitment therapy; CAU: care as usual; CBA: cost-benefit analysis; CBT: cognitive behavioural therapy; CEA: cost-effectiveness analysis; CUA: cost-utilisation analysis; DALY: disability-adjusted life-year; ICER: incremental cost-effectiveness ratio; RCT: randomised controlled trial; RTW: return to work; TAU: treatment as usual; WDI: workplace dialogue intervention; WTE: whole time equivalent; QALY: quality-adjusted life year

- (a) The first part (sessions 1 to 3) emphasized mindfulness, cognitive defusion, and acceptance. During sessions 4 to 6, the focus was on exploring and clarifying personal values and committing to pursuing valued life activities.
- (b) The first step was an individual interview with the participant at the clinic followed by an interview with the participant's supervisor at the workplace. These meetings, lasting up to 60 minutes, aimed to investigate the participants' and the supervisors' views upon causes of the sickness absence, and what might facilitate RTW.
- (c) There was no integration or coordination of the two interventions, and no interaction between therapists.
- (d) This typically included psychotherapy, CBT and/or pharmacological treatments, physical therapy and counselling.

Study	Jensen 2005			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Jensen 2005 (Sweden)	Population:	Total costs:	Working days lost	Full incremental analysis
Economic analysis:	Blue-collar and service/care workers (aged 18 to 60 years)	At 3 years (mean cost per person): female; male BM: €107,703; €130,015	per year, pre- intervention: male; female	For women, BM, PT and CBT were all less expensive over 3 years vs TAU with BM having the lowest cost per person vs CBT

Study	Jensen 2005			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Study details Cost-benefit analysis (CBA) Study design: An economic evaluation using data directly from a trial. No economic model was constructed. Approach to analysis: An economic evaluation was conducted alongside a randomised controlled trial (RCT) where healthcare costs, lost days of work and disability pension costs were captured.	The state of the s		BM: 67; 72 PT: 76; 92 CBT:115; 109 TAU: 65; 80 Working days lost per year, post-intervention BM: 99; 123 PT: 95; 110 CBT:109; 101 TAU: 82; 86	BM vs CBT: €50,097 less expensive BM vs PT: \$82,057 less expensive BM vs TAU: €137,509 less expensive For men, CBT and PT were both more expensive over 3 years than TAU with BM being less expensive than TAU by €63,224 Analysis of uncertainty Not undertaken
Perspective: Societal Time horizon: 3 years Treatment effect duration: Not relevant Discounting: Not conducted	female) to improve pain management. b Intervention 3: Behavioural medicine (BM) (PT+CBT) (n=63, mean age: 43, 48% female) Intervention 4: Treatment-as-usual (TAU) (n=48, mean age: 44, 58% female):	of absence measured using human capital approach), disability pensions. Intervention costs.		

Study	Jensen 2005			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
	No additional interventions outside the normal routines in health care. This was not otherwise described.			

Health outcomes: Within trial analysis using retrospective questionnaires. Absence from work from National Social Insurance Board. **Quality-of-life weights:** SF-36 data were collected but not reported. **Cost sources:** Costs were from published sources but not clearly reported.

Comments

Source of funding: AFA Insurance and Alecta Insurance. **Limitations:** Author recognised limitations: there was a small sample size with low power and wide confidence intervals; the intention to treat results were not statistically significant. **Other:** None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: BM: behavioural medicine; CBA: cost-benefit analysis; CBT: cognitive behavioural therapy; GP: general practitioner;; NA: not applicable; PT: physiotherapy; RCT: randomised controlled trial

- (a) This consisted of approximately 20 scheduled hours per week aimed at enhancing the physical functioning and facilitating lasting behaviour change. Each participant was assigned to an individually tailored training program.
- (b) This was an average of 13 to 14 scheduled hours per week aimed at improving the participants' ability to manage their pain and resume a normal level of activity.

Study	Lambeek 2010				
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness	
Lambeek 2010 (Netherlands)	Population: Workers (aged 18 to 65 years) on full or	Total costs: At 12 months IC: £13,165 (£1,479 direct	Days until sustainable return to work	Full incremental analysis CEA £3 extra cost for every day earlier return to work	
Economic analysis:	partial sick leave for 12 weeks to two years	costs, £11,686 indirect costs)	IC: 129 UC: 197	with IC. Only direct costs were considered.	

Study	Lambeek 2010			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Cost-benefit analysis (CBA), cost- effectiveness analysis (CEA) and cost- utilisation analysis(CUA) Study design: An economic evaluation using data directly from a trial. No economic model was constructed. Approach to analysis: An economic evaluation conducted alongside a randomised controlled trial where primary and secondary healthcare costs, lost days of work and utility values were captured. CEA had sustainable return to work as effectiveness measure. Perspective: Societal Time horizon: 12 months	due to non-specific lower back pain (LBP) Cohort settings: Integrated Care (IC) (Intervention): mean age 45.5, 44% female Usual care (UC): mean age 46.8, 40% female Intervention 1: IC (n=66, mean age 45.5, 44% female), a graded activity protocol at the workplace. a Intervention 2: UC (n=68, mean age 46.8, 40% female): referred to occupational physician and GP with a letter containing the advice to treat them according to the Dutch guidelines for patients with LBP.	UC: £18,475 (£1,262 direct costs, £17,213 indirect costs) Currency & cost year: 2007 UK£ Cost components incorporated: Health care costs (primary and secondary care physicians and specialists, hospital stays and diagnostic tests, drug costs). Intervention costs. Productivity loss from absenteeism.	QALYs IC: 0.74 UC: 0.65	IC vs UC: dominant IC cost saving over UC, per person: £5,310 QALY gain, per person: 0.09 (direct and indirect costs considered) CBA IC, return on investment for every £1 spent: £26 (direct and indirect costs considered) Analysis of uncertainty Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA and CEA. This showed that in 98% of iterations IC dominated UC. For the CUA and for the CEA that if there was a willingness to pay of £10 for one day earlier return to work there was a 95% chance that IC was cost effective. Scenario analysis showed that if only complete cases were considered then there was no statistical difference in costs between IC and UC. A further scenario explored the impact of the intervention only for people aged under 55 years which resulted in a doubling in the costs of IC. A final scenario analysis showed that varying productivity levels did not impact on results.

Study	Lambeek 2010			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Treatment effect duration: Not relevant				
Discounting: Not conducted				

Health outcomes: Within trial analysis was used, using retrospective questionnaires. Absence from work data were from the National Social Insurance Board. **Quality-of-life weights:** SF-36 data were collected but not reported beyond a statement that there was no difference between groups. **Cost sources:** Costs were from published sources but not clearly reported.

Comments

Source of funding: AFA Insurance and Alecta Insurance. **Limitations:** Author recognised limitations: small sample size with low power and wide confidence intervals; intention to treat results were not statistically significant. **Other:** None

Overall applicability: Partially applicable

Overall quality: Potentially serious limitations

Abbreviations: CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; CUA: cost-utility analysis; IC: ; LBP: low back pain; UC: usual care; QALY: quality-adjusted life year

(a) Workers were referred to a clinical occupational physician who was responsible for the coordination of the care and for communication with the other healthcare professionals in the team. The workplace intervention protocol formulated a consensus based plan for adaptations at work to facilitate return to work. The integrated care team consisted of a medical specialist, occupational therapist, physiotherapist, and clinical occupational physician.

Study	Loisel 2002			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Loisel 2002 (Canada)	Population:	Total costs:	Mean number of DFB	Full incremental analysis
	Workers absent for	At 12 months		CEA (cost per DFB saved compared to UC)
Economic analysis:	more than 4 weeks	CI: \$12,038	At 12 months	
Cost-benefit analysis	with occupational back	OI: \$9,569	CI: 114.9	At 12 months
(CBA) and cost-	pain	Sherbrooke: \$12,137	OI: 116.1	CI: \$187.40

Study Study details	Loisel 2002			
	Population & interventions	Costs	Health outcomes	Cost-effectiveness
effectiveness analysis (CEA) Study design: Economic evaluation using data directly from a trial. No economic model was constructed. Approach to analysis: Economic evaluation conducted alongside a randomised controlled trial where healthcare and sick pay costs were captured. CEA had days on full benefits (DFB) as effectiveness measure. Perspective: Health and employment insurer Time horizon: 6.4 years Treatment effect duration: Not relevant Discounting: Not conducted	Cohort settings: Mean age ranged from 37.4 for "Sherbrooke Model" to 44.5 with occupational intervention (OI). Gender split ranged from 60% female with "Sherbrooke model" to 19% with usual care (UC). The study had fewer than 30 people in each study arm and no statistical significance testing of results was performed Intervention 1: OI (n=22, mean age: 44.5, 41% female): visits to the study occupational medicine physician and a participatory ergonomics intervention with the study ergonomist, the injured worker, there supervisor, and management and	At 6.4 years CI: \$16,902 OI: \$16,252 Sherbrooke: \$14,494 UC: \$33,079 Currency & cost year: 1998 Canadian\$ Cost components incorporated: Health care costs, intervention costs and income replacement costs	Sherbrooke: 115.9 UC: 126.9 At 6.4 years CI: 178.7 OI: 228.0 Sherbrooke: 125.6 UC: 418.3	OI: -\$20.40 (dominated UC) Sherbrooke: \$213.50 At 6.4 years CI: -\$67.50 (dominated UC) OI: -\$88.40 (dominated UC) Sherbrooke: -\$63.50 (dominated UC) CBA, cost differential compared to UC At 12 months CI: -\$2,250 OI: \$220 Sherbrooke: -\$2,348 At 6.4 years CI: \$16,176 OI: \$16,827 Sherbrooke: \$18,585 Analysis of uncertainty Sensitivity analyses were performed but varying the total healthcare costs by 60% to 190% and income per capita by 85% to 125%. Over a 6.4 year time period all interventions remained dominant compared to UC over the cost ranges considered.

Study	Loisel 2002			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
	union representatives.			
	Intervention 2: Clinical rehabilitation intervention (CRI) (n=31, mean age: 40.2, 42% female)): with a back pain specialist and potentially a multidisciplinary work rehabilitation intervention at 12 weeks of absence. b			
	Intervention 3: OI+CRI ("Sherbrooke Model") (n=10, mean age: 37.4, 60% female))			
	Intervention 4: Usual care (UC) (n=26, mean age: 41.6, 19% female): with worker's physician receiving no advice about return to work.			

Study	Loisel 2002			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness

Health outcomes: Quebec Workers Compensation Database. Quality-of-life weights: NA. Cost sources: Quebec Workers Compensation Database.

Comments

Source of funding: Institut de Recherche en Santé et Sécurité au Travail du Québec (IRSST). **Limitations:** Author recognised limitations Salaries in control arm were higher than intervention arms. Costs of job modifications were not recorded. Workers may have had subsequent back problems not considered to be work related. **Other:** None

Overall applicability: Partially applicable Overall quality: Minor limitations

Abbreviations: CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; CI: clinical intervention; CRI: clinical rehabilitation intervention; DFB: days on full benefits; OI: occupational intervention; UC: usual care

- (a) This participatory ergonomics intervention was limited in scope and duration with job modifications recommended to the employer who was at liberty to implement them or not.
- (b) Clinical examination by a back pain medical specialist, participation in a back school after eight weeks of absence from regular work and, if necessary, a multidisciplinary work rehabilitation intervention after 12 weeks of absence from work.

Study	Meijer 2006			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Meijer 2006 (Netherlands) Economic analysis: Cost-effectiveness analysis (CEA) Study design: An economic evaluation using data directly from a trial. No economic model was constructed.	Population: Bank and university workers on at least 50% contracts with 50% sick leave in last 4 to 20 weeks due to non-specific upper extremity musculoskeletal disorders (MSDs) Cohort settings:	Total costs per week 2 months MDT: €1,335 UC: €448 6 months MDT: €664 UC: €359 12 months	Percentage of regular hours worked Baseline MDT: 29% UC: 29% 6 months MDT: 82% UC: 72%	MDT was more expensive compared to UC and did not increase the proportion of days worked. MDT was not cost-effective. Analysis of uncertainty Not undertaken

Study	Meijer 2006	Meijer 2006				
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		
Approach to analysis: An economic evaluation was conducted alongside a randomised controlled trial (RCT) where healthcare costs, productivity losses (days of work lost and time at work lost from extra breaks) and free time lost were captured. CEA used half days returned to work as the effectiveness measure. Perspective: Societal Time horizon: 12 months Treatment effect duration: Not relevant Discounting: Not conducted	Intervention 1: Multidisciplinary treatment (MDT) (n=20, mean age: 38.3, 70% female): an outpatient training programme carried out at Dutch rehabilitation centres. a Intervention 2: Usual care (UC) (n=14, mean age: 37.9, 64% female): supervision by occupational health services. UC could include treatment at the workplace and in the regular health care system, initiated by a general practitioner, or medical specialist.	MDT: €430 UC: €315 Currency & cost year: Euro (€) 2004 Cost components incorporated: Health care costs (medical services and medications), productivity loss (sickness days adjusted by productivity loss from hours of absence measured using human capital approach (HCA)), free time costs. Intervention costs.	12 months MDT: 86% UC: 73% There was no statistical difference between MDT and UC at any time point.			

Health outcomes: Within trial analysis using retrospective questionnaires. **Quality-of-life weights:** NA. **Cost sources:** Dutch board on medical tariffs, Dutch medicines compensation system, self-reported costs in questionnaire.

Comments

Study	Meijer 2006			
	Population &	Costs	Health outcomes	Cost-effectiveness
Study details	interventions			

Source of funding: The Netherlands Organization for Health Research and Development (ZONMw) and a supplementary grant from the UWV. **Limitations:** Author recognised limitations: the study was not adequately powered failing to recruit the targeted number of participants. **Other:** None

Overall applicability: Partially applicable
Overall quality: Potentially serious limitations

Abbreviations: CEA: cost-effectiveness analysis; HCA: human capital approach; MDT: multidisciplinary treatment; MSD: musculoskeletal disorders; NA: not available; RCT: randomised controlled trial; UC: usual care

(a) The main part of the intervention took 13 full days, 5 return-to-work sessions and 1 feedback session, all of which took place within 2 months. Each day's schedule consisted of four (1.5 hours) sessions: two physical sessions and two psychological sessions, twice a week supplemented with a fifth session consisting of 30 minutes of relaxation exercises.

Study	Radford 2012			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Radford 2012 (UK) Economic analysis: Cost-benefit analysis (CBA), cost- effectiveness analysis (CEA) and cost- utilisation analysis (CUA) Study design: An economic evaluation using data directly from a cohort study. No economic model was constructed.	Population: Patients in paid or voluntary work or in full time education hospitalised for at least 48 hours due to traumatic brain injury (TBI) Cohort settings: Total cohort: male (80%), mean age 34.3 Intervention 1: TBI Vocational Rehabilitation (TBI-	At 12 months (health and social care perspective) TBI-VR: £2,106.94 UC: £2,031.71 At 12 months (societal perspective) TBI-VR: £8,786 UC: £10,648 Currency & cost year: 2007 UK£ Cost components incorporated:	Quality-adjusted life years (QALYs) at 12 months TBI-VR: 0.1938 UC: 0.1763 Return to work or education at 12 months TBI-VR: 75% UC: 60%	Per person returned to work, TBI-VR vs UC (health and social care perspective): £501.33 Societal perspective: TBI-VR was more effective and saved money CBA From health and social care perspective, TBI-VR vs UC was £75.23 more costly. From societal perspective TBI-VR vs UC was £1,863 less expensive. Neither difference was statistically significant CUA TBI-VR vs UC, health and social care perspective: £4,299 per QALY gained.

Study	Radford 2012			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Approach to analysis: Economic evaluation conducted alongside a trial where healthcare, return to work or study outcomes and utility values were captured. The intervention was compared to usual care by looking at costs and outcomes in patients in surrounding areas. CEA had return to paid or voluntary work or study as outcome. Perspective: CBA: health and social care CEA and CUA: societal Time horizon: 12 months Treatment effect duration: Not relevant Discounting: Not conducted	VR), provided by an occupational therapist. Intervention 2: Usual care (UC) (n=54): participants in hospitals without TBI-VR. b	Health and social care perspective: Social worker and rehabilitation therapy costs, primary care costs. Societal perspective added included lost wages for participant and carer and benefits advisor costs.		Neither the QALY gain nor cost difference were statistically significant Analysis of uncertainty Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CUA. The boot strapped incremental cost-effectiveness ratio (ICER) was £2,567 lower than the deterministic ICER. Sensitivity analysis using imputed data for missing values more than doubled the cost per person returned to work in the CEA and increased the ICER per QALY gained in the CUA to £35,873 with TBI-VR.

Study	Radford 2012					
	Population &	Costs	Health outcomes	Cost-effectiveness		
Study details	interventions					

Health outcomes: Within trial analysis using retrospective questionnaires. **Quality-of-life weights:** EQ-5D visual analogue scale (VAS). **Cost sources:** NHS reference costs, PSSRU and Jobcentre plus.

Comments

Source of funding: College of Occupational Therapists. **Limitations:** Author recognised limitations: Incomplete follow up data; not an RCT; not properly powered resulting in wide confidence intervals. **Other:** There were no statistically significant results. It was difficult to isolate the effect of the intervention from the wider effects of the MDT that operates within Nottingham. It included people who worked, were students and were unemployed.

Overall applicability: Not applicable Overall quality: Potentially serious limitations

Abbreviations: CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; CMD: ; CUA: cost-utilisation analysis; GP: general practitioner; ICER: incremental cost-effectiveness ratio; MDT: multidisciplinary treatment; NHS: national health service; OT: occupational therapy; PSSRU: Personal Social Services Research Unit; QALY: quality-adjusted life year; RCT: randomised controlled trial; TBI: traumatic brain injury; UC: usual care; VAS: visual analogue scale; VR: vocational rehabilitation;

- (a) Vocational rehabilitation involved: assessing the impact of TBI on the participant, family and their roles; community reintegration training; pre-work training; liaison with employers, tutors or employment advisors.
- (b) Local differences in service provision meant that this varied widely between participants, but potentially involved support from Headway (a voluntary organization providing advice and support to TBI people and their families), community occupational therapy (OT) or physiotherapy and routine GP follow-up.

Study	Rebergen 2009					
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		
Rebergen 2009 (Netherlands) Economic analysis: Cost-benefit analysis (CBA) and cost- effectiveness analysis (CEA)	Population: Police workers on sick leave due to common mental disorders (CMDs) Cohort settings: Not applicable (NA)	At 12 months (health care perspective) GBC: €2,145 UC: €2,664 At 12 months (societal perspective using HCA for productivity loss)	Days of sick leave GBC: 113 UC: 114 These were not statistically significantly different	CEA GBC vs UC, incremental cost-effectiveness ratio (ICER) per sick day avoided: -€736 CBA Estimated net monitory benefit of GBC, per person: €3,582		

Study	Rebergen 2009			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Study design: Economic evaluation using data directly from a trial. No economic model was constructed. Approach to analysis: Economic evaluation conducted alongside a randomised controlled trial where healthcare costs, days of sick leave and productivity loss were captured. CEA had days of sick leave as outcome. Perspective: CEA: societal CBA: employer Time horizon: 12 months Treatment effect duration: Not relevant Discounting: Not conducted	Intervention 1: Guideline based care (GBC) (n=125, mean age: 38.8, 48.8% female), treatment by OPs according to the Dutch guideline for workers with mental health problems. a Intervention 2: Usual care (UC) (n=115, 40.0, 39.5% female): minimal involvement of the OP and easy access to counselling by a psychologist.	Traumatic brain injury - vocational rehabilitation (TBI-VR): €14,114 UC: €14,202 No costs were statistically significantly different Currency & cost year: 2003 Euros (€) Cost components incorporated: Healthcare perspective: Primary care (general practitioner (GP) visits, tests and medications), occupational healthcare, hospital care and psychological treatment. Societal perspective added in productivity loss using human capital approach (HCA) approach. Frictional cost method (FCM) approach was also used but only partially reported.		Outcomes are similar between GBC and UC, but direct costs were lower with GBC. The authors concluded that GBC could be cost-effective. Analysis of uncertainty Stochastic uncertainty in the data were dealt with using nonparametric bootstraps for CEA. The iterations showed there was never more than a 50% chance of GBC being cost effective per day of sick leave avoided regardless of the value of the day of work lost Different approaches to measuring productivity loss were analysed but did not affect the main findings.

Study	Rebergen 2009					
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		
Data sources						

Health outcomes: Within trial analysis using retrospective questionnaires. **Quality-of-life weights:** EQ-5D visual analogue scale (VAS). **Cost sources:** NHS reference costs, PSSRU and Jobcentre plus.

Comments

Source of funding: College of Occupational Therapists. **Limitations:** Author recognised limitations: Incomplete follow up data, not an RCT and not properly powered resulting in wide confidence intervals. **Other:** No statistically significant results. Difficult to isolate effect of the intervention from the wider effects of the MDT that operates within Nottingham. Included people who worked, were students and were unemployed.

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; CMD: common mental disorder; GBC: guideline based care; GP: general practitioner; FCM: friction cost method; HCA: human capital approach; ICER: incremental cost-effectiveness ratio; NA: not available; OP: occupational physician; TBI: traumatic brain injury; UC: usual care; VR: vocational rehabilitation

(a) The course focused on an early start of the intervention by OPs, in which they operated as an activating counsellor using CBT to enhance the problem-solving capacity of workers, especially in relation to their work environment. This consisted of clinical management according to the APA Guideline (2000) and antidepressants and/or CBT with senior psychiatric residents. Visits lasted 30 minutes every 2 to 3 weeks.

Study	Schene 2007					
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		
Schene 2007 (Netherlands) Economic analysis: Cost-benefit analysis	Population: Workers (aged 18 years or over) with at least 50% absence over 10 weeks to 2 years due to work	At 12 months (cost of intervention only) OT: \$3,149 TAU: \$1,891	No health or employment outcomes reported beyond earnings over 12 month period	Difference in total earnings minus costs of intervention (cost of intervention only) OT vs TAU: \$3,952 higher (not statistically significant)		
(CBA) Study design: An economic evaluation using data directly from	related major depressive disorder (WRMDD)	Currency & cost year: US\$, cost year not reported Cost components incorporated:		Analysis of uncertainty The base case was a bootstrapped analysis to account for stochastic uncertainty. The only sensitivity analysis performed was on the value of an hours work. As the value reduces the		

Study	Schene 2007			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
a trial. No economic model was constructed. Approach to analysis: An economic evaluation conducted alongside a randomised controlled trial (RCT) where healthcare costs (outpatient, GP and medication) and hours of work were captured. For the CBA, hours of work were multiplied by Dutch average hourly wage. Perspective: Societal Time horizon: 12 months for economic evaluation Treatment effect duration: Not relevant Discounting: Not conducted	Cohort settings: Not applicable (NA) Intervention 1: Treatment as usual (TAU) (n=30, mean age: 45.2, 53% female): out-patient treatment for depression. This consisted of clinical management according to the American Psychiatric Association Guideline (2000) and antidepressants and/or cognitive behavioural therapy (CBT) with senior psychiatric residents. Visits lasted 30 minutes every 2–3 weeks. Intervention 2: TAU + occupational therapy (OT) (n=32, mean age: 46.6, 50% female), the addition of OT, which had	Outpatient treatments, OT, medications, GP visits, hospitalisations, travelling and parking and earnings		probability of that OT is more cost effective than TAU falls. In the base case it is 75.5% at \$36.88 per hour and falls to 52.5% at \$10 per hour.

Study	Schene 2007				
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness	
	diagnostic and therapeutic phases. ^a				

Health outcomes: Within trial analysis were used, using retrospective questionnaires. Quality-of-life weights: NA. Cost sources: Not reported.

Comments

Source of funding: Landelijk Instituut Sociale Verzekering (LISV). Limitations: Author recognised limitations: small sample size and limited follow up data.

Other: None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: CBA: cost-benefit analysis; CBT: cognitive behavioural therapy; GP: general practitioner; NA: not applicable; OT: occupational therapy; RCT: randomised controlled trial; TAU: treatment as usual; WRMDD: work related major depressive disorder;

(a) TAU included antidepressants, if indicated and accepted by patients. They were treated by senior psychiatric residents with visits lasting 30 minutes every 2 to 3 weeks. OT consisted of two skilled occupational therapists providing the intervention over three manual-based phases: diagnostic phase (4 weeks) – five contacts with an occupational physician from the patient's employer and a plan for work reintegration; therapeutic phase (24 weeks) – 24 weekly group sessions (8 to 10 patients) and 12 individual sessions; follow-up phase (20 weeks) – three individual visits.

Study	Squires 2011				
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness	
Squires 2011 (UK)	Population: Workers on sick leave for 1 week to 6 months	Not reported	Not reported	CUA (from societal perspective so includes costs to NHS and from lost wages)	
Economic analysis: Cost-utilisation analysis (CUA)	due to musculoskeletal disorders (MSDs)	Currency & cost year: 2007 UK£		WI and PAEW are both cheaper than UC and more effective	
Study design: An economic evaluation using a Markov model using data from	Cohort settings: Mean age: 41; gender not reported	Cost components incorporated: Costs to NHS of MSD: general practitioner (GP) visits, prescriptions, allied		PAI is more costly but more effective than UC PAEW dominates all interventions. CEA, cost per sick day avoided	

Study	Squires 2011			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
published studies on three interventions Approach to analysis: Economic evaluation using effectiveness data from previously published studies, utilities from the British Household Panel Survey and costs from published sources. Perspective: NHS and person shaped support (PSS), societal and employer. Time horizon: Lifetime Treatment effect duration: If an individual did not return to work in 6 months, the probability of return to work was assumed to be same for both interventions and usual care. Discounting: Not	Intervention 1: Workplace intervention (WI): a workplace assessment and work modifications based on participative ergonomics involving all relevant stakeholders. Intervention 2: Physical activity and education intervention (PAE): any form of physical activity and education around how to deal with pain and body mechanics. Intervention 3: Physical activity, education and workplace visit (PAEW): WI+PAE plus a workplace visit by the employee and the physical therapist to inform rehabilitation. a	health professionals, hospital outpatients, Cost of interventions, salaries.		Results were presented on the costeffectiveness plane and not in a detailed table or text. Analysis of uncertainty Sensitivity and scenario analyses were undertaken. PAEW was not dominant if only the employer perspective was taken and the probability of sick leave recurring was doubled. In a threshold analysis, if the intervention costs were less than an additional £3,000 and returns at least an additional 3% of people to work (32/1,000) in comparison to UC, then it is likely to result in a cost per QALY gained below £20,000.
conducted				

Study	Squires 2011					
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		
	Usual care (UC): treatment of MSDs in the UK ^b					

Health outcomes: From published studies. **Quality-of-life weights:** SF-36 data from British Household Panel Survey. **Cost sources:** DWP, PSSRU and published sources.

Comments

Source of funding: National Institute for Health and Care Excellence. **Limitations:** Author recognised limitations: Evidence on effectiveness of interventions was poor and not necessarily generalizable to the UK. No long term (post 12 month) follow up was available. No probabilistic sensitivity analysis was possible. Large amount of structural uncertainty. Relationships between variables was simplified. **Other:** None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: CEA: cost-effectiveness analysis; CUA: cost-utilisation analysis; GP: general practitioner; MSD: musculoskeletal disorders; NHS: national health service; PAEW: physical activity, education and workplace visit; PAE: physical activity and education intervention; PSS: person shaped support; QALY: quality-adjusted life year; UC: usual care; WI: workplace intervention;

- (a) This enabled the employer to become actively involved in the rehabilitation process. The PAEW intervention did not include a workplace assessment and work modifications, as was part of the WI.
- (b) UC included 4.5 GP visits, 4.5 prescriptions, 3 packs of pain relief medication, 4 half hour sessions of physiotherapy (in 7% of cases), 2.5 sessions of osteopathy (in 5% of cases), 2.5 sessions of chiropractic treatment (in 2% of cases) and a hospital outpatient visit (in 10% of cases).

Study	Steenstra 2006	Steenstra 2006				
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		
Steenstra 2006 (Netherlands) Economic analysis:	Population: Workers on sick leave for 2 to 6 weeks due to lower back pain (LBP)	Mean total costs WI: €8,993 CI: €10,537 WI+CI: €12,391	Actual quality-adjusted life years (QALY) values not reported	CEA (per one day less of sick leave) WI vs UC: €19 WI+CI vs WI: €11		
	Cohort settings:	UC: €9,109	Sick leave (calendar days)	CI vs UC: €29		

Study	Steenstra 2006			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Cost-effectiveness analysis (CEA) and cost-utilisation analysis (CUA) Study design: An economic evaluation using data directly from a trial. No economic model was constructed. Approach to analysis: An economic evaluation was conducted alongside a randomised controlled trial (RCT) where healthcare costs, productivity losses and utility values were captured. CEA had days sick leave as outcome. Perspective: Societal Time horizon: 52 weeks Treatment effect duration: Not relevant Discounting: Not conducted	Intervention 1: Clinical intervention (CI) (n=28, mean age: 39.2, 79% female): a graded activity programme of 26 x 1-hour sessions, with a frequency of 2 sessions per week. a Intervention 2: Workplace intervention (WI) (n=96, mean age: 44, 47% female): UC, a workplace assessment and modification and communication between OP and GP in order to discuss how to counsel the worker to RtW. b Intervention 3: Usual care (UC) (n=100, mean age: 41.2, 67% female): Dutch OP guidelines for LBP delivered by a	Currency & cost year: Euro (€), cost year not stated Cost components incorporated: Direct healthcare costs (occupational physician, allied health professionals, hospitalisations), interventions, absenteeism	WI: 108.5 CI: 155.9 WI+CI: 181.7 UC: 135.2	CUA (cost per QALY) WI vs UC: -€1483 WI+CI vs WI: €24416 CI vs UC: €5447 Analysis of uncertainty The base case was a bootstrapped analysis to account for stochastic uncertainty. The cost-effective planes (and confidence intervals of point estimates) suggested that WI and UC were likely similar in cost but that WI was more effective. CI is likely less effective and more costly than both WI and UC. Scenario analyses suggested using a fixed sum per day of production lost, net rather than calendar sick days and using a HCA approach to productivity loss did not significantly influence results.

Study	Steenstra 2006					
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		
	GP. Resuming daily activities and work within two weeks is encouraged and a clinical intervention recommended after 12 weeks.					
	Intervention 4: WI + CI (n=27, mean age: 43.6, 47% female) where workers had WI in the first 8 weeks.					

Data sources

Health outcomes: Within trial analysis was used using retrospective questionnaires and computerised medical records. **Quality-of-life weights:** EQ-5D VAS. **Cost sources:** Dutch costing guidelines, health care charges and professional organisations.

Comments

Source of funding: The Netherlands Organisation for Health Research and Development (ZonMw), Dutch Ministries of Health, Welfare and Sports and of Social Affairs. **Limitations:** None discussed. **Other:** None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: CEA: cost-effectiveness analysis; CI: clinical intervention; CUA: cost-utilisation analysis; GP: general practitioner; LBP: lower back pain; LP: lumbar puncture; NA: not applicable; OP: occupational therapy; PT: physiotherapy; RCT: randomised controlled trial; RTW: return to work; UC: usual care; WI: work intervention

(a) A graded activity programme based on operant behavioural therapy principles based on the findings from patient history, physical examination, functional capacity evaluation, the demands from the patients' work and the patients' expectations on time to RTW. The entire programme consisted of a maximum of 26 one-hour sessions, with a frequency of two sessions a week. The first session took half an hour more since taking the patients' history and a physical examination were part of this session. The programme ended as soon as a full RTW had been established, according to an earlier agreed upon individual schedule. During the programme the worker had an active role in RTW and the physiotherapist acted as a coach and supervisor, using a hands-off approach.

Study	Steenstra 2006					
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		

(b) The WI started at baseline, at least before 8 weeks of sick-leave. The intervention consisted of: Dutch OP guidelines for LBP; A workplace assessment and work modifications based on participative ergonomics, which involved all important stakeholders: the occupational health service's ergonomist or occupational health nurse, the worker on sick-leave, the workers supervisor and other communication between the OP and the GP, to reach consensus on counselling the worker in RTW.

Study	Uegaki 2010			
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness
Uegaki 2010 (Netherlands) Economic analysis: Cost-utilisation analysis	Population: Workers with partial sick leave over 3 months due to distress	At 12 months MISS: €12,538 UC: €12,722 Currency & cost year:	Quality-adjusted life years (QALYs) over 12 months MISS: 0.78	Incremental cost-effectiveness ratio (ICER) MISS vs UC: -€7,356 per QALY gained Neither change in costs nor change in QALYs were statistically significantly different between MISS and UC.
(CUA)	Cohort settings: Not applicable (NA)	Euro (€) 2004	UC: 0.76	Analysis of uncertainty
Study design: An economic evaluation using data directly from a trial. No economic model was constructed. Approach to analysis: An economic evaluation was conducted alongside a randomised controlled trial (RCT) where healthcare costs, productivity losses and	Intervention 1: Minimal intervention for stress-related mental disorders with sick leave (MISS) (n=109, mean age: 42.0, 67% female): a GP customized version of an activating approach a Intervention 2:	Cost components incorporated: Primary care costs (general practitioner (GP), diagnostic tests, psychologist). Secondary care costs (mental health care, medical specialists, home help). Occupational physician (OP). Productivity losses (sick leave per hour assessed using frictional cost method (FCM) approach to productivity		The base case was a bootstrapped analysis to account for stochastic uncertainty. Cost-effectiveness planes showed that in the base case 77% of bootstrapped pairs would be considered cost-effective at a willingness to pay (WTP) threshold of €25,600 per QALY. Sensitivity analysis explored different approaches to costing lost productivity but did not significantly influence the results. Subgroup analysis suggested MISS may be most cost-effective for patients with stress

Study	Uegaki 2010					
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		
utility values were captured.	Usual care (UC) (n=83, mean age: 39.6, 65% female):	loss). Intervention costs (training for MISS).		related mental disorders, which was the only analysis which had statistically significant improvement in QALYs vs UC: -€28,278		
Perspective: Societal	comparable to usual care in real life					
Time horizon: 12 months	managed by a general practitioner (GP). No further information given.					
Treatment effect duration: Not relevant	9.70111					
Discounting: Not conducted						

Data sources

Health outcomes: Within trial analysis using retrospective questionnaires and computerised medical records. **Quality-of-life weights:** EQ-5D using Dutch tariff. **Cost sources:** Dutch costing guidelines, health care charges and professional organisations.

Comments

Source of funding: Health Research and Development Council (ZonMw) in The Netherlands. **Limitations:** Author recognised limitations
Usual care may not have mirrored the real world, lack of statistical power from small sample size, retrospective nature of data collection, QALYs may not be an appropriate estimate of benefit for patients with mental health problems. Presenteeism was not measured. **Other:** None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: CUA: cost-utilisation analysis; FCM: friction cost method; GP: general practitioner; ICER: incremental cost-effectiveness ratio; MISS: minimal Intervention for stress-related mental disorders with sick leave; NA: not applicable; OP: occupational physician; QALY: quality-adjusted life year; RCT: randomised controlled trial; UC: usual care; WTP: willingness to pay

(a) This was developed on the basis of three consultations over a time span of four weeks, and encompassed the following five key tasks: 1 diagnosing stress-related mental disorders; 2 providing education about the problem and importance of taking an active role in one's functional recovery; 3 advising patients on how to reflect, cope and problem-solve; 4 monitoring progress; 5 referring to specialists.

Study	van Oostrom 2009			
	Population & interventions	Costs	Health outcomes	Cost-effectiveness
(Netherlands) Economic analysis: Cost-benefit analysis (CBA), cost- effectiveness analysis (CEA) and cost- utilisation analysis (CUA) Study design: An economic evaluation was used, using data directly from a trial. No economic model was constructed. Approach to analysis: An economic evaluation was conducted alongside a randomised controlled trial (RCT) where healthcare, occupational health services, productivity	Population: Workers on sick leave for 2 to 8 weeks due to distress. Cohort settings: Not applicable (NA) Intervention 1: Workplace intervention (WI) (n=73, mean age 48.6, 23.3% female): consisted of a stepwise communication process to identify and solve obstacles to return to work (RTW). a Intervention 2: Usual care (UC) (n=72, mean age 49.2, 19.4% female): treatment by the OP according to the Dutch Guidelines. b	Societal perspective (12 months) WI: €3,201 UC: €2,758 Employer perspective (12 months) WI: €1,386 UC: €802 Currency & cost year: Euro (€) 2008 Cost components incorporated: Societal perspective: healthcare costs (primary care, occupational physician, home healthcare, medication, allied health professionals). Productivity loss (Human capital approach (HCA) and friction cost analysis (FCA) approaches). Employer perspective: occupational health services provided by employer, productivity loss as per societal perspective.	Mean duration of sick leave, over 12 months CEA WI: 133 days UC: 134 days Mean quality-adjusted life year (QALY) CUA WI: 0.77 UC: 0.78	CEA WI vs UC: £627 per sick day avoided Change in costs or change in sick days was not statistically different between WI and UC CBA Net Monetary Benefit WI (HCA): -€1,987 WI (FCA): -€1,700 WI was statistically significantly more costly than UC and changes in costs of productivity loss whilst favouring UC were not statistically significant regardless of productivity measure. CUA WI vs UC, incremental cost-effectiveness ratio (ICER) (HCA): -€18,4562 per QALY gained WI vs UC, ICER (FCA): -€155,850 WI dominates UC. Neither cost differences with WI or QALY gains were statistically significant. Analysis of uncertainty The base case was a bootstrapped analysis to account for stochastic uncertainty. Cost effectiveness planes showed substantial uncertainty in results which reflects the statistical uncertainty in the point estimates of cost differences and effectiveness measures between WI and UC.

Study	van Oostrom 2009					
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness		
Perspective: CEA and CUA: societal CBA: employer				Subgroup analysis suggested WI may be most cost-effective for patients with an intention to return to work but findings were still limited in statistical significance.		
Time horizon: 12 months						
Treatment effect duration: Not relevant						
Discounting: Not conducted						

Data sources

Health outcomes: Within trial analysis using retrospective questionnaires and computerised medical records. **Quality-of-life weights:** EQ-5D using Dutch tariff. **Cost sources:** Dutch costing guidelines, health care charges and professional organisations.

Comments

Source of funding: Dutch Ministry of Social Affairs and Employment and the participating occupational health services. **Limitations:** Author recognised limitations: costs of workplace adaptations were not registered. Presenteeism was not considered. Small sample size with effects skewed to a small number of costly participants. 20 out of 73 participants did not receive the WI. **Other:** None

Overall applicability: Partially applicable Overall quality: Potentially serious limitations

Abbreviations: CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; CUA: cost-utilisation analysis; FCA: frictional cost approach; HCA: human capital approach; ICER: incremental cost-effectiveness ratio; NA: not applicable; OP: occupational physician; QALY: quality-adjusted life year; RCT: randomised controlled trial; RTW: return to work; UC: usual care; WI: work intervention

- (a) Three meetings were planned to take place within 2 weeks. The purpose of the first meeting between the sick-listed employee and the RTW coordinator was to identify obstacles for RTW from the perspective of the employee. The second meeting was between the supervisor and the RTW coordinator, where obstacles to the employee's RTW were identified from the perspective of the supervisor. In the third meeting, which was generally the longest, the employee, supervisor and RTW coordinator discussed solutions and formulated a consensus-based plan for their implementation.
- (b) According to the evidence-based guideline of the Dutch Association of Occupational Physicians (NVAB) published in 2000 and updated in 2007. This guideline aims to facilitate the optimal functioning of employees with mental health problems and to prevent long-term sick leave and frequent

Study	van Oostrom 2009				
Study details	Population & interventions	Costs	Health outcomes	Cost-effectiveness	

recurrences. An early start to the treatment by occupational physicians is recommended. Occupational physicians act as motivating counsellors using cognitive behavioural elements to enhance the problem-solving capacity of employees. In addition, the Improved Gatekeeper Act requires that both the employer and employee take responsibility for a RTW plan.

Appendix J – Health economic analysis

See separate economic modelling report.

Appendix K – Excluded studies

Economic studies

Table 6: Summary of studies excluded from the economic evidence review for the workplace health interventions

Reference	Reason for exclusion	RQs
Aas RW, Holte KA, Tuntland H, Roe C, Labriola M, Lund T, et al. Workplace interventions for neck pain in workers. Cochrane Database Syst Rev. 2011; (4): CD008160. Available from: https://www.cochranelibrary.com/cdsr/doi/10.1002/14651858.CD008160/full	Ineligible study design	1a, 2a, 3a
Arends I, Bultmann U, van Rhenen W, H G, van der Klink JJL. Economic evaluation of a problem solving intervention to prevent recurrent sickness absence in workers with common mental disorders. PLoS ONE. 2013;8(8):e71937.	Ineligible outcomes	1a, 2a
Bedell W, Kaszkin-Bettag M. Coherence and health care costRCA actuarial study: a cost-effectiveness cohort study. Altern Ther Health Med. 2010;16(4):26-31.	Ineligible study design	1a, 2a, 3a
Bergstrom G, Bergstrom C, Hagberg J, Bodin L, Jensen I. A 7-year follow-up of multidisciplinary rehabilitation among chronic neck and back pain patients. Is sick leave outcome dependent on psychologically derived patient groups? Eur J Pain. 2010;14(4):426-33.	Ineligible intervention	1a, 2a, 3a
Bernaards CM, Bosmans JE, Hildebrandt VH, van Tulder MW, Heymans MW. The cost-effectiveness of a lifestyle physical activity intervention in addition to a work style intervention on recovery from neck and upper limb symptoms and pain reduction in computer workers. Occup Environ Med. 2011;68(4):265-72.	Ineligible patient population	1a, 2a, 3a
Braun T, Bambra C, Booth M, Adetayo K, Milne E. Better health at work? An evaluation of the effects and cost-benefits of a structured workplace health improvement programme in reducing sickness absence. J Public Health. 2015;37(1):138-42.	Ineligible patient population	1a, 2a, 3a
Brown KC, Sirles AT, Hilyer JC, Thomas MJ. Cost-effectiveness of a back school intervention for municipal employees. Spine. 1992;17(10):1224-8.	Ineligible patient population	1a, 2a, 3a
Brouwers E, de Bruijne M, Terluin B, Tiemens BG, Verhaak PFM. Cost-effectiveness of an activating intervention by social workers for patients with minor mental disorders on sick leave: a randomized controlled trial. Eur J Public Health. 2007;17(2):214-20.	Ineligible outcomes	1a, 2a

Reference	Reason for exclusion	RQs
Bultmann U, Sherson D, Olsen J, Hansen CL, Lund T, Kilsgaard J. Coordinated and tailored work rehabilitation: a randomized controlled trial with economic evaluation undertaken with workers on sick leave due to musculoskeletal disorders. J Occup Rehabil. 2009;19(1):81-93.	Ineligible outcomes	1a, 2a
de Jong PHP, Hazes JM, Buisman LR, Barendregt PJ, van Zeben D, van der Lubbe PA, et al. Best cost-effectiveness and worker productivity with initial triple DMARD therapy compared with methotrexate monotherapy in early rheumatoid arthritis: cost-utility analysis of the tREACH trial. Rheumatol. 2016;55(12):2138-47.	Ineligible patient population	1a, 2a, 3a
Dewa CS, Hoch JS. Estimating the net benefit of a specialized return-to-work program for workers on short-term disability related to a mental disorder: an example exploring investment in collaborative care. J Occup Environ Med. 2014;56(6):628-31.	Ineligible patient population	1a, 2a, 3a
Dewa CS, Hoch JS, Carmen G, Guscott R, Anderson C. Cost, effectiveness, and cost-effectiveness of a collaborative mental health care program for people receiving short-term disability benefits for psychiatric disorders. Can J Psychiatry. 2009;54(6):379-88.	Ineligible patient population	1a, 2a, 3a
Driessen M, Bosmans J, Proper K, Anema J, Bongers P, van der Beek A. The economic evaluation of a participatory ergonomics programme to prevent low back and neck pain. Work. 2012;41(Suppl 1):2315-20.	Ineligible patient population	1a, 2a, 3a
Finnes A, Enebrink P, Sampaio F, Sorjonen K, Dahl J, Ghaderi A, et al. Cost-Effectiveness of Acceptance and Commitment Therapy and a Workplace Intervention for Employees on Sickness Absence due to Mental Disorders. J Occup Environ Med. 2017;59(12):1211-20.	Ineligible outcomes	1a, 2a
Flanagan H, Barwell F, Mazelan P, Spurgeon P. A Better Model of Managing Sickness Absence. In: Spurgeon P, Burke RJ, Cooper CL, editors. The Innovation Imperative in Health Care Organisations. Cheltenham: Edward Elgar Publishing; 2012. p. 113-34.	Ineligible study design	1a, 2a, 3a
Geraedts AS, Van Dongen JM, Kleiboer AM, Wiezer NM, Van Mechelen W, Cuijpers P, et al. Economic evaluation of a web-based guided self-help intervention for employees with depressive symptoms: Results of a randomized controlled trial. J Occup Environ Med. 2015;57(6):666-75.	Ineligible patient population	1a, 2a, 3a
Globe D, Mazonson P, Santas C, Murphy R, Cheng A, Huang X, et al. Impact of etanercept treatment on absenteeism and productivity: The work loss and productivity survey. Am Health Drug Benefits. 2010;3(4):191-200.	Ineligible patient population	1a, 2a, 3a

Reference	Reason for exclusion	RQs
Haldorsen EM, Kronholm K, Skouen JS, Ursin H. Predictors for outcome of a multi-modal cognitive behavioural treatment program for low back pain patients-a 12-month follow-up study. Eur J Pain. 1998;2(4):293-307.	Ineligible study design	1a, 2a, 3a
Hartfiel N, Clarke G, Havenhand J, Phillips C, Edwards RT. Cost-effectiveness of yoga for managing musculoskeletal conditions in the workplace. Occup Med. 2017;67(9):687-95.	Ineligible patient population	1a, 2a, 3a
Hlobil H, Staal JB, Twisk J, Koke A, Ariens G, Smid T, et al. The effects of a graded activity intervention for low back pain in occupational health on sick leave, functional status and pain: 12-month results of a randomized controlled trial. J Occup Rehabil. 2005;15(4):569-80.	Ineligible study design	1a, 2a, 3a
lijima S, Yokoyama K, Kitamura F, Fukuda T, Inaba R. Cost-benefit analysis of comprehensive mental health prevention programs in Japanese workplaces: a pilot study. Ind Health. 2013;51(6):627-33.	Ineligible intervention	1a, 2a, 3a
Jensen IB, Bergstrom G, Ljungquist T, Bodin L. A 3-year follow-up of a multidisciplinary rehabilitation programme for back and neck pain. Pain. 2005;115(3):273-83.	Ineligible outcomes	1a, 2a
Jensen C, Nielsen CV, Jensen OK, Petersen KD. Cost-effectiveness and cost-benefit analyses of a multidisciplinary intervention compared with a brief intervention to facilitate return to work in sick-listed patients with low back pain. Spine. 2013;38(13):1059-67.	Ineligible setting	1a, 2a, 3a
Kim P, Hayden JA, Mior SA. The cost-effectiveness of a back education program for firefighters: a case study. J Can Chiropr Assoc. 2004;48(1):13-9.	Ineligible patient population	1a, 2a, 3a
Lambeek LC, Bosmans JE, Van Royen BJ, Van Tulder MW, Van Mechelen W, Anema JR. Effect of integrated care for sick listed patients with chronic low back pain: economic evaluation alongside a randomised controlled trial. BMJ. 2010;341:c6414.	Ineligible outcomes	1a, 2a
Lammerts L, van Dongen JM, Schaafsma FG, van Mechelen W, Anema JR. A participatory supportive return to work program for workers without an employment contract, sick-listed due to a common mental disorder: an economic evaluation alongside a randomized controlled trial. BMC Public Health. 2017;17(1):162.	Ineligible patient population	1a, 2a, 3a
Loisel P, Abenhaim L, Durand P, Esdaile JM, Suissa S, Gosselin L, et al. A population-based, randomized clinical trial on back pain management. Spine. 1997;22(24):2911-8.	Ineligible study design	1a, 2a, 3a
Loisel P, Lemaire J, Poitras S, Durand MJ, Champagne F, Stock S, et al. Cost-benefit and cost-effectiveness analysis of a disability prevention model for back pain management: a six year follow up study. Occup Environ Med. 2002;59(12):807-15.	Ineligible outcomes	1a, 2a

Reference	Reason for exclusion	RQs
McLaren CF, Reville RT, Seabury SA. How Effective Are Employer Return to Work Programs? International Review of Law and Economics. 2017;52(C):58-73.	Ineligible patient population	1a, 2a, 3a
Meijer EM, Sluiter JK, Heyma A, Sadiraj K, Frings-Dresen MH. Cost-effectiveness of multidisciplinary treatment in sick-listed patients with upper extremity musculoskeletal disorders: a randomized, controlled trial with one-year follow-up. Int Arch Occup Environ Health. 2006;79(8):654-64.	Ineligible outcomes	1a, 2a
Mewes JC, Steuten LMG, Groeneveld IF, de Boer AGEM, Frings-Dresen MHW, Ijzerman MJ, et al. Return-to-work intervention for cancer survivors: budget impact and allocation of costs and returns in the Netherlands and six major EU-countries. BMC Cancer. 2015; 15: 899. Available from: https://bmccancer.biomedcentral.com/articles/10.1186/s12885-015-1912-7	Ineligible patient population	1a, 2a, 3a
Michaels CN, Greene AM. Worksite wellness: increasing adoption of workplace health promotion programs. Health Promot Pract. 2013;14(4):473-9.	Ineligible study design	1a, 2a, 3a
Molde Hagen E, Grasdal A, Eriksen H R. Does early intervention with a light mobilisation program reduce longterm sick leave for low back pain: a 3- year follow-up study. Spine. 2003;28(20):2309-16.	Ineligible setting	1a, 2a, 3a
Moyneur E, Bookhart BK, Mody SH, Fournier A-A, Mallett D, Duh MS. The economic impact of pre-dialysis epoetin alpha on health care and work loss costs in chronic kidney disease: an employer's perspective. Disease Management. 2008;11(1):49-58.	Ineligible patient population	1a, 2a, 3a
Niemisto L, Rissanen P, Sarna S, Lahtinen-Suopanki T, Lindgren KA, Hurri H. Cost-effectiveness of combined manipulation, stabilizing exercises, and physician consultation compared to physician consultation alone for chronic low back pain: a prospective randomized trial with 2-year follow-up. Spine. 2005;30(10):1109-15.	Ineligible patient population	1a, 2a, 3a
Noben C, Evers S, Genabeek Jv, Nijhuis F, de Rijk A. Improving a web-based employability intervention for work-disabled employees: results of a pilot economic evaluation. Disability and rehabilitation. Assistive technology. 2017;12(3):280-89.	Ineligible outcomes	1a, 2a, 3a
Noben C, Hoefsmit N, Evers S, de Rijk A, Houkes I, Nijhuis F. Economic Evaluation of a New Organizational RTW Intervention to Improve Cooperation Between Sick-Listed Employees and Their Supervisors: A Field Study. J Occup Environ Med. 2015;57(11):1170-7.	Ineligible outcomes	1a, 2a, 3a
Noben C, Smit F, Nieuwenhuijsen K, Ketelaar S, Gartner F, Boon B, et al. Comparative cost-effectiveness of two interventions to promote work functioning by targeting mental health complaints among nurses: pragmatic cluster randomised trial. Int J Nurs Stud. 2014;51(10):1321-31.	Ineligible patient population	1a, 2a, 3a

Reference	Reason for exclusion	RQs
Olofsson S, Wickstrom A, Hager Glenngard A, Persson U, Svenningsson A. Effect of treatment with natalizumab on ability to work in people with multiple sclerosis: Productivity gain based on direct measurement of work capacity before and after 1 year of treatment. BioDrugs. 2011;25(5):299-306.	Ineligible study design	1a, 2a, 3a
Oude Hengel KM, Bosmans JE, Van Dongen JM, Bongers PM, Van der Beek AJ, Blatter BM. Prevention program at construction worksites aimed at improving health and work ability is cost-saving to the employer: results from an RCT. Am J Ind Med. 2014;57(1):56-68.	Ineligible patient population	1a, 2a, 3a
Radford K, Phillips J, Drummond A, Sach T, Walker M, Tyerman A, et al. Return to work after traumatic brain injury: cohort comparison and economic evaluation. Brain Inj. 2013;27(5):507-20.	Ineligible outcomes	1a, 2a
Rantonen J, Karppinen J, Vehtari A, Luoto S, Viikari-Juntura E, Hupli M, et al. Cost-effectiveness of providing patients with information on managing mild low-back symptoms in an occupational health setting. BMC Public Health. 2016; 16: 316. Available from: https://bmcpublichealth.biomedcentral.com/articles/10.1186/s12889-016-2974-4	Ineligible patient population	1a, 2a, 3a
Rebergen DS, DJ B, van Tulder MW, van der Beek AJ, van Mechelen W. Cost-effectiveness of guideline-based care for workers with mental health problems. J Occup Environ Med. 2009;51(3):313-22.	Ineligible outcomes	1a, 2a
Rittle C. Can increasing adult vaccination rates reduce lost time and increase productivity? Workplace Health Saf. 2014;62(12):508-16.	Ineligible patient population	1a, 2a, 3a
Roelofs PDDM, Bierma-Zeinstra SMA, van Poppel MNM, van Mechelen W, Koes BW, van Tulder MW. Cost-effectiveness of lumbar supports for home care workers with recurrent low back pain: an economic evaluation alongside a randomized-controlled trial. Spine. 2010;35(26):E1619-26.	Ineligible patient population	1a, 2a, 3a
Schene AH, Koeter MW, Kikkert MJ, Swinkels JA, McCrone P. Adjuvant occupational therapy for work-related major depression works: randomized trial including economic evaluation. Psychol Med. 2007;37(3):351-62.	Ineligible outcomes	1a, 2a
Schneider U, Linder R, Verheyen F. Long-term sick leave and the impact of a graded return-to-work program: evidence from Germany. Eur J Health Econ. 2016;17(5):629-43.	Ineligible study design	1a, 2a, 3a
Shephard RJ. Do work-site exercise and health programs work? Phys Sportsmed. 1999;27(2):48-72. Spekle EM, Heinrich J, Hoozemans MJM, Blatter BM, van der Beek AJ, van Dieen JH, et al. The cost-effectiveness of the RSI QuickScan intervention programme for computer workers: Results of an economic evaluation alongside a randomised controlled trial. BMC Musculoskelet Disord. 2010; 11:	Ineligible study design Ineligible intervention	1a, 2a, 3a 1a, 2a, 3a

Reference	Reason for exclusion	RQs
259. Available from: https://bmcmusculoskeletdisord.biomedcentral.com/articles/10.1186/1471-2474-		
11-259		
Squires H, Rick J, Carroll C, Hillage J. Cost-effectiveness of interventions to return employees to work	Ineligible outcomes	1a, 2a
following long-term sickness absence due to musculoskeletal disorders. J Public Health.		
2012;34(1):115-24.		
Steenstra IA, Anema JR, van Tulder MW, Bongers PM, de Vet HC, van Mechelen W. Economic	Ineligible outcomes	1a, 2a
evaluation of a multi-stage return to work program for workers on sick-leave due to low back pain. J		
Occup Rehabil. 2006;16(4):557-78.		4 0 0
Suoyrjo H, Oksanen T, Hinkka K, Kivimaki M, Klaukka T, Pentti J, et al. The effectiveness of	Ineligible patient population	1a, 2a, 3a
vocationally oriented multidisciplinary intervention on sickness absence and early retirement among		
employees at risk: an observational study. Occup Environ Med. 2009;66(4):235-42. Suryahadi A, Sambodho P. An Assessment of Policies to Improve Teacher Quality and Reduce	Ineligible patient population	10.20.20
Teacher Absenteeism. In: Suryadarma D, Jones GW, editors. Education in Indonesia. Singapore	mengible patient population	1a, 2a, 3a
Institute of Southeast Asian Studies; 2013. p. 139-59.		
Taimela S, Justen S, Aronen P, Sintonen H, Laara E, Malmivaara A, et al. An occupational health	Ineligible patient population	1a, 2a, 3a
intervention programme for workers at high risk for sickness absence. Cost effectiveness analysis	mengiolo patient population	14, 24, 64
based on a randomised controlled trial. Occup Environ Med. 2008;65(4):242-8.		
Targett P, Wehman P. Return to Work after Traumatic Brain Injury: A Supported Employment	Ineligible outcomes	1a, 2a, 3a
Approach. In: Schultz IZ, Rogers S, editors. Work Accommodation and Retention in Mental Health.	G	
New York: Springer; 2011. p. 277-94.		
Theodore BR, Mayer TG, Gatchel RJ. Cost-effectiveness of early versus delayed functional restoration	Ineligible patient population	1a, 2a, 3a
for chronic disabling occupational musculoskeletal disorders. J Occup Rehabil. 2015;25(2):303-15.		
Torstensen TA, Ljunggren AE, Meen HD, Odland E, Mowinckel P, Geijerstam S. Efficiency and costs	Ineligible setting	1a, 2a, 3a
of medical exercise therapy, conventional physiotherapy, and self-exercise in patients with chronic low		
back pain. A pragmatic, randomized, single-blinded, controlled trial with 1-year follow-up. Spine.		
1998;23(23):2616-24.		
Uegaki K, Bakker I, de Bruijne M, van der Beek A, Terluin B, van Marwijk H, et al. Cost-effectiveness	Ineligible outcomes	1a, 2a
of a minimal intervention for stress-related sick leave in general practice: results of an economic		
evaluation alongside a pragmatic randomised control trial. J Affect Disord. 2010;120(1-3):177-87.		

Reference	Reason for exclusion	RQs
Van Der Beek AJ. Primary preventive effects of a multifaceted workplace intervention on low back pain. Pain. 2015;156(9):1583-84.	Ineligible study design	1a, 2a, 3a
van Duijn M, Eijkemans MJ, Koes BW, Koopmanschap MA, Burton KA, Burdorf A. The effects of timing on the cost-effectiveness of interventions for workers on sick leave due to low back pain. Occup Environ Med. 2010;67(11):744-50.	Ineligible intervention	1a, 2a, 3a
van Oostrom SH, Heymans MW, de Vet HCW, van Tulder MW, van Mechelen W, Anema JR. Economic evaluation of a workplace intervention for sick-listed employees with distress. Occup Environ Med. 2010;67(9):603-10.	Ineligible outcomes	1a, 3a
Vogt J, Leonhardt J, Koper B, Pennig S. Economic evaluation of CISMa pilot study. Int J Emerg Ment Health. 2004;6(4):185-96.	Ineligible intervention	1a, 2a, 3a
Welch LS. Improving work ability in construction workers - Let's get to work. Scand J Work Environ Health. 2009;35(5):321-24.	Ineligible study design	1a, 2a, 3a
Wynne-Jones G, Artus M, Bishop A, Lawton SA, Lewis M, Jowett S, et al. Effectiveness and costs of a vocational advice service to improve work outcomes in patients with musculoskeletal pain in primary care: A cluster randomised trial (SWAP trial ISRCTN 52269669). Pain. 2018;159(1):128-38.	Ineligible outcomes	1a, 2a, 3a
Yassi A, Kettner J, Hammond G, Cheang M, McGill M. Effectiveness and cost-benefit of an influenza vaccination program for health care workers. Can J Infect Dis. 1991;2(3):101-8.	Ineligible patient population	1a, 2a, 3a
Yermakov S, Davis M, Calnan M, Fay M, Cox-Buckley B, Sarda S, et al. Impact of increasing adherence to disease-modifying therapies on healthcare resource utilization and direct medical and indirect work loss costs for patients with multiple sclerosis. J Med Econ. 2015;18(9):711-20.	Ineligible patient population	1a, 2a, 3a

Appendix M – Health economic quality assessment

Study identification		
Arends I, Bultmann U, van Rhenen W, Groen H, van der Klink JJL. Econor sickness absence in workers with common mental disorders. PLoS ONE.		solving intervention to prevent recurrent
Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	Unclear	Not clear about the length of time people have been off of work
1.2 Are the interventions appropriate for the review question?	Yes	Intervention targeted at return to work
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in the Netherlands where the occupational support offered is differently organised to the UK - for example there are occupational physicians
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Societal for CEA and employer for CBA
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	Effects included were appropriate for the analysis chosen
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	NA	QALYs were not used, but the effectiveness measures chosen were reasonable for the CEA performed
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Yes	Costs to all relevant sectors were considered
1.9 Overall judgement: Partially applicable		

Section 2: Study limitations (the level of methodological quality)	NA	No decision model was used
2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	INA	no decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was only 12 months so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Partly	Impact on QALYs was not considered
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	The underlying trial was used for resource use data and published sources were used for costs data
2.8 Are the unit costs of resources from the best available source?	Yes	The underlying trial was used for resource use data and published sources were used for costs data
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Yes	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps. Deterministic SA performed on key parameters
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		
CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; QALY: quality-ad	diusted life-vear: RC	CT: randomised controlled trial: SA: sensitivity analysis

Brouwers EPM, de Bruijne MC, Terluin B, Tiemens BG, Verhaak PFM. Cost-effectiveness of an activating intervention by social workers for patients with minor mental disorders on sick leave: a randomized controlled trial. Eur J Public Health. 2007;17(2):214-20.

	Question no: 3.1 & 3.3
Yes/partly/no/unclear/NA	Comments
Partly	Patients were on sick leave for less than 3 months
Yes	Intervention was targeted at return to work
Partly	The study was conducted in the Netherlands where the occupational support offered is differently organised to that in the UK - for example there are occupational physicians
Yes	Societal for CEA, public health insurer for CBA
Yes	Effects included were appropriate for the analysis chosen
NA	Not conducted but a short time horizon was used
Partly	Health states were derived using SF36. The sources of utility values for SF36 health states were not provided.
Partly	Indirect costs were not considered
NA	No decision model was used
No	The time horizon was only 18 months so it is unclear whether return to work was sustained
Yes	QALYs were reported
	Partly Yes Partly Yes Yes NA Partly Partly NA No

2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	Underlying trial for resource use and published sources for costs
2.8 Are the unit costs of resources from the best available source?	Yes	The underlying trial was used for resource use data and published sources were used for costs data
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Partly	Stochastic uncertainty in the data were dealt with using nonparametric bootstraps. No deterministic sensitivity analysis (SA) was performed on key parameters
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		

form 36

Bultmann U, Sherson D, Olsen J, Hansen CL, Lund T, Kilsgaard J. Coordinated and tailored work rehabilitation: a randomized controlled trial with economic evaluation undertaken with workers on sick leave due to musculoskeletal disorders. J Occup Rehabil. 2009;19(1):81-93.

CBA: cost-benefit analysis; CUA: cost-utility analysis; QALY: quality-adjusted life-year; RCT: randomised controlled trial; SA: sensitivity analysis; SF36: short-

Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	Yes	Workers were on sick leave for 4-12 weeks
1.2 Are the interventions appropriate for the review question?	Yes	Intervention was targeted at return to work
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in Denmark where the organisation of sickness benefits system is

		similar to the UK, but may still be different enough to limit the generalisability of findings
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Societal
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	NA	QALYs were not used but the effectiveness measures chosen were reasonable
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Yes	Costs to all relevant sectors were considered
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)		
2.1 Does the model structure adequately reflect the nature of the topic under	NA	No decision model was used
evaluation?	TV C	The decision model was ased
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was only 12 months so it's unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Partly	Impact on QALYs was not considered
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	Resource use and cost data were collected from national Danish registries
2.8 Are the unit costs of resources from the best available source?	Yes	Resource and cost data collected from national Danish registries for trial participants

2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Partly	One way deterministic sensitivity analysis. No PSA was undertaken.
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		
PSA: probabilistic sensitivity analysis; QALY: quality-adjusted life-year; RCT: randomised controlled trial		

Study identification Finnes A, Enebrink P, Sampaio F, Sorjonen K, Dahl J, Ghaderi A, et al. Cost-Effectiveness of Acceptance and Commitment Therapy and a Workplace Intervention for Employees on Sickness Absence due to Mental Disorders. J Occup Environ Med. 2017;59(12):1211-20.			
Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3	
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments	
1.1 Is the study population appropriate for the review question?	Unclear	Workers on sickness absence due to anxiety	
1.2 Are the interventions appropriate for the review question?	Yes	Intervention targeted at return to work	
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in Sweden where the organisation of sickness benefits system is similar to the UK but may still be different enough to limit the generalisability of findings	
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Healthcare and separate societal analysis including sickness benefit costs	
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen	
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted but a short time horizon was used	

1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	Yes	Health states were derived using EQ-5D utility data valued using the English valuation set (Swedish in scenario analysis)
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	No	The societal perspective did not include impacts on employers and drug costs were not included in either perspective.
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)		
2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	NA	No decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was 12 months so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Yes	Health states were derived using EQ-5D data valued using the UK valuation set (Swedish in scenario analysis)
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Partly	Medication costs were not included
2.7 Are the estimates of resource use from the best available source?	Yes	The volumes of each cost category were obtained from study records, and unit costs were obtained from national public databases and websites
2.8 Are the unit costs of resources from the best available source?	Yes	The volumes of each cost category were obtained from study records, and unit costs were obtained from national public databases and websites
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given

2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Yes	Stochastic uncertainty in the data were dealt with using nonparametric bootstrapping. Scenario analysis for key model assumptions
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		

Jensen IB, Bergstrom G, Ljungquist T, Bodin L. A 3-year follow-up of a multidisciplinary rehabilitation programme for back and neck pain. Pain. 2005:115(3):273-83

EQ5D:euroqol 5 dimensions; DALY: disability-adjusted life-year; QALY: quality-adjusted life-year; RCT: randomised controlled trial

2005;115(3):273-83.		
Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	Yes	On sick leave for 1 to 6 months
1.2 Are the interventions appropriate for the review question?	Yes	Intervention targeted at return to work
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in Sweden where the organisation of sickness benefits system is similar to the UK, but may still be different enough to limit the generalisability of findings
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Societal
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted, but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	NA	SF-36 data were collected, but not reported beyond a statement that there was no difference between groups

1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued? 1.9 Overall judgement: Partially applicable Section 2: Study limitations (the level of methodological quality) 2.1 Does the model structure adequately reflect the nature of the topic under NA Appropriate costs and outcomes were considered, for the perspective taken NA No decision model was used
Section 2: Study limitations (the level of methodological quality)
2.1 Does the model structure adequately reflect the nature of the topic under NA No decision model was used
evaluation?
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes? Partly The time horizon was 3 years so some estimate of long-term costs and effectiveness could be made
2.3 Are all important and relevant outcomes included? Partly SF-36 data were collected, but not reported beyond a statement that there was no difference between groups
2.4 Are the estimates of baseline outcomes from the best available source? Yes Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source? Yes Taken from a RCT
2.6 Are all important and relevant costs included? Yes All relevant costs were considered
2.7 Are the estimates of resource use from the best available source? Yes Resource use from a trial
2.8 Are the unit costs of resources from the best available source? Partly Costs were from published sources
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data? NA Not relevant (CBA)
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis? No Not undertaken
2.11 Is there any potential conflict of interest? No There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations
Other comments: None
CBA: cost-benefit analysis; PSA: probabilistic sensitivity analysis; QALY: quality-adjusted life-year; RCT: randomised controlled trial; SA: sensitivity analysis; SF36: short form 36

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Lambeek LC, Bosmans JE, Van Royen BJ, Van Tulder MW, Van Mechelen W, Anema JR. Effect of integrated care for sick listed patients with chronic low back pain: economic evaluation alongside a randomised controlled trial. BMJ (Clinical research ed.). 2010;341:c6414.

Yes/partly/no/unclear/NA	Comments
Partly	Includes people on partial sick leave so it may not be continuous leave
Yes	Intervention was targeted at return to work
Partly	The study was conducted in the Netherlands where the occupational support offered is differently organised to the UK, for example there are occupational physicians
Yes	Societal
Yes	The effects included were appropriate for analysis chosen
NA	Not conducted, but a short time horizon was used
Yes	Health states were derived using EQ-5D data valued using the Dutch tariff
Partly	Cost of work modifications were not included
NA	No decision model was used
	Yes Partly Yes Yes Yes NA Yes Partly

2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was 12 months so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Yes	QALYs were reported
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	From a retrospective questionnaire and Dutch manual for costing economic evaluations
2.8 Are the unit costs of resources from the best available source?	Yes	From a retrospective questionnaire and Dutch manual for costing economic evaluations
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Yes	Stochastic uncertainty in the data were dealt with using nonparametric bootstrapping. Scenario analysis was undertaken for key model assumptions.
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		
EQ5D: euroqol 5 dimensions; QALY: quality-adjusted life-year; RCT: randomis	ed controlled trial	

Study identification Loisel P, Lemaire J, Poitras S, Durand MJ, Champagne F, Stock S, et al. Comodel for back pain management: a six year follow up study. Occup Envir		
Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	Yes	Workers were absent for 4 or more weeks

4.0. And the interpreting among night for the review was the no	Yes	Intervention was targeted at return to work
1.2 Are the interventions appropriate for the review question?1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in Canada where the organisation of sickness benefits system is similar to the UK, but may still be different enough to limit the generalisability of findings
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Partly	Health and employment insurer
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	NA	QALYs were not used, but the effectiveness measures chosen were reasonable
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Partly	Costs of job modifications were not recorded.
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)		
2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	NA	No decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	Yes	The time horizon of 6.4 years showed significant changes in costs and outcomes over time
2.3 Are all important and relevant outcomes included?	Partly	The impact on QALYs was not considered
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	Quebec Workers Compensation Database

2.8 Are the unit costs of resources from the best available source?	Yes	Quebec Workers Compensation Database
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Partly	Deterministic SA was performed. No probabilistic sensitivity analysis (PSA) was performed.
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Minor limitations		
Other comments: None		

PSA: probabilistic sensitivity analysis; QALY: quality-adjusted life-year; RCT: randomised controlled trial; SA: sensitivity analysis

Study identification Meijer EM, Sluiter JK, Heyma A, Sadiraj K, Frings-Dresen MH. Cost-effectiveness of multidisciplinary treatment in sick-listed patients with upper extremity musculoskeletal disorders: a randomized, controlled trial with one-year follow-up. Int Arch Occup Environ Health. 2006;79(8):654-64.			
Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3	
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments	
1.1 Is the study population appropriate for the review question?	Partly	At least 50% sick leave over past 4 to 20 weeks	
1.2 Are the interventions appropriate for the review question?	Yes	Intervention was targeted at return to work	
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in the Netherlands where the occupational support offered is differently organised to the UK, for example there are occupational physicians	
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Societal	
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen	
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted but a short time horizon was used	

1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	NA	QALYs were not used but the effectiveness measures chosen were reasonable
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Yes	Appropriate costs and outcomes for the perspective were considered
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)		
2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	NA	No decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was 12 months so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Partly	The impact on QALYs was not considered
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	Dutch board on medical tariffs, Dutch medicines compensation system, self-reported costs in questionnaire
2.8 Are the unit costs of resources from the best available source?	Yes	Dutch board on medical tariffs, Dutch medicines compensation system, self-reported costs in questionnaire
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	No	Not undertaken
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		

PSA: probabilistic sensitivity analysis; QALY: quality-adjusted life-year; RCT: randomised controlled trial

Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	No	Includes unpaid workers and students. It is not clear how long injury had caused the workers to be out of work nor is it clear if they are able to return to work.
1.2 Are the interventions appropriate for the review question?	Yes	Intervention targeted at return to work
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Yes	UK study within past 6 years
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Health and social care for CBA, societal for CEA and CUA
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	No	EQ-5D VAS was used
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Yes	Appropriate costs and outcomes for the perspective were considered
1.9 Overall judgement: Not applicable		

2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	NA	No decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was 12 months so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Partly	Impact on QALYs was reported using VAS
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Partly	The estimates were taken from a clinical trial but it was only an observational study
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	Resource use from a trial. NHS reference costs, PSSRU and Jobcentre plus.
2.8 Are the unit costs of resources from the best available source?	Yes	NHS reference costs, PSSRU and Jobcentre plus
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Yes	Stochastic uncertainty in the data were dealt with using nonparametric bootstrapping. Scenario analysis was used for key model assumptions
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		

Other comments:

CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; cost-utility analysis; EQ5D: euroqol 5 dimensions; NHS: National Health Service; PSSRU: Personal Social Services Research Unit; QALY: quality-adjusted life-year; RCT: randomised controlled trial; VAS - visual analogue scale

Study identification

Rebergen DS, Bruinvels DJ, van Tulder MW, van der Beek AJ, van Mechelen W. Cost-effectiveness of guideline-based care for workers with mental health problems. J Occup Environ Med. 2009;51(3):313-22.

Guidance topic: Indoor Air Quality At Home Question no: 3.1 & 3.3

Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	Unclear	The length of time people have been off work is unclear.
1.2 Are the interventions appropriate for the review question?	Yes	Intervention was targeted at return to work
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was conducted in the Netherlands where the occupational support offered is differently organised to the UK, for example there are occupational physicians
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Societal for CEA, employer for CBA
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted, but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	NA	QALYs were not used, but the effectiveness measures chosen were reasonable
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Partly	Patient and family health care costs outside of occupational healthcare costs were not included
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)	NIA	No. 1. Colombia de la constanta de la colombia del colombia del colombia de la colombia del colombia del colombia de la colombia de la colombia de la colombia de la colombia del colombia
2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	NA	No decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was 12 months, so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Partly	Impact on QALYs was not considered
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants

2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Partly	Patient and family health care costs outside of occupational healthcare costs were not included
2.7 Are the estimates of resource use from the best available source?	Yes	Insurance company records of Dutch police force on trial participants
2.8 Are the unit costs of resources from the best available source?	Yes	Insurance company records of Dutch police force on trial participants
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Yes	Stochastic uncertainty in the data were dealt with using nonparametric bootstrapping. Scenario analysis was used for key model assumptions
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		

Schene AH, Koeter MW, Kikkert MJ, Swinkels JA, McCrone P. Adjuvant occupational therapy for work-related major depression works: randomized trial including economic evaluation. Psychol Med. 2007;37(3):351-62.

CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; QALY: quality-adjusted life-year; RCT: randomised controlled trial

Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	Partly	Workers with at least 50% absence over last 10 weeks to 2 years
1.2 Are the interventions appropriate for the review question?	Yes	Intervention was targeted at return to work

1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in the Netherlands where the occupational support offered is differently organised to the UK, for example there are occupational physicians
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Societal
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	No	No health or employment outcomes were reported beyond earnings over a 12-month period
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted, but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	Partly	QALYs were not used, but the effectiveness measures chosen were reasonable
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Yes	Appropriate costs and outcomes for the perspective were considered
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)		
2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	NA	No decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was only 12 months so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Partly	Impact on QALYs was not considered
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	From trial
2.8 Are the unit costs of resources from the best available source?	No	Not reported

2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	NA	Was a CBA
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Yes	Stochastic uncertainty in the data were dealt with using nonparametric bootstrapping. Scenario analysis was used for key model assumptions.
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		
CBA: cost-benefit analysis; QALY: quality-adjusted life-year; RCT: randomised	controlled trial	

Squires H, Rick J, Carroll C, Hillage J. Cost-effectiveness of interventions to return employees to work following long-term sickness absence due to musculoskeletal disorders. Journal of public health (Oxford, England). 2012;34(1):115-24.

Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	Partly	Sick leave was 1 week to 6 months
1.2 Are the interventions appropriate for the review question?	Yes	Intervention was targeted at return to work
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	Although this is a UK study, it is based upon effectiveness studies that were conducted outside of the UK and the authors stated that this may limit the generalisability of the results
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	NHS and PSS, societal and employer.
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Partly	Whilst QALYs and increased likelihood of return to work are included, they are not well-reported
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted, but a short time horizon was used

1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	Partly	SF-36 data from British Household Panel Survey
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Yes	Appropriate costs and outcomes for the perspective were considered
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)		
2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	Yes	The Markov model that was developed was adequate to answer the current topic
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	Partly	Although a lifetime time horizon was used, the model was populated with effectiveness evidence over 12 months
2.3 Are all important and relevant outcomes included?	Yes	QALYs were reported
2.4 Are the estimates of baseline outcomes from the best available source?	NA	Hypothetical cohort in the model
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from RCTs
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	DWP, PSSRU and published sources
2.8 Are the unit costs of resources from the best available source?	Yes	DWP, PSSRU and published sources
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Partly	The cost-effectiveness plane was given, but full incremental results were not provided and could not be calculated
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Partly	Deterministic SA was performed. No PSA was undertaken.
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		

DWP: Department of Work and Pensions; NHS: National Health Service; PSA: probabilistic sensitivity analysis; PSS: person shaped support; PSSRU: Personal Social Services Research Unit; QALY: quality-adjusted life-year; RCT: randomised controlled trial; SA: sensitivity analysis; SF36: short form 36

Study identification

Steenstra IA, Anema JR, van Tulder MW, Bongers PM, de Vet HC, van Mechelen W. Economic evaluation of a multi-stage return to work program for workers on sick-leave due to low back pain. J Occup Rehabil. 2006;16(4):557-78.

Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	Partly	Sick leave was 2 to 6 weeks
1.2 Are the interventions appropriate for the review question?	Yes	The intervention was targeted at return to work
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in the Netherlands where the occupational support offered is differently organised to the UK, for example there are occupational physicians
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Societal
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted, but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	No	EQ-5D VAS data were used
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Yes	A full range of costs and outcomes were considered for the perspective chosen
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)		

2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	NA	No decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was 12 months so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Partly	Impact on QALYs was reported using VAS
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	Dutch costing guidelines, health care charges and professional organisations for trial participants
2.8 Are the unit costs of resources from the best available source?	Yes	Dutch costing guidelines, health care charges and professional organisations for trial participants
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Yes	Stochastic uncertainty in the data were dealt with using nonparametric bootstrapping. Scenario analysis was used for key model assumptions.
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest
2.12 Overall assessment: Potentially serious limitations		
Other comments: None		
EQ5D: euroqol 5 dimensions; QALY: quality-adjusted life-year; RCT: randomised controlled trial; VAS: visual analogue scale		

Uegaki K, Bakker I, de Bruijne M, van der Beek A, Terluin B, van Marwijk H, et al. Cost-effectiveness of a minimal intervention for stress-related sick leave in general practice: results of an economic evaluation alongside a pragmatic randomised control trial. J Affect Disord. 2010;120(1-3):177-87.

Guidance topic: Indoor Air Quality At Home Question no: 3.1 & 3.3

Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments
1.1 Is the study population appropriate for the review question?	Partly	Partial sick leave was over 3 months
1.2 Are the interventions appropriate for the review question?	Yes	The intervention was targeted at return to work
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in the Netherlands where the occupational support offered is differently organised to the UK, for example there are occupational physicians
1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Societal
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted, but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	Yes	EQ-5D data using the Dutch tariff
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Yes	A full range of costs and outcomes were considered for the perspective chosen
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)		
2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	NA	No decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was 12 months so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Yes	QALYs were reported
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT

2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered	
2.7 Are the estimates of resource use from the best available source?	Yes	Dutch costing guidelines, health care charges and professional organisations for trial participants	
2.8 Are the unit costs of resources from the best available source?	Yes	Dutch costing guidelines, health care charges and professional organisations	
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given	
2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Yes	Stochastic uncertainty in the data were dealt with using nonparametric bootstrapping. Scenario analysis was used for key model assumptions	
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest	
2.12 Overall assessment: Potentially serious limitations			
Other comments: None			
EQ5D: euroqol 5 dimensions; QALY: quality-adjusted life-year; RCT: randomised controlled trial			

Study identification van Oostrom SH, Heymans MW, de Vet HCW, van Tulder MW, van Mechelen W, Anema JR. Economic evaluation of a workplace intervention for sick-listed employees with distress. Occup Environ Med. 2010;67(9):603-10.				
Guidance topic: Indoor Air Quality At Home		Question no: 3.1 & 3.3		
Section 1: Applicability (relevance to specific review questions and the NICE reference case)	Yes/partly/no/unclear/NA	Comments		
1.1 Is the study population appropriate for the review question?	Partly	Workers had sick leave for 2 to 8 weeks		
1.2 Are the interventions appropriate for the review question?	Yes	The intervention targeted return to work		
1.3 Is the system in which the study was conducted sufficiently similar to the current UK context?	Partly	The study was in the Netherlands where the occupational support offered is differently organised to the UK, for example there are occupational physicians		

1.4 Are the perspectives clearly stated and are they appropriate for the review question?	Yes	Societal for CEA and CUA, employer perspective for CBA
1.5 Are all direct effects on individuals included, and are all other effects included where they are material?	Yes	The effects included were appropriate for the analysis chosen
1.6 Are all future costs and outcomes discounted appropriately?	NA	Not conducted, but a short time horizon was used
1.7 Is QALY used as an outcome, and was it derived using NICE's preferred methods? If not, describe rationale and outcomes used in line with analytical perspectives taken (item 1.4 above).	Yes	EQ-5D data using the Dutch tariff
1.8 Are costs and outcomes from other sectors fully and appropriately measured and valued?	Partly	The costs of workplace adaptations were not considered
1.9 Overall judgement: Partially applicable		
Section 2: Study limitations (the level of methodological quality)		
2.1 Does the model structure adequately reflect the nature of the topic under evaluation?	NA	No decision model was used
2.2 Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?	No	The time horizon was 12 months so it is unclear whether return to work was sustained
2.3 Are all important and relevant outcomes included?	Yes	QALYs were reported
2.4 Are the estimates of baseline outcomes from the best available source?	Yes	Drawn from trial participants
2.5 Are the estimates of relative intervention effects from the best available source?	Yes	Taken from a RCT
2.6 Are all important and relevant costs included?	Yes	All relevant costs were considered
2.7 Are the estimates of resource use from the best available source?	Yes	Dutch costing guidelines, health care charges and professional organisations for trial participants
2.8 Are the unit costs of resources from the best available source?	Yes	Dutch costing guidelines, health care charges and professional organisations
2.9 Is an appropriate incremental analysis presented or can it be calculated from the data?	Yes	Incremental results were given

2.10 Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?	Yes	Stochastic uncertainty in the data were dealt with using nonparametric bootstrapping. Scenario analysis was undertaken for key model assumptions	
2.11 Is there any potential conflict of interest?	No	There were no notable conflicts of interest	
2.12 Overall assessment: Potentially serious limitations			
Other comments: None			
CBA: cost-benefit analysis; CEA: cost-effectiveness analysis; CUA: cost-utility analysis; EQ5D: euroqol 5 dimensions; QALY: quality-adjusted life-year; RCT: randomised controlled trial; VAS: visual analogue scale			