

Improving question formulation for use in evidence appraisal in a tertiary care setting: a randomised controlled trial

Submission date 07/11/2001	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 07/11/2001	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 07/03/2008	Condition category Other	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

Study objectives

The study sought to determine whether adding simple instructions and examples on clinical question formulation would increase the specificity of the question being submitted by the health care professional compared to using a standard form without instructions and examples.

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Question formulation by health care professionals

Interventions

New participants were invited to reformulate clinical queries. The control group was given no instructions. The intervention group was given a brief explanation of proper formulation, written instructions, and diagrammatic examples.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

The change in the proportion of reformulated questions that described each the dimensions of specificity.

Secondary outcome measures

The differences in the degree by which a particular dimension was specified.

Overall study start date

01/07/2000

Completion date

30/06/2001

Eligibility**Key inclusion criteria**

Health care professionals affiliated with a tertiary care health care network in Melbourne, Australia

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

39

Key exclusion criteria

Previous users of the service

Date of first enrolment

01/07/2000

Date of final enrolment

30/06/2001

Locations**Countries of recruitment**

Australia

Study participating centre

Deputy Director

Victoria

Australia
3168

Sponsor information

Organisation

Centre for Clinical Effectiveness (Australia)

Sponsor details

Monash Institute of Health Services Research
246 Clayton Road
Clayton
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Sponsor type

Research organisation

Website

<http://www.med.monash.edu.au>

ROR

<https://ror.org/02t1bej08>

Funder(s)

Funder type

Not defined

Funder Name

Self-funded through internal research funds

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

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	01/11/2001		Yes	No