Bipolar and Childwish: Considerations and thoughts of women with bipolar disorder about family planning and pregnancy.

Published: 18-07-2012
Last updated: 26-04-2024

Purpose of the study is to understand women with bipolar disorder in their decision-making process concerning family planning or prior to any pregnancy.

Ethical review	Approved WMO
Status	Recruitment stopped
Health condition type	Manic and bipolar mood disorders and disturbances
Study type	Observational non invasive

Summary

ID

NL-OMON37332

Source
ToetsingOnline

Brief title
Biplar and Childwish

Condition

- Manic and bipolar mood disorders and disturbances

Synonym
Bipolar disorder, Family planning

Research involving
Human

Sponsors and support

Primary sponsor: Dimence (Deventer)
Source(s) of monetary or material Support: Ministerie van OC&W

Intervention

Keyword: Bipolar disorder, Childwish, Pregnancy

Outcome measures

Primary outcome

Insight into the perception of the decision whether or not to have children of

women with bipolar disorder.

Secondary outcome

Understanding what women need and expect from their clinician during the

decision whether or not to become pregnant.

Study description

Background summary

In the Netherlands, approximately 1.3% of the population between 18 and 65 have the diagnose bipolar disorder (de Graaf, 2011). For young adults of childbearing age, the diagnose can affect the pregnancy wish. For women, play both the biological and psychological factors a role (Yonkers, 2004). The way women experience the decision-making process about pregnancy and family planning is not known from the literature. It could be important for the treatment and support during this process to understand the thoughts and considerations of women about family planning and prior to any pregnancy.

Study objective

Purpose of the study is to understand women with bipolar disorder in their decision-making process concerning family planning or prior to any pregnancy.

Study design

The study conducts a general qualitative design.

Study burden and risks

Participation is free of significant risks to health or physical function.

Respondents may withdraw from the study at any time. The interview will take 45 - 60 min. Participants who may be treated by a specialist, will be advised her therapist to be aware of participation.

Contacts

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age
Adults (18-64 years)
Elderly (65 years and older)

Inclusion criteria

Bipolar disorder
Women in the childbearing age without childeren

Exclusion criteria

Women with childeren

Study design

Design

Study type: Observational non invasive

Masking:	Open (masking not used)
Control:	Uncontrolled
Primary purpose:	Prevention

Recruitment

NL

Recruitment status:	Recruitment stopped
Start date (anticipated):	31-07-2012
Enrollment:	15
Type:	Actual

Ethics review

Approved WMO

Date:	18-07-2012
Application type:	First submission
Review commission:	METC Universitair Medisch Centrum Utrecht (Utrecht)

Approved WMO

Date:	05-09-2012
Application type:	Amendment
Review commission:	METC Universitair Medisch Centrum Utrecht (Utrecht)

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
CCMO	NL38614.041.12