ISRCTN18030685 https://doi.org/10.1186/ISRCTN18030685

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

Registration date
04/05/2007

Last Edited
23/09/2021

Recruitment status
No longer recruiting

Overall study status
Completed

Condition category
Mental and Behavioural Disorders

[ ] Prospectively registered
[ ] Protocol
[ ] 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

Contact name
Ms Cheryl Carling

Contact details
PO box 7004
St Olavs Plass
Oslo
Norway
N-0130
+47 98627327
cheryl.carling@kunnskapssenteret.no

Additional identifiers

Protocol serial number
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

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