Leiden Improvement of MIgraine Therapy in general practice.

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Ethische beoordeling	Niet van toepassing
Status	Werving gestart
Type aandoening	-
Onderzoekstype	Interventie onderzoek

Samenvatting

ID

NL-OMON24827

Bron
NTR

Verkorte titel
LIMIT-study

Aandoening

migraine

Ondersteuning

Overige ondersteuning: ZonMw doelmatigheidsfonds
Stichting Nuts Ohra

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

HIT-6 (Headache Impact Test)-score at baseline and after 3, 6, 9 and 12 months.

Toelichting onderzoek

Achtergrond van het onderzoek

We aim to study the costs and effects of a pro-active approach of migraine patients by the General Practitioner. The design is a pragmatic cluster randomised controlled trial with the general practice as the unit of randomisation. Patients using >=24 DDD's triptans are selected from the EMD and invited by mail to consult their GP. Diagnosis and treatment plan of migraine will be evaluated and discussed. Patients will be offered prophylactic treatment according to the Dutch Guidelines (NHG-standaard). The control group will receive usual care. The primary outcome is the score on the HIT-6 questionnaire.

Doel van het onderzoek

The aim of the study is to optimize therapy of migraine patients, according to the Dutch GP Guideline for headache and consequently reduce the use of triptans. The project will explore the costs and effects of a proactive approach of patients with triptan use by GPs.

Onderzoeksproduct en/of interventie

Proactive stepped approach based on the Dutch GP Guideline versus usual care. Step 1: a letter to invite patients for consultation. Step 2: a visit to the GP, who can give information about headache and therapy, reduce/stop the triptans, prescribe prophylactic therapy or reconsider the diagnosis of migraine.
Control group: care as usual.

Contactpersonen

Publiek

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Wetenschappelijk

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

Patients in general practice using ¡Ý24 DDD triptans (or ¡Ý6 DDDs in the last 3 months), enlisted in 60 general practices, that are part of LEON (the Leiden Eerstelijns Onderzoeksnetwerk, managed by the department of Public Health and Primary Care).

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

1. Younger than 18 years;
2. Cognitive impairment;
3. Psychiatric illness;
4. Terminal illness;
5. Non-Dutch speaking.

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Blindering:	Open / niet geblindeerd
Controle:	Geneesmiddel

Deelname

Nederland
Status: Werving gestart

(Verwachte) startdatum:	01-03-2007
Aantal proefpersonen:	600
Type:	Verwachte startdatum

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	NL827
NTR-old	NTR840
Ander register	: N/A
ISRCTN	ISRCTN72421511

Resultaten

Samenvatting resultaten

N/A