

Corticosteroid injections in patients with trochanteric pain syndrome: open label randomized clinical trial in general practice.

Gepubliceerd: 27-03-2006 Laatste bijgewerkt: 13-12-2022

Local corticosteroid injections combined with usual care will have a positive effect on experienced recovery and will reduce pain at 3 and 12 months follow up compared to usual care alone.

Ethische beoordeling	Positief advies
Status	Werving gestopt
Type aandoening	-
Onderzoekstype	Interventie onderzoek

Samenvatting

ID

NL-OMON20947

Bron

NTR

Verkorte titel

TIS = Trochanter Injection Study

Aandoening

Trochanteric pain, also known as trochanteric bursitis or pseudotrochanteric bursitis

Ondersteuning

Primaire sponsor: Erasmus Medical Center, Department of General Practice

Overige ondersteuning: ZON-MW, The Netherlands Organization for Health Research and Development

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

Experienced recovery (7 points Likert scale) and severity of pain (0-10 VAS) at 3 months follow up.

Toelichting onderzoek

Achtergrond van het onderzoek

A remarkable part of patients with hip pain suffer from trochanteric pain syndrome. In this study (cost) effectiveness of local corticosteroid injections will be determined regarding to normal policy in patients with local trochanteric pain syndrome in general practice.

Doel van het onderzoek

Local corticosteroid injections combined with usual care will have a positive effect on experienced recovery and will reduce pain at 3 and 12 months follow up compared to usual care alone.

Onderzoeksopzet

N/A

Onderzoeksproduct en/of interventie

1. Intervention group: Usual care combined with corticosteroid injections (triamcetonacetaat 40 mg, lidocaine 1%);

2. Control group: Usual care.

Contactpersonen

Publiek

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

Patients aged 18 till 80 consulting the general practitioner with trochanteric hip pain existing longer then one week, with the following signs:

1. Severe local pain by pressure at the trochanter major, but unsure whether the patient recognizes this as the pain he/she consulted for;
2. Local pain by pressure at the trochanter major, which the patient recognizes as the pain he/she consulted for.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

1. Patients who don't understand the questionnaires because of inadequate mastering of Dutch language or cognitive disorders;
2. Patients who presented themselves in the previous year to the GP with the same complaints;
3. Patients with sensibility disorders due to meralgia paresthetica, patients who previously have undergone hip surgery at the same side and patients with systematic rheumatologic or neurologic disorders.

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Toewijzing:	Gerandomiseerd
Blinding:	Open / niet geblindeerd
Controle:	Geneesmiddel

Deelname

Nederland	
Status:	Werving gestopt
(Verwachte) startdatum:	01-05-2006
Aantal proefpersonen:	150
Type:	Werkelijke startdatum

Ethische beoordeling

Positief advies	
Datum:	27-03-2006
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

NTR-new

NTR-old

Ander register

ISRCTN

ID

NL584

NTR640

: N/A

ISRCTN16994576

Resultaten

Samenvatting resultaten

N/A