Societal participation and overcoming adversity

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

Samenvatting

ID

NL-OMON26346

Bron
NTR

Aandoening

victimization
societal participation
severe mental illness
victimisatie
maatschappelijke participatie
ernstige psychische aandoeningen

Ondersteuning

Primaire sponsor: Tilburg University
Stichting Rehabilitatie '92
GGz Eindhoven
Dijk & Duin
Overige ondersteuning: The Netherlands Organisation for Scientific Research (NWO)

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

The primary study parameters will be safety and societal participation.

Toelichting onderzoek

Achtergrond van het onderzoek

Background
People with severe mental illness (SMI) in the Netherlands are more often victims than perpetrators of violence, and are also more often victimized than other Dutch citizens. This victimization appears to be one of the risks that these service users of mental health care have to deal with in their efforts to participate in the community. Other risks in societal participation are rejection, failure, loss and social stigma with possible adverse consequences.

Objective of the study
The main objective of the study is to gain insight in the effectiveness of the Victoria modules, in terms of acknowledgement for these experiences and risk awareness. These experiences and insights can be used to take acceptable risks in rehabilitation process. The focus is on increasing societal participation, reducing victimization and other forms of adversity.

Intervention
The first part of the module aims at exploring victimization experiences or other forms of adversity that service users have experienced in societal participation. The second part focuses on the evaluation of rehabilitation crises, with specific attention to the social context in which the crisis has occurred and environmental triggers that brought the service user off balance.

Study design and methods
The research will be carried out as a multicentre RCT, in which 346 outpatients will be included at baseline. Randomization is at a team level. A total of 8 FACT-teams of 2 institutes for mental health care are included. There are three moments of measurement: at the start (T0), after 9 months (T1), and after 18 months of intervention (T2).
Generalized linear mixed modelling is used to examine whether there is an effect on primary and secondary outcome parameters (societal participation, victimization, self-confidence,

stigmatization and discrimination, and quality of life).

Doel van het onderzoek

The Victoria modules are thought to have a positive effect on acknowlegdement for adversities and risk awareness. This can be used to take acceptable risks in rehabilitation process. This will then lead to more societal participation and less victimization and other forms of adversity.

Onderzoeksopzet

T0 = no intervention, starts Feb. 2016.

training in Victoria.

T1 = experiment and control group, 9 months after T0.

T2 = experiment and control group, 9 months after T1.

Onderzoeksproduct en/of interventie

Victoria is a methodology to explore victimization (risks) attached to societal participation with severe mental illness, in the form of a conversation module.

The experimental teams receive the Victoria modules, and the control groups receive care as usual.

Contactpersonen

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

- having a Severe Mental Illness

- receiving care from a Flexible Assertive Community Treatment team of the Mental Health Care Centres Dijk en Duin or Eindhoven

- Older than 17 years of age

- Willing to participate in the research

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

- younger than 18 years of age

- insufficient command of the Dutch language

- incapable to answer questions due to cognitive impairments, severe symptomatology, or psychoorganic disorders

- prolonged admission to psychiatric hospital, or staying in prison

Onderzoeksopzet

Opzet

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

Deelname

Nederland

Status:	Werving gestopt
(Verwachte) startdatum:	01-02-2016
Aantal proefpersonen:	346
Type:	Werkelijke startdatum

Voornemen beschikbaar stellen Individuele Patiënten Data (IPD)

Wordt de data na het onderzoek gedeeld: Nee

Ethische beoordeling

Positief advies

Datum:	11-01-2016
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	NL4172
NTR-old	NTR5585
Ander register	CCMO NL53845.028.15 : NWO 432-12-808

Resultaten