Content and concurrent validity of the Patient Specific Functional Scale (PSFS) on patients with neck pain.

Gepubliceerd: 29-08-2018 Laatst bijgewerkt: 18-08-2022

We expect to make recommendations in choosing between the PSFS or PSFS-D answering option and to describe important items from a patient's perspective regarding relevance, comprehensiveness and comprehensibility. We expect high correlation...

Ethische beoordeling	Positief advies
Status	Werving gestart
Type aandoening	-
Onderzoekstype	Observationeel onderzoek, zonder invasieve metingen

Samenvatting

ID

NL-OMON22924

Bron
Nationaal Trial Register

Verkorte titel
CROMM study

Aandoening

neck pain, validity, activities

Ondersteuning

Primaire sponsor: Fysio-Experts
Rijndijk 137
2394AG Hazerswoude Rijndijk
070-7400840
info@fysio-experts.nl
Overige ondersteuning: initiator

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

The primary outcome of this study is to evaluate the content and concurrent validity of the patient specific functional scale on patients with neck pain.

Toelichting onderzoek

Doel van het onderzoek

We expect to make recommendations in choosing between the PSFS or PSFS-D answering option and to describe important items from a patient's perspective regarding relevance, comprehensiveness and comprehensibility.

We expect high correlation coefficients ($r \geq 0.60$) between the PSFS(-D) and the NDI.

Onderzoeksopzet

At baseline and 12 week follow-up the NRS, PSFS-D, and NDI will be measured.

Onderzoeksproduct en/of interventie

Before entering the study, participants will be screened to make sure that they meet all of the inclusion criteria. Informed consent is obtained from all participants. The first ten patient entering the study will be asked to join the content validity study as well.

Individual in person interviews following a semi structured discussion will be performed, on the following topics: understanding, meaning, relevance, response options, etc. and the preference regarding the answering option and an example list.

Interviews will be audio-recorded and transcribed for analysis.
After the content validity study is finalized all participants entering the study will receive a questionnaire including the NRS, PSFS(-D) and NDI

Contactpersonen

Publiek

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Wetenschappelijk

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

Patients with neck pain are eligible if they are over 18 years of age, adequately understand Dutch and are classified as Grade I or II as described by the Neck Pain Task Force

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

Patients are excluded in the presence of serious pathology (such as infection, cancer, fracture or rheumatoid arthritis) and previous surgery.

Onderzoeksopzet

Opzet

Type:	Observationeel onderzoek, zonder invasieve metingen
Onderzoeksmodel:	Anders

Controle: N.v.t. / onbekend

Deelname

Nederland

Status:	Werving gestart
(Verwachte) startdatum:	01-07-2018
Aantal proefpersonen:	100
Type:	Verwachte startdatum

Ethische beoordeling

Positief advies

Datum:	29-08-2018
Soort:	Eerste indiening

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

Geen registraties gevonden.

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register	ID
NTR-new	NL7265
NTR-old	NTR7463
Ander register	Erasmus Medical Center, ethics committee : MEC-2018-129

Resultaten