

# Second round screening form

\* Required

1. Author \*

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2. Year

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3. Assessor \*

*Mark only one oval.*

☐ SE

☐ MK

☐ SK

☐ HW

☐ UK

## Eligibility Criteria

### DESIGN

4. the study is a randomized controlled trial or a longitudinal observational study

*Mark only one oval.*

☐ Yes

☐ No

## 5. explanation

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## PATIENTS

6. the study solely includes adults (i.e. 18 years or older) OR 95% of the patients (2x SD) is 18 years or older

*Mark only one oval.*

☐ Yes

☐ No

7. all patients have 'chronic primary pain that is perceived in bone(s), joint(s), muscle(s) or related soft tissue(s)

*Mark only one oval.*

☐ Yes

☐ No

8. at least 75% of the patients have chronic 'chronic primary pain that is perceived in bone(s), joint(s), muscle(s) or related soft tissue(s)

*Mark only one oval.*

☐ Yes

☐ No

## 9. explanation

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## INTERVENTION

## 10. the intervention includes

Additional information: 1) the treatment program includes a common philosophy of rehabilitation in line with the BPS model of pain; 2) includes active patient involvement (tasks, training and/or exercise); 3) an interdisciplinary team of at least 3 different professions; 4) interprofessional communication is embedded in the protocol; 5) All care is provided at the same facility. \*\*\*Note 1\*\*\* both inpatient and outpatient are eligible; we will include interventions of any intensity. \*\*\*Note 2\*\*\* We will not exclude interventions on criterium 4, because most programs do not explicitly report on this. Instead, we assume that HCPs within a single facility will have some form of structured communication.

*Mark only one oval per row.*

	yes	no	not reported
common BPS philosophy of rehabilitation	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
active patient involvement	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
team of 3 different HCPs	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
embedded interprof. communication	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
single facility	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

## 11. explanation

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OUTCOMES

12. the study includes at minimum one of the following outcome measures as primary outcome

Note: pain intensity is a secondary outcome. HRQOL: overall measure of how individuals feel and function in daily life, by assessing multiple domains of health; PF/PI: (self reported capabilities or pain related interference on activities or daily living); Social health: measure the ability to participate in social roles and activities.

Mark only one oval per row.

	yes	no	not reported
hrqol	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
physical functioning or pain interference	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
anxiety, anger, self-efficacy or depression	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
social health	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

13. explanation

TIMEFRAME

## 14. The study includes the following measurement timepoints

Note: We decided not to exclude treatment programs that do not have a post-intervention measurement. Follow-up should

*Mark only one oval per row.*

	Yes	No
baseline	<input type="radio"/>	<input type="radio"/>
post intervention	<input type="radio"/>	<input type="radio"/>
follow-up (at minimum 12 months)	<input type="radio"/>	<input type="radio"/>

## 15. explanation

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## EXCLUSION CRITERIA

## 16. the study focused on patients with post-surgical pain or cancer pain

*Mark only one oval.*

☐ Yes

☐ No

## 17. the study focused on patients of a specific comorbidity (e.g. psychiatric or obese patients)

*Mark only one oval.*

☐ Yes

☐ No

18. explanation

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## FINAL DECISION

19. decision \*

*Mark only one oval.*

☐ include

☐ exclude

20. general comments

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