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The utility of standardised pre-clinical assessment based on the Development And Well-Being Assessment (DAWBA) in a child and adolescent mental health service

Submission date
01/08/2007

Recruitment status
No longer recruiting

Registration date
05/09/2007

Overall study status
Completed

Last Edited
21/03/2013

Condition category
Mental and Behavioural Disorders

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[X] Results
[ ] Individual participant data

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

Study objectives
1. The agreement between pre-clinical assessment (DAWBA based) and blind clinical assessment is incomplete.
2. The disclosure of pre-clinical assessment diagnoses will have an effect on clinical diagnoses including comorbidities.
3. Parents and adolescents will display acceptance and satisfaction with pre-clinical assessment based on the DAWBA.

Ethics approval required
Old ethics approval format

Ethics approval(s)
The trial was approved by the Ethical Committee of the Canton of Zurich, Specialized Subcommittee Psychiatry, Neurology, Neurosurgery, on the 1st June 2007.

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Diagnostic

Participant information sheet

Health condition(s) or problem(s) studied
Childhood psychiatric disorders

Interventions
The provision of diagnostic information based on the DAWBA to the clinicians treating the intervention group. The control group will undergo a normal clinical assessment. The diagnostic information will be gathered from parents, adolescents above the age of 11 years, and teachers.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
The level of agreement between the pre-clinical assessment and the clinical assessment.

Secondary outcome measures
Clinical routine ratings by the clinicians and parents.

Overall study start date
01/09/2007

Completion date
31/08/2008

Eligibility

Key inclusion criteria
The trial will include referrals from four outpatient clinics of the service. Patients will be aged 5-18 years.

Participant type(s)
Patient

Age group
Child

Lower age limit
5 Years

Upper age limit
18 Years

Sex
Both

Target number of participants
500

Key exclusion criteria
Children of parents with insufficient unterstanding of German and emergency referrals will be excluded.

Date of first enrolment
01/09/2007

Date of final enrolment

31/08/2008

Locations

Countries of recruitment
Switzerland

Study participating centre
Department of Child and Adolescent Psychiatry
Zurich
Switzerland
CH-8032

Sponsor information

Organisation
University of Zurich (Switzerland)

Sponsor details
Neumuensterallee 9
Postfach
Zurich
Switzerland
CH-8032

Sponsor type
University/education

Website
http://www.uzh.ch/about_en.html

ROR
https://ror.org/02crff812

Funder(s)

Funder type
University/education

Funder Name
Research budget of the Department of Child and Adolescent Psychiatry, University of Zurich (Switzerland)

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	01/10/2012		Yes	No