Encouraging physician appropriate prescribing of non-steroidal anti-inflammatory therapies

Submission date	Recruitment status No longer recruiting	Prospectively registered		
02/08/2004		[X] Protocol		
Registration date 10/08/2004 Last Edited	Overall study status Completed Condition category	Statistical analysis plan		
		Results		
		Individual participant data		
29/10/2021	Signs and Symptoms	Record updated in last yea		

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

 ${\bf Clinical Trials. gov\ number}$

Secondary identifying numbers

N/A

Study information

Scientific Title

Encouraging physician appropriate prescribing of non-steroidal anti-inflammatory therapies

Acronym

MAAUI

Study objectives

Not provided at time of registration

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

GP practice

Study type(s)

Quality of life

Participant information sheet

Health condition(s) or problem(s) studied

Chronic pain

Interventions

All primary care physicians in Manitoba, Canada have been randomly assigned to a control group or an intervention study group. The educational intervention being evaluated consists of an audit and feedback mechanism combined with optional participation in a Continuing Medical Education interactive workshop. The primary outcome of the study is the change, from pre- to post- intervention, in physicians' appropriate prescribing of non-steroidal anti-inflammatory therapies for patients requiring chronic treatment. Three classes of non-steroidal anti-inflammatory therapies have been identified: a new class of non-steroidal anti-inflammatory agents known as coxib therapy, traditional non-steroidal anti-inflammatory drug (NSAID) monotherapy, and traditional NSAID therapy combined with gastro-protective agents. Appropriate prescribing is defined based on international clinical practice guidelines and the provincial drug reimbursement policy in Manitoba.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/11/2000

Completion date

01/01/2004

Eligibility

Key inclusion criteria

All primary care physicians in Manitoba, Canada

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/11/2000

Date of final enrolment

01/01/2004

Locations

Countries of recruitment

Canada

Study participating centre Primary Health Care Research Unit Winnipeg, Manitoba

Canada R2H 2A6

Sponsor information

Organisation

Merck Frosst Canada and Co. (Canada)

Sponsor details

C.P. 1005 Pointe-Claire-Dorval Quebec Canada H9R 4P8

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lorijean_manness@merck.com

Sponsor type

Industry

ROR

https://ror.org/028979q34

Funder(s)

Funder type

Industry

Funder Name

Merck Frosst Canada Ltd (Canada)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

Not provided at time of registration

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article	Protocol	24/08/2004		Yes	No