

Surgical treatment of joint degeneration of the thumb, a cohort of all treated patients.

No registrations found.

| | |
|------------------------------|------------------|
| Ethical review | Positive opinion |
| Status | Recruiting |
| Health condition type | - |
| Study type | - |

Summary

ID

NL-OMON29592

Source

NTR

Brief title

Primary CMC1 osteoarthritis

Health condition

carpometacarpal osteoarthritis thumb (carpometacarpale artrose duimbasis)

Sponsors and support

Primary sponsor: Erasmus MC Rotterdam, the Netherlands

Xpert Clinic, Hilversum, the Netherlands

Source(s) of monetary or material Support: Fonds NutsOhra

Intervention

Outcome measures

Primary outcome

1. Pain (VAS), pain during activities, rest and overall pain in the last week

2. Function (MHQ)

Secondary outcome

3. Satisfaction with the hand (VAS)

4. Return to work

5. Strength (kg), grip/pinch

6. Complications

Study description

Background summary

In this prospective cohort, we include all surgically treated patients for CMC OA for data-analysis, efficacy analysis or comparison with other therapies such as conservative therapy or different surgical procedures

Study objective

- surgical treatment for CMC OA is effective in terms of pain, function and satisfaction
- There is a relatively large group of patients with significant residual pain and functional impairment or deterioration of their complaints
- Baseline factors like MCP-hyperextension, sex, age and pre-operative pain levels can predict succes or failure after surgery
- een significant deel van de patienten die conservatief zijn behandeld worden alsnog chirurgisch behandeld (%?)

Study design

Preoperatively and at 3 and 12 months postoperative.

Intervention

Trapeziectomy + LRTI

Contacts

Public

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Eligibility criteria

Inclusion criteria

1. Patients > 18 years
2. Patient is diagnosed with primary CMC I osteoarthritis and planned for operation;
3. Right and left handed;
4. One or both hands are involved;
5. Surgical treatment with trapeziectomy and LRTI
6. Eaton and Glickel stage II-IV

Exclusion criteria

1. previous surgical treatment
2. posttraumatic osteoarthritis

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3. surgery, primarily for STT osteoarthritis

Study design

Design

| | |
|---------------------|-------------------------|
| Intervention model: | Other |
| Allocation: | Non controlled trial |
| Masking: | Open (masking not used) |
| Control: | N/A , unknown |

Recruitment

| | |
|---------------------------|-------------|
| NL | |
| Recruitment status: | Recruiting |
| Start date (anticipated): | 01-11-2011 |
| Enrollment: | 1000 |
| Type: | Anticipated |

Ethics review

| | |
|-------------------|------------------|
| Positive opinion | |
| Date: | 01-08-2016 |
| Application type: | First submission |

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

| Register | ID |
|----------|--|
| NTR-new | NL5859 |
| NTR-old | NTR6039 |
| Other | Fonds NutsOhra project 1402-056 : MEC-2015-691 |

Study results