# Hybrid single-photon emission computed tomographycomputed tomography (SPECT-CT) imaging results in chronic low back pain patients as compared to an asymptomatic control group

Submission date	<b>Recruitment status</b> No longer recruiting	<ul><li>Prospectively registered</li></ul>		
26/12/2013		☐ Protocol		
Registration date	Overall study status Completed	Statistical analysis plan		
17/02/2014		[X] Results		
Last Edited	Condition category	[] Individual participant data		
16/01/2020	Musculoskeletal Diseases			

# Plain English summary of protocol

Background and study aims

For patients with low chronic low back pain that is not cured by conservative treatment, more invasive treatments can be considered. Those treatments may consist in injection, application of high frequency current or surgery. Success is dependent on how precisely the painful structure is identified. The new molecular imaging modality SPECT-CT detects increased bone metabolism during the degeneration process. It might be helpful in identifying the chronic low back pain phenotype. We want to evaluate the sensitivity of this molecular imaging modality in a group of patients suffering from chronic low back pain, as compared to a control group with no chronic back pain.

# Who can participate?

Patients older than 18 years with chronic (> 3 months) low back pain without a specific pain phenotype on MRI, CT and classical X-Ray are referred for SPECT-CT imaging. In the control group, the patients were referred for SPECT-CT for other reasons than chronic low back pain.

## What does the study involve?

Patients are asked if they consent to the analysis of their data. All patients undergo a SPECT-CT as planned within the diagnostic process for their condition.

What are the possible benefits and risks of participating?

This study is about data analysis only and there are no benefits or additional risks in participating. On rare occasions patients may be allergic to the product that is injected for imaging.

Where is the study run from? AZ Nikolaas Hospital, Sint Niklaas, Belgium. When is the study starting and how long is it expected to run for? August 2013 to January 2014.

## Who is funding the study

The SPECT-CT is part of the normal diagnostic process and is paid for by the standard procedure (health insurance). Costs for data recording and analysis are covered by the investigator.

#### Who is the main contact?

Dr Erik Van de Kelft, Director of the Neurosurgery Department, erik.vandekelft@aznikolaas.be Dr Koen Melis, Director of the Neuroradiology Department, koen.melis@aznikolaas.be

# Contact information

# Type(s)

Scientific

#### Contact name

Dr Erik Van de Kelft

#### Contact details

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# Additional identifiers

# Protocol serial number

2013-06/02

# Study information

#### Scientific Title

Evaluation of the prevalence of increased Phosphorus uptake at lumbar level during SPECT-CT in a group of patients with low back pain as compared to an asymptomatic a control group

# Study objectives

SPECT-CT shows hotspots at places with an increased phosphate metabolism due to mechanical stress. This is indicative for increased bone metabolism as result of the degeneration process. We hypothesize that the prevalence of hotspots observed during SPECT-CT will be higher in the group of patients with chronic low back pain as compared to an asymptomatic control group.

## Ethics approval required

Old ethics approval format

# Ethics approval(s)

Medical Ethics Committee of AZ Nikolaas, Sint Niklaas Belgium, Ref: EC 13022

## Study design

Prospective comparative study

## Primary study design

Observational

## Study type(s)

Diagnostic

## Health condition(s) or problem(s) studied

Sensitivity of SPECT-CT in the identification of the pain phenotype when analyzing chronic low back pain

#### **Interventions**

Patients with chronic low back pain (more than 3 months) are referred for SPECT-CT imaging. In the control group, the patients were referred for SPECT-CT for other reasons than chronic low back pain.

All patients undergo a SPECT-CT as planned within the diagnostic process for their condition. The imaging data will be analyzed.

## Intervention Type

Procedure/Surgery

## Primary outcome(s)

Number of patients with hotspots on SPECT-CT in the group with chronic low back pain, compared to the number of patients with hotspots on SPECT-CT in the group without chronic low back pain.

## Key secondary outcome(s))

Analysis of the structures that show activity in both groups. The SPECT-CT images are read by the neuroradiologist immediately after the investigation. As the value of SPECT-CT as diagnostic tool for spinal pain is measured the interpretation of the images is done only once.

# Completion date

30/01/2014

# **Eligibility**

# Key inclusion criteria

- 1. Patients older than 18 years
- 2. Referred to the department of medical imaging for SPECT-CT
- 3. Having signed the informed consent

## Participant type(s)

Patient

## Healthy volunteers allowed

No

## Age group

Adult

## Lower age limit

18 years

#### Sex

All

## Total final enrolment

200

## Key exclusion criteria

- 1. Recent vertebral fractures
- 2. History of lumbar spine surgery
- 3. Diagnosed malignancy
- 4. Pregnancy
- 5. Contraindication for injection 99mTc hydroxymethane diphosphonate (99MTc-HDP)

## Date of first enrolment

08/08/2013

## Date of final enrolment

30/01/2014

# Locations

## Countries of recruitment

Belgium

# Study participating centre

Moerlandstraat, 1

Sint Niklaas Belgium 9100

# Sponsor information

# Organisation

**AZ Nikolaas** 

# Funder(s)

## Funder type

Industry

## Funder Name

Neuro-surgery.org (Belgium)

## Funder Name

The investigation (SPECT-CT) is part of the normal diagnostic process for the patients, thus the costs involved are carried by the patients individual health insurance.

# **Results and Publications**

Individual participant data (IPD) sharing plan

# IPD sharing plan summary

Not provided at time of registration

## **Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/10/2017	16/01/2020	Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes