

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	☐ Prospectively registered
Registration date 05/09/2007	Overall study status Completed	☐ Protocol
Last Edited 21/03/2013	Condition category Mental and Behavioural Disorders	☐ Statistical analysis plan
		☒ Results
		☐ Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

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.

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Diagnostic

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(s)

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

Key secondary outcome(s)

Clinical routine ratings by the clinicians and parents.

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

Healthy volunteers allowed

No

Age group

Child

Lower age limit

5 Years

Upper age limit

18 Years

Sex

All

Key exclusion criteria

Children of parents with insufficient understanding 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)

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

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