Clinical relevance of WARP by patients whit recidive complaints after lumbar spinal fusion

No registrations found.

Ethical review	Positive opinion
Status	Pending
Health condition type	-
Study type	Observational non invasive

Summary

ID

NL-OMON25359

Source
Nationaal Trial Register

Health condition

Recidive complaints after lumbar spinal fusion

Sponsors and support

Primary sponsor: Siemens
Source(s) of monetary or material Support: -

Intervention

Outcome measures

Primary outcome

Increased signal-noise ratio, contrast-noise ratio, standard deviation

Secondary outcome

Better clinical diagnostic value

Study description

Background summary

-

Study objective

A better diagnostic image will be created after using a MRI metal artifact reduction sequence.

Study design

-

Intervention

none

Contacts

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Eligibility criteria

Inclusion criteria

Patient whit a lumbar spinal fusion. Recidive complaints after lumbar spinal fusion. All kinds of non-ferromagnetic metal.

Exclusion criteria

Ferromagnetic metal

Study design

Design

Study type:	Observational non invasive
Intervention model:	Parallel
Allocation:	Non-randomized controlled trial
Masking:	Open (masking not used)
Control:	Active

Recruitment

NL

Recruitment status:	Pending
Start date (anticipated):	01-12-2014
Enrollment:	20
Type:	Anticipated

Ethics review

Positive opinion

Date:	01-12-2014
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	NL4872
NTR-old	NTR4990
Other	: 14N119

Study results

Summary results

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