Randomised controlled trial of Triple P intervention for multi-problem families [Opvoedingsondersteunings programma Triple P]

Submission date 22/11/2006	Recruitment status No longer recruiting	Prospectively registered
Registration date	Overall study status	 Protocol Statistical analysis plan
22/11/2006	Completed	[] Results
Last Edited 14/10/2008	Condition category Mental and Behavioural Disorders	 Individual participant data Record updated in last year
14/10/2000		

Plain English summary of protocol

Not provided at time of registration

Study website http://www.trimbos.nl

Contact information

Type(s) Scientific

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Contact details

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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

The tested intervention (level five intervention of Triple P, an intensive individual training for the family of children with behavioural problems) will cause a decrease in behavioural problems in children, dysfunctional parenting styles in parents, and depression, anxiety and stress in parents. Furthermore it will cause an increase in parenting competences in parents.

Ethics approval required

Old ethics approval format

Ethics approval(s) Received from the local medical ethics committee

Study design Multicentre, randomised, active controlled, parallel group trial

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) Not specified

Study type(s) Treatment

Participant information sheet

Health condition(s) or problem(s) studied Behavioural and emotional problems in children

Interventions

The Triple P (Positive Parenting Program) consists of five levels, ranging from brief and broad support to parents in dealing with a child who exhibits behavioural problems to an intensive family intervention (level five). This intervention is aimed at family dysfunction and serious behavioural problems in a child. It offers intensive individual training for the family (around eleven sessions of one hour each, divided in modules), focusing on the childs behavioural problems (in combination with problems of the parents such as conflict between the partners, depression and stress).

Intervention Type

Other

Phase Not Specified

Primary outcome measure Decrease of behavioural and emotional problems in children.

Secondary outcome measures

1. Decrease of dysfunctional parenting styles

2. Depression, anxiety and stress in parents

3. Increase of parenting competences

Overall study start date 01/10/2006

Completion date 31/12/2008

Eligibility

Key inclusion criteria

Multi-problem family
 Children between four to 13 years old

Participant type(s)

Patient

Age group Child

Lower age limit 4 Years

Upper age limit 13 Years

Sex Both

Target number of participants 200

Key exclusion criteria Does not comply with the above inclusion criteria

Date of first enrolment 01/10/2006

Date of final enrolment

31/12/2008

Locations

Countries of recruitment Netherlands

Study participating centre Trimbos Institute - Netherlands Institute of Mental Health and Addiction Utrecht Netherlands 3500 AS

Sponsor information

Organisation Trimbos Institute - Netherlands Institute of Mental Health and Addiction (The Netherlands)

Sponsor details

P.O. Box 725 Utrecht Netherlands 3500 AS +31 (0)30 2971100 info@trimbos.nl

Sponsor type Research organisation

ROR https://ror.org/02amggm23

Funder(s)

Funder type Research organisation

Funder Name The Netherlands Organization for Health Research and Development (ZonMw) (The Netherlands)

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