

Vitamin D and neuralgia in multiple myeloma

Gepubliceerd: 28-07-2016 Laatst bijgewerkt: 18-08-2022

Decreased levels of vitamin D is associated with the occurrence of peripheral neuropathy in multiple myeloma patients.

Ethische beoordeling	Niet van toepassing
Status	Anders
Type aandoening	-
Onderzoekstype	Observationeel onderzoek, zonder invasieve metingen

Samenvatting

ID

NL-OMON27690

Bron

NTR

Aandoening

Peripheral neuropathy - perifere neuropathie

Multiple myeloma - multipel myeloom

Vitamin D - vitamine D

Ondersteuning

Primaire sponsor: Medical Centre Leeuwarden

Overige ondersteuning: Wetenschapsfonds MCL

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

The correlation between PN and the vitamin D status in MM patients.

Toelichting onderzoek

Achtergrond van het onderzoek

Vitamin D status and peripheral neuropathy in multiple myeloma patients

Background: Randomized controlled trials have shown that the introduction of the novel agents

bortezomib, thalidomide and lenalidomide have improved response rates, progression-free survival

and overall survival. However, chemotherapy-induced peripheral neuropathy (CIPN), especially when

using bortezomib, is a common adverse event. In addition, several studies have found that up to 54%

of MM patients have peripheral neuropathy (PN) at diagnosis, indicating that the disease itself can

also induce PN. PN decreases quality of life and requires dose adjustment, delay or premature

termination of the treatment, resulting in a negative influence on time to progression and survival.

Vitamin D was found to reduce polyneuropathy in diabetes mellitus type 2 patients and a possible

mechanism was found in animal trials, where the investigators found an increase of nerve growth

factor in diabetic rats after supplementation of vitamin D. Recently, it was found that vitamin D

deficient MM patients were more likely to have severe CIPN (>grade 2) of both motor and sensory

PN. In this study we want to confirm the relationship of vitamin D status and the occurrence and

severity of PN in MM patients.

Objectives: The primary objective of this study is to determine the correlation between the 25-

hydroxyvitamin D serum levels and PN in patients with multiple myeloma. Secondary objectives are

to evaluate the influence of different vitamin D levels on the severity of PN; to search for differences

in prevalence of PN for each drug; to determine the correspondence of the ICPNQ results and patients' records and to gain insight in the number of patients with inadequate 25-hydroxyvitamin D

serum levels (< 75 nmol/l).

Study design: This is an observational study. Blood samples will be drawn to determine vitamin D

levels and the ICPNQ Questionnaire, a validated questionnaire to distinguish different PN grades in

MM patients, will be completed. VAS scores will be used to grade the intensity of PN.

Study population: Patients with smoldering or symptomatic multiple myeloma in the Medical Centre

Leeuwarden are included in the study. Patients must be over 18 years of age and must be able to

give informed consent.

Burden and risks: Preventing CIPN is of great importance for the continuation of chemotherapy and a

prolonged exposure will presumably result in a higher survival rate and an improved quality of life.

When this study confirms the relationship between vitamin D and neuropathy, new opportunities for

the prevention of PN may arise. As blood sampling is performed frequently in myeloma patients and

the questionnaire can be filled in within minutes, the burden for the patients is minimal.

Doel van het onderzoek

Decreased levels of vitamin D is associated with the occurrence of peripheral neuropathy in multiple myeloma patients.

Onderzoeksopzet

Vitamin D levels and neuropathy grading will be determined at one timepoint.

Onderzoeksproduct en/of interventie

- Blood samples will be drawn to determine 25-hydroxyvitamin D levels.
- All patients will complete the self-assessment survey ICPNQ and when necessary the VAS score, to distinguish different PN grades in MM patients.

Contactpersonen

Publiek

Medisch Centrum Leeuwarden

Berdien Oortgiesen
Afdeling Klinische Farmacie & Farmacologie - Postbus 888

Leeuwarden 8901 BR
The Netherlands
Tel: 058- 286 1918

Wetenschappelijk

Medisch Centrum Leeuwarden

Berdien Oortgiesen
Afdeling Klinische Farmacie & Farmacologie - Postbus 888

Leeuwarden 8901 BR
The Netherlands
Tel: 058- 286 1918

Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

Patients with smoldering or symptomatic multiple myeloma

Patients 18 years or older

Able to give informed consent

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

None

Onderzoeksopzet

Opzet

Type: Observationeel onderzoek, zonder invasieve metingen

Onderzoeksmodel: Anders

Controle: N.v.t. / onbekend

Deelname

Nederland

Status: Anders

(Verwachte) startdatum: 26-09-2016

Aantal proefpersonen: 120

Type: Onbekend

Ethische beoordeling

Niet van toepassing

Soort: Niet van toepassing

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

Geen registraties gevonden.

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register	ID
NTR-new	NL5835
NTR-old	NTR5990
Ander register	: RTPO 985

Resultaten