Treatment of PTSD in children

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The present study aims at the improvement of the treatment of children with PTSD by examining the effectiveness of a brief intensive exposure therapy (IET) for children with PTSD using a multiple baseline design (patients are randomly assigned to...

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

Samenvatting

ID

NL-OMON21205

Bron
NTR

Aandoening

Posttraumatic stress disorder (PTSD)

Ondersteuning

Primaire sponsor: Prof. dr. R.C.M.E. Engels
Behavioural Science Institute, Radboud University Nijmegen
Overige ondersteuning: ZonMw

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

PTSD symptoms, self-report by child: UCLA PTSD Reaction Index (PTSD-RI)

Toelichting onderzoek

Achtergrond van het onderzoek

The present study aims to examine the effectiveness of a brief intensive exposure therapy for children with PTSD. The new treatment program makes use of proven effective therapy techniques in processing the trauma and decreasing PTSD symptoms in children, whereas the delivery of the treatment is using a new format: one week of intensive trauma-focused treatment while including a trauma-related social support program for family members.

Country of Recruitment: The Netherlands

Doel van het onderzoek

The present study aims at the improvement of the treatment of children with PTSD by examining the effectiveness of a brief intensive exposure therapy (IET) for children with PTSD using a multiple baseline design (patients are randomly assigned to baseline length).

Onderzoeksopzet

Time points of primary interest (primary outcome)

Weekly during:

- Baseline length (varying from 4 to 8 weeks before treatment)

- Treatment length (5 weeks)

- Post treatment length (varying from 4 to 8 weeks after treatment)

- Follow up length (during 4 weeks, 3 months after treatment)

Time points secondary outcomes

- before treatment (A0)

- three weeks after baseline (A1)

- three months after baseline (A2)

- and six months after baseline (A3)

Onderzoeksproduct en/of interventie

Brief intensive modified exposure therapy and a trauma-related social support program for family members:

5 days (one week), offered in three blocks of 90 minutes each day and up to three follow up appointments (90 minutes each)

Contactpersonen

Publiek

GGZ Nijmegen
Tarweweg 2

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The Netherlands
024-3837820

Wetenschappelijk

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The Netherlands
024-3837820

Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

(1) Current DSM-IV diagnosis of PTSD established with the PTSD section of the Kiddie-Schedule for Affective Disorders and Schizophrenia-Present and Lifetime Version (K-SADS-PL)

(2) Age between 7 and 18 years (boys & girls)

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

(1) Psychosis or delusion disorders (current or in the past)

(2) Suicidality

(3) Mental retardation

(4) Substance abuse or dependence or alcohol abuse or dependence

(5) Insufficient ability to speak and write Dutch

(6) Trauma caused by a caregiver who is part of the current primary care system

(7) Current DSM-IV diagnosis of PTSD of the caregiver established with a structured diagnostic interview (M.I.N.I.) and the PTSD Symptom Scale Self-Report (PSS-SR)

Onderzoeksopzet

Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Cross-over
Toewijzing:	Gerandomiseerd
Blindering:	Open / niet geblindeerd
Controle:	N.v.t. / onbekend

Deelname

Nederland

Status:	Werving gestopt
(Verwachte) startdatum:	20-12-2012
Aantal proefpersonen:	10
Type:	Werkelijke startdatum

Ethische beoordeling

Positief advies

Datum:	02-12-2013
Soort:	Eerste indiening

Registraties

Opgevolgd door onderstaande (mogelijk meer actuele) registratie

Geen registraties gevonden.

Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

In overige registers

Register	ID
NTR-new	NL4149
NTR-old	NTR4301
Ander register	*ZonMW & **CCMO : *80-82470-98-006-04 & **NL36971.091.11
ISRCTN	ISRCTN wordt niet meer aangevraagd.

Resultaten

Samenvatting resultaten
N/A