Effectiveness of the (POPP) treatment program in sexually abused preschool girls: a randomized controlled trial

Gepubliceerd: 22-10-2015 Laatst bijgewerkt: 15-05-2024

POPP treatment will lead to more PTSD symptom reduction compared to wait list.

Ethische beoordeling	Positief advies
Status	Werving gestart
Type aandoening	-
Onderzoekstype	Interventie onderzoek

Samenvatting

ID

NL-OMON20992

Bron
Nationaal Trial Register

Verkorte titel
POPP treatment for sexually abused preschool girls

Aandoening

PTSD symptoms

Ondersteuning

Primaire sponsor: GGZ Oost Brabant
Overige ondersteuning: GGZ Oost Brabant

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

PTSD symptoms from baseline to post-intervention. The change in PTSD symptoms between the experimental and control group is compared at post-intervention.

Toelichting onderzoek

Achtergrond van het onderzoek

The main objectives of the present study are: assessing the effectiveness of the POPP-treatment program in girls aged 3-6 years with posttraumatic stress reactions following sexual abuse

Doel van het onderzoek

POPP treatment will lead to more PTSD symptom reduction compared to wait list.

Onderzoeksopzet

The assessments will be conducted at baseline (T1), three months later at post-intervention (post-immediate POPP treatment, or at the end of waiting list: T2), and six months later (follow-up of immediate POPP treatment or post-treatment in the waiting list condition: T3)

Onderzoeksproduct en/of interventie

The POPP treatment program is based on Trauma Focused-Cognitive Behavior Therapy (TF-CBT) techniques, with a newly described central position for projection and healing play. Ten treatment sessions (for a maximum of 60 minutes each) are offered, along with parental guidance. Under the waiting list condition, the POPP treatment program starts after three months.

Contactpersonen

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Wetenschappelijk

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

The criteria for inclusion are a) increased levels of PTSD symptoms, b) PTSD symptoms related to sexual abuse, and c) the child has reached the developmental level of symbolic play.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

Exclusion criteria are a) the absence of parental permission, b) severe psychiatric conditions that require an emergency response (suicidal intent, psychotic symptoms, or severe dissociations), c) the most recent episode of sexual abuse occurred more than 12 months before referral to the study, and d) lack of a long-term caretaker or severe family problems (lack of stability for the ongoing six months).

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Toewijzing:	Gerandomiseerd
Blindering:	Enkelblind
Controle:	Geneesmiddel

Deelname

Nederland

Status:	Werving gestart
(Verwachte) startdatum:	01-07-2015
Aantal proefpersonen:	40
Type:	Verwachte startdatum

Ethische beoordeling

Positief advies

Datum:	22-10-2015
Soort:	Eerste indiening

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

ID: 41070
Bron: ToetsingOnline
Titel:

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register	ID
NTR-new	NL5307
NTR-old	NTR5416
CCMO	NL50473.091.14
OMON	NL-OMON41070

Resultaten

Samenvatting resultaten
n/a