Posttraumatic osteoarthritis following distal radius fractures in young patients and the correlation with subjective and objective outcome measures

Published: 23-07-2013 Last updated: 15-05-2024

To determine the incidence of posttraumatic osteoarthritis following distal radius fractures in a cohort of young non-osteoporotic patients and correlation with objective and subjective outcome measures.

Ethical review Approved WMO

Status Recruitment stopped

Health condition type Fractures

Study type Observational invasive

Summary

ID

NL-OMON39658

Source

ToetsingOnline

Brief title

Osteoarthritis following distal radius fractures in young patients

Condition

- Fractures
- Bone and joint therapeutic procedures

Synonym

distal radius fracture, wrist fracture

Research involving

Human

Sponsors and support

Primary sponsor: Universitair Medisch Centrum Groningen

Source(s) of monetary or material Support: Ministerie van OC&W,Fonds de Gavere

Intervention

Keyword: Fracture, Osteoarthritis, Radius, Subjective

Outcome measures

Primary outcome

The incidence of posttraumatic osteoarthritis.

Secondary outcome

Subjective outcome measures objectivied with the use of validated questionnaires (PRWE, DASH, SF-36 and MHQ). Objective outcomes measured with functional tests such as range of motion and gripstrength. Radiological findings measured with X-rays.

Study description

Background summary

The development of posttraumatic osteoarthritis (PA) following distal radius fractures (DRFs) has been commonly described. Direct and indirect joint impact loading, soft tissue injuries, joint dislocation and intra-articular fractures, increase the risk of progressive joint degeneration that causes PA. It is thought posttraumatic osteoarthritis develops less in younger patients. However, it might be more invalidating for a young non-osteoporotic patient to develop posttraumatic osteoarthritis and loss of function following a distal radius fracture than for an older patient. The extend of the loss of function can be objectived using functional measures, such as range of motion and grip strength. Subjective measures to objectivy loss of function as experienced by the patient can be performed using validated questionnaires. In this study, the incidence of posttraumatic osteoarthritis following a distal radius fracture is in young patients is determined. Also, the question arises what the correlation between the existence of post traumatic osteoarthritis and the objective and subjective outcome measures is following a distal radius fracture in young

patients.

Study objective

To determine the incidence of posttraumatic osteoarthritis following distal radius fractures in a cohort of young non-osteoporotic patients and correlation with objective and subjective outcome measures.

Study design

Descriptive cohort study

Study burden and risks

According to the international commission on radiological protection (ICRP) the estimated risk for the radiological procedure (4 x-rays per subject) is categorized aslla, very low level of risk of radiogenic cancer, >0.2 and <2 mSv. In comparison to the * Rijks Instituut Voor Milieuhygiëne (RIVM)*, the annual radiation dose in the Netherlands is 1.7 mSv. The number of visits is once for the functional physical tests combined with the radiological procedure. The physical discomfort during the functional tests and the radiological investigation is minimal. The subject will be asked to sit still for the radiological investigations for 4 times less than 1 minute.

Contacts

Public

Universitair Medisch Centrum Groningen

Hanzeplein 1 Groningen 9713GZ NL

Scientific

Universitair Medisch Centrum Groningen

Hanzeplein 1 Groningen 9713GZ NL

Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years) Elderly (65 years and older)

Inclusion criteria

All patients treated in the period 2005 untill 2011 in the Medical Center Leeuwarden with a distal radius fracture of which the x-ray after the trauma and after 6 weeks is present in the archives of the Radiology Department. Men aged between the age 18-50 years and women between the age of 18-40 years at the time of presentation.

Exclusion criteria

Fractures treated more than 7 days following initial trauma, open fractures and preexistent osteoartritis.

Study design

Design

Study type: Observational invasive

Masking: Open (masking not used)

Control: Uncontrolled

Primary purpose: Diagnostic

Recruitment

NL

Recruitment status: Recruitment stopped

Start date (anticipated): 01-09-2013

Enrollment: 80

Type:	Actual
1 7 0 0 1	, , , ,

Ethics review

Approved WMO

Date: 23-07-2013

Application type: First submission

Review commission: RTPO, Regionale Toetsingscie Patientgebonden Onderzoek

(Leeuwarden)

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

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

ID: 22608 Source: NTR

Title:

In other registers

Register ID

Other Aangemeld bij NTR CCMO NL41587.099.13 OMON NL-OMON22608