

Versterken van eigen emotionele veerkracht tijdens behandeling met antidepressiva.

Gepubliceerd: 31-08-2009 Laatste bijgewerkt: 18-08-2022

1. Individualized feedback on the experience of positive emotions in daily life situations (activities, events and social interactions) increases the ability of patients to generate positive emotions. In addition, we hypothesize therefore that...

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

Samenvatting

ID

NL-OMON25509

Bron

Nationaal Trial Register

Verkorte titel

REMID-ID

Aandoening

depression (depressie)

Ondersteuning

Primaire sponsor: J.J. Van Os, head of department Psychiatry & Neuropsychology, University of Maastricht

Overige ondersteuning: Zon-Mw

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

Main outcomes are pre-post intervention change in:

1. Depressive symptomatology;

2. Daily life emotional experience (in terms on NA and PA) and person-context interaction;

3. Pre-post intervention

change on the empowerment questionnaire;

4. Health care

consumption and costs. In addition, long-term outcome will be examined.

Toelichting onderzoek

Achtergrond van het onderzoek

Rationale:

Recent evidence indicates that individual daily life person-context interactions determine vulnerability for depression, and predict relapse as well as recovery. We hypothesize that momentary assessment technology (with the Psy-mate) to monitor these daily life person-context interactions during treatment will enrich passive antidepressant pharmacotherapy with an active resource-mobilising psychotherapeutic context, thus enhancing therapeutic efficacy (reducing symptoms and relapse rate).

Objective:

(1) Does feedback on continuous monitoring with the Psy-mate during pharmacotherapy result in a better treatment response (immediate and follow-up reductions of depressive symptoms, decreased relapse risk)? (2) Do measurements of daily life person-context interactions improve the prediction of future courses of depressive symptoms? (3) Can individual differences in (1) the effect of feedback on daily life person-context interaction and (2) the predictive value of these person-context interactions be traced back to genetic differences?

Study design:

Randomized clinical trial with three groups of 40 subjects each (2 experimental groups and 1 control group). The first group receives a 5-day pre- and post Psy-mate assessment and a continuous Psy-mate assessment (3 days of Psy-mate measurements during a 6-weeks period) with weekly feedback (to both patient and therapist) during treatment as usual (TAU). The second group also receives a 5-day pre- and post Psy-mate assessment and a continuous Psy-mate assessment (3 days of Psy-mate measurements during a 6-weeks period) but without feedback during treatment as usual (TAU). The third group receives a

5-day pre- and post Psy-mate assessment but no additional intervention during TAU. This is the control group.

Study population:

A sample of

patients (n=120) with a depressive disorder will be recruited within (i) SEARCH, the research network of mental health institutions within Limburg and South-Brabant, The Netherlands, (ii) general practices and (iii) the general population.

Intervention (if applicable):

The Psy-mate

is a recently developed wearable interactive palmtop suitable for the Experienced Sampling Method (ESM) to study subjects in their daily life.

The intervention group receives feedback on Psy-mate measurements of the experience of positive emotions in daily life.

Main study

parameters/endpoints:

The rate of change in reward experience measured with the Psy-mate, and the change in depression symptomatology measured with the Structured Clinical Interview for DSM-IV Axis I Disorders (SCID-I) and the Hamilton Depression Rating Scale (HDRS).

Doel van het onderzoek

1. Individualized feedback on the experience of positive emotions in daily life situations (activities, events and social interactions) increases the ability of patients to generate positive emotions. In addition, we hypothesize therefore that the feedback condition combined with antidepressant treatment compared to antidepressant treatment alone is more effective in reducing immediate and follow-up depressive symptomatology and future relapse;

2. Giving individualized feedback on relevant daily life person-context interactions during pharmacotherapy is cost-effective compared to treatment as usual;

3. Measurement of relevant dynamic daily life behaviours significantly contributes to the prediction of the future course of depressive symptomatology;

4. Individual differences in (i) the effect of feedback on daily life person-context interactions and (ii) the predictive value of these person-context interactions can be traced back to genetic differences;

5. Subjects who receive feedback on their experience of positive

emotions in daily life will develop an increase in feelings of empowerment compared to subjects who do not receive feedback.

Onderzoeksopzet

Screening:

Hamilton Depression Rating Scale (HAM-17), Structured Clinical Interview for DSM-IV Axis I Disorders (SCID-I), personality inventory (NEO-PI), Interview for depression symptomatology (IDS), trauma (Childhood Trauma Questionnaire and Recent life Event Scale), PRODISQ, TiC-P, Euroquol-5D and a saliva sample.

Baseline:

5 days of measurement with the Psy-mate, and after completion of this period: the depression part of the SCID-I, the HAM-17, and the IDS.

Intervention period (week 2 to 8):

1. Intervention group: 3 days/week during a 6 week period of Psy-mate measurements combined with weekly feedback during treatment as usual (TAU);
2. Control group 1: 3 days/week during a 6 week period of Psy-mate measurements without weekly feedback during TAU;
3. Control group 2: no Psy-mate measurements, no feedback during TAU.

At the end of the intervention period, all 3 groups have to complete the depression part of the SCID-I, the HAM-17, the IDS, and the Empowerment questionnaire.

Post intervention measurement:

5 days of measurement with the Psy-mate, and after completion of this period: the depression part of the SCID-I, the HAM-17, and the IDS.

Follow-up measurements:

1. 4 weeks after the intervention period: the depression part of the SCID-I, the HAM-17, and the IDS;
2. 8 weeks after the intervention period: the depression part of the SCID-I, the HAM-17, and

the IDS;

3. 12 weeks after the intervention period: the depression part of the SCID-I, the HAM-17, the IDS, TiC-P, PRODISQ, EQ-5D;

4. 24 weeks after the intervention period: the depression part of the SCID-I, the HAM-17, the IDS, TiC-P, PRODISQ, EQ-5D, Interview for Recent Life Events.

Onderzoeksproduct en/of interventie

The Psy-mate is a recently developed wearable interactive palmtop suitable for the Experience Sampling Method (ESM) to study subjects in their daily life. The intervention group receives feedback on Psy-mate measurements of positive affect in their daily life.

Contactpersonen

Publiek

Universiteit Maastricht
Capaciteitsgroep Psychiatrie en Neuropsychologie, UM/FHML
Locatie Vijverdal, SN-gebouw
Postbus 616

Marieke Wichers
Maastricht 6200 MD
The Netherlands
+31-43 - 36 88 669

Wetenschappelijk

Universiteit Maastricht
Capaciteitsgroep Psychiatrie en Neuropsychologie, UM/FHML
Locatie Vijverdal, SN-gebouw
Postbus 616

Marieke Wichers
Maastricht 6200 MD
The Netherlands
+31-43 - 36 88 669

Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

1. 18-65 years;
2. DSM IV diagnosis of current major depressive disorder or a past DSM-IV diagnosis of major depression with residual symptoms (HAM-17>7);
3. Use of antidepressants or moodstabilizers;
4. Adequate vision;
5. Sufficient Dutch language skills.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

No (hypo) manic or mixed episode within the past month.

Onderzoeksopzet

Opzet

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

Deelname

Nederland	
Status:	Werving gestart
(Verwachte) startdatum:	01-01-2010
Aantal proefpersonen:	120

Type:

Verwachte startdatum

Voornemen beschikbaar stellen Individuele Patiënten Data (IPD)

Wordt de data na het onderzoek gedeeld: Nog niet bepaald

Ethische beoordeling

Positief advies

Datum:

31-08-2009

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	NL1862
NTR-old	NTR1974
Ander register	METC Unimaas : M09-1935
ISRCTN	ISRCTN wordt niet meer aangevraagd.

Resultaten

Samenvatting resultaten

N/A