ISRCTN43532635 https://doi.org/10.1186/ISRCTN43532635

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

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

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.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
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Sponsor information

Organisation
Merck Frosst Canada and Co. (Canada)

Sponsor details
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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