Long-term effects of a standard treatment of a depomedrol / lidocaine injection in carpal tunnel syndrome; a pilot study of the clinical and neurophysiological changes in the median nerve.

Published: 07-05-2014 Last updated: 21-04-2024

The purpose of this study is to see if we can predict which patients with CTS are eligible for an injection, and which should be directly referred to a surgeon. This to shorten the period of the patient's symptoms and to save healthcare costs.

Ethical review	Not approved
Status	Will not start
Health condition type	Peripheral neuropathies
Study type	Observational non invasive

Summary

ID

NL-OMON40966

Source ToetsingOnline

Brief title Effect of an injection in a carpal tunnel syndrome

Condition

• Peripheral neuropathies

Synonym

Median nerve compression in the carpal tunnel, nerve compression in the wrist

Research involving

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Human

Sponsors and support

Primary sponsor: HagaZiekenhuis **Source(s) of monetary or material Support:** Haga ziekenhuis neurologie

Intervention

Keyword: Carpal tunnel syndrome, depomedrol/lidcoaine, EMG, n. medianus

Outcome measures

Primary outcome

Clinical and neurophysiological changes of the median nerve after an injection

of depomedrol / lidocaine in carpal tunnel syndrome.

Secondary outcome

Study description

Background summary

Many patients with a CTS will be treated with a corticosteroïdinjectie at the carpal tunnel. After 6 weeks, the patient is monitored and reviewed for clinical improvement. It is not routinely performed to do a clinical neurophysiological research after the injection. But give the complaints of the patient the improvement in the function of the nerve and the nerve itself? And this treatment is not only a delay of surgery, and can we save time and healthcare costs by sending someone directly for surgery?

Study objective

The purpose of this study is to see if we can predict which patients with CTS are eligible for an injection, and which should be directly referred to a surgeon. This to shorten the period of the patient's symptoms and to save healthcare costs.

Study design

Prospectively studied with twenty patients with CTS symptoms at the neurology of Haga Hospital. All patients, in the period of 02-01-2014, with a the carpal tunnel and have had an injection will undergo a follow-up study. This will consist of (if not routine), an additional EMG (conduction)test, a nerve ultrasound and a questionnaire at 6 weeks and 6 months.

Study burden and risks

Contacts

Public HagaZiekenhuis

Leyweg 275 den haag 2545CH NL **Scientific** HagaZiekenhuis

Leyweg 275 den haag 2545CH NL

Trial sites

Listed location countries

Netherlands

Eligibility criteria

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

Inclusion criteria

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Carpal tunnel syndrome >18 years

Exclusion criteria

- Not Dutchspeaking/reading
- previous treatment of the CTS ipsilateral
- traumatic nerve injury
- weakness and directly surgery is indicated
- pregnancy

Study design

Design

Study type: Observational non invasive	
Masking:	Open (masking not used)
Control:	Uncontrolled
Primary purpose:	Diagnostic

Recruitment

NL	
Recruitment status:	Will not start
Enrollment:	20
Туре:	Anticipated

Ethics review

Not approved	
Date:	07-05-2014
Application type:	First submission
Review commission:	METC Leiden-Den Haag-Delft (Leiden)
	metc-ldd@lumc.nl

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 CCMO **ID** NL48797.098.14