

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

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

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.

Primary study design

Interventional

Study design

Randomized controlled trial

Study type(s)

Not Specified

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(s)

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

Key secondary outcome(s)

Preferred presentation.

Completion date

05/02/2006

Eligibility

Key inclusion criteria

At least 18 years of age and fluent in Norwegian.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 Years

Sex

Not Specified

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)

ROR

<https://ror.org/01thff661>

Funder(s)**Funder type**

Research council

Funder Name

Norwegian Research Council (Norway)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration