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What is the effect of alternative formats presenting the effects of antidepressants for moderate depression on decisions about whether to use them?

Submission date	Recruitment status	[ ] Prospectively registered
09/03/2007	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
04/05/2007	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
23/09/2021	Mental and Behavioural Disorders	[ ] Record updated in last year

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
What is the effect of alternative formats presenting the effects of antidepressants for moderate depression on decisions about whether to use them?

Acronym
HIPPO (Health Information Project, Presentation Online)

Study objectives
The null hypotheses for this study are that in terms of congruence between peoples' values and treatment decisions there is no difference between:
1. Words without frequencies and tabular information
2. Words without frequencies and graphically presented information, and
3. Graphically presented information and tabular information.

Ethics approval required
Old ethics approval format

Ethics approval(s)
The University at Buffalo (New York, USA), Health Sciences Institutional Review Board approved on 15 May 2002.

Study design
Randomized 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
Moderate depression (hypothetical scenario)

Interventions
Participants were randomized to view one of the following presentation formats of the effects of antidepressants for moderate depression:
1. Graphically presented information

2. Tabular information
3. Words without frequencies
4. No information

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Hypothetical treatment decision to take or not to take anti-depressants (Selective Serotonin Reuptake Inhibitors [SSRIs]).

Secondary outcome measures
Preferred presentation.

Overall study start date
17/10/2005

Completion date
05/02/2006

Eligibility

Key inclusion criteria
At least 18 years of age and fluent in Norwegian.

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Sex
Not Specified

Target number of participants
1400

Key exclusion criteria
Previous participation in this trial.

Date of first enrolment
17/10/2005

Date of final enrolment

05/02/2006

Locations

Countries of recruitment
Norway

Study participating centre
PO box 7004
Oslo
Norway
N-0130

Sponsor information

Organisation
Norwegian Knowledge Centre for the Health Services (Norway)

Sponsor details
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Sponsor type
Other

Website
http://www.kunnskapssenteret.no/

ROR
https://ror.org/01thff661

Funder(s)

Funder type
Research council

Funder Name

Norwegian Research Council (Norway)

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