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Brief Report

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Introducing a Group Therapy Program (PLAN D) for Young Outpatients with Derealization and Depersonalization: A Pilot Study

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Keywords

Derealization · Depersonalization · Cognitive behavioral therapy · Group intervention · Outpatients

Abstract

Depersonalization and derealization (DD) cause significant distress and are associated with poor role and social functional outcomes. Despite the relatively high prevalence of DD symptoms and the chronic course in those suffering from a DD disorder, there still exists a need for effective interventions. Preliminary evidence indicates that cognitive behavioral therapy (CBT) delivered in an individual setting demonstrates some positive intervention effects for patients with DD regarding their symptom levels. By considering DD-specific treatment needs, a group therapy program was developed as an add-on therapy based on CBT techniques called PLAN D comprising the following elements: psychoeducation, lifestyle interventions, acceptance and mindfulness training, and new patterns of DD-related cognitions. In a pilot study, we present an 8-week group intervention for adolescents and young adults with DD disorder. To our knowledge, no standardized group intervention program for DD exists so far. Thus, this novel intervention represents a promising opportunity to positively influence long-term outcomes and course of DD.

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Introduction

"I feel like I am watching my life from behind glass, as if I am living in a dream" – this is a typical statement from someone experiencing derealization and/or depersonalization (DD). DD is a perceptional reaction and psychological phenomenon, especially occurring when individuals are highly distressed, traumatized, very tired, anxious, or intoxicated [1]. In the Diagnostic and Statistical Manual of Mental Disorders, Fifth Edition (DSM-5) [2], clinically relevant DD is diagnosed as depersonalization/ derealization disorder in the chapter of dissociative disorders. Therein, it is defined as a persistent or recurrent experience of unreality and detachment from oneself or their surroundings, while reality testing remains intact and the symptoms cause clinically significant distress or impairment in functioning [2].

Prevalence rates of 1.2–2.4% were found for clinically significant DD symptoms in the community and of 30– 82% in clinical samples, in a systematic review [3]. Furthermore, DD (disorder) often exists as a comorbidity, especially in depression and anxiety [4]. Most DD experiences are transient and disappear as situational factors ease. However, in a relevant minority of cases, DD persists for days, weeks, or months, reoccurs, or remains permanent [3]. Age of onset is adolescence, and an earlier onset is associated with higher severity and poorer prog-

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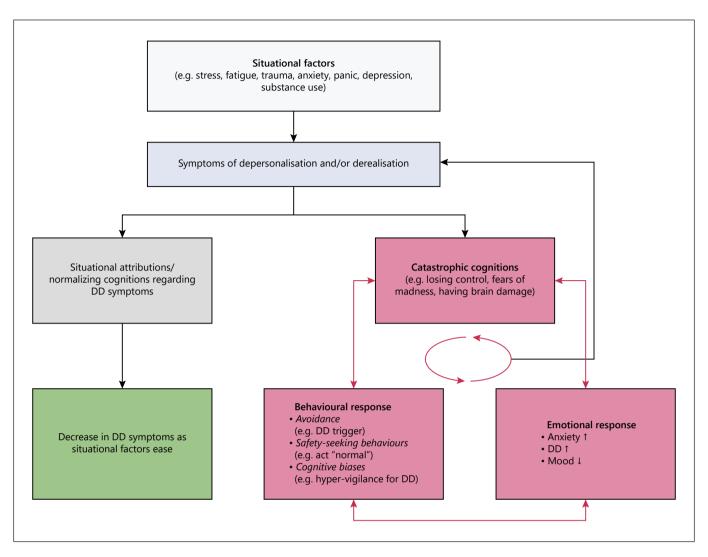


Fig. 1. Cognitive behavioral model of DD adapted from Hunter [14] explaining development and maintenance of DD disorder. DD, derealization/depersonalization.

nosis [4, 5]. Individuals with DD frequently are distressed, worry about their mental state, and are frightened of becoming crazy, a common fear not only in patients who experience DD but also in patients experiencing ultrahigh-risk (UHR) symptoms. Furthermore, chronic DD often is associated with functional impairments [1, 6, 7]. Apart from an isolated DD disorder or comorbid phenomena, DD commonly appears in subjects at clinical high risk (CHR) for psychosis. Thus derealization, but not depersonalization, is one of the 14 basic symptoms that were shown to be specific to the development of firstepisode psychosis [8]; basic symptoms defined as self-experienced disturbance of "normal" mental processes [9, 10] are employed in 2 CHR criteria [11–13]: cognitive disturbances (COGDIS) and cognitive-perceptive (COP- ER) basic symptoms. They are part of the 2 complementary early detection approaches to the characterization of the CHR state of psychoses: the UHR and the basic symptoms criteria [13]. The (attenuated) positive symptoms as part of the UHR criteria are distinct from the basic symptoms since they are experienced by the individual – depending on the insight into its abnormal nature – as real, normal thinking, and feeling [9, 10]. Derealization as part of the COPER criterion is rated as below attenuated positive symptom level (for more details to this distinction, see [10]). Taken together, the high prevalence of DD symptoms, its role as a psychosis risk symptom (according to basic symptoms), and the chronic course in those suffering from a DD disorder highlight the need for effective targeted interventions.

Therefore, Hunter et al. [1, 6] conceptualized a cognitive behavioral model for the development and maintenance of DD (Fig. 1) based on etiological models for anxiety disorders (e.g., panic, health anxiety). The model suggests that chronic DD may result from catastrophic interpretation of common, normally transient symptoms of DD as being indicative of a serious mental disorder. This leads to an exacerbation and perpetuation of DD symptoms through the development of cognitive biases and behaviors forming a maintenance cycle [1, 14]. Although the cognitions are disorder specific, the similarity between anxiety disorder and DD lies in this misinterpretation [6, 15]. Accordingly, Hunter et al. [6] modified cognitive behavioral therapy (CBT) for anxiety disorders to DD, respectively, for their specific thought contents which resemble a "mental health anxiety." They created a CBT protocol in 3 phases with the following interventions: (1) psychoeducation and normalizing and diary keeping, (2) reducing avoidance and safety behavior as well as self-focused attention and symptom monitoring, and (3) focusing on maintaining progress and relapse. They investigated this CBT protocol in an open trial investigating a sample of 21 patients with DD disorder (mean age 38 years) and with a mean duration of 14 years of DD. In an open 2-year study, the intervention was carried out with a mean number of 13 sessions. After intervention, 29% of the participants no longer met DD criteria. While they found a significant reduction in clinicianrated DD symptoms, to our knowledge no replication study currently exists.

Considering the paucity of literature on psychological interventions in DD, there are several specific interventions considered to be successful in patients with DD: DD assessment and exploration of relevant factors which enhance and reduce DD symptoms, psychoeducation, training of grounding strategies or mindfulness, lifestyle interventions (e.g., sleep, substance use, activity), reducing avoidance, safety behavior, symptom monitoring and self-focused attention, cognitive restructuring techniques to challenge negative automatic DD-related cognitions, techniques to facilitate the controlled re-experience of emotions as well as targeting the high arousal and anxiety level [5, 7, 14, 16–19]. An important clinical feature in DD patients is the often-reported emotional detachment. It has been shown that chronic DD disorder patients showed different skin conductance response to emotional stimuli compared to healthy or clinical controls meaning a reduced autonomic response to unpleasant stimuli, suggesting a selective inhibitory mechanism on emotional processing [20]. A replication study with DD patients

suggested that the cognitive evaluation of emotional stimuli is disconnected from their bodily or autonomic response, respectively [21], which reflects an increased introversion and control of emotional impulses. In the same study, mindfulness exercises were found to be helpful in modulating this autonomic response since they immediately decreased DD intensity [21]. In sum, primarily CBT elements with a special focus on DD-specific cognitions, enriched by emotion regulation techniques such as mindfulness and relaxation training, are applied and conferred to be helpful in DD. Additionally, acceptance and commitment therapy (ACT) [22] is also discussed as a useful element in DD, especially because ACT targets experiential avoidance, which is assumed in dissociative conditions such as DD [23]. Despite this detailed DD conceptualization and DD-specific knowledge, to date, no standardized treatment protocol for DD has been established, and the need for (novel) therapeutic approaches is still ongoing. To address this, we report on the development and implementation of a DD-specific group program targeting young patients with DD disorder by giving them a first CBT-based intervention.

Materials and Methods

Rationale and Conceptualization of PLAN D

In the "Bern Early Detection and Intervention Center for Mental Crises" (FETZ Bern), where we apply an indicated prevention approach of psychoses, DD is a frequently reported phenomenon among our help-seeking outpatients (30% with intermittent DD symptoms, 8% with DD disorder). Based on our indicated prevention approach, we aimed to develop a structured psychotherapy manual for individuals with DD and to deliver this psychotherapy as a group program, in order to bring DD patients together and foster group processes, which is known to facilitate normalization of experiences and receive early psychological intervention. In the FETZ Bern, the approach of normalizing experiences and fears related to these symptoms is generally used by the FETZ therapists not only in the treatment of DD but also in CBT offered to patients with UHR symptoms, where an evidence-based psychotherapy by van der Gaag et al. [24] is used. Almost every UHR patient fears the loss of control of his/her psyche, and the therapeutic approach in the FETZ Bern aims to emphasize that the patient is not crazy and to create hope for recovery. The program's name "PLAN D" not only encompasses the various elements and intervention techniques (Table 1) but also takes into account that many DD patients often try a lot of, and struggle with, other plans to eliminate DD experiences before seeking help. To that end, PLAN D should present an alternative, more reasonable plan. The group program follows a CBT approach that considers the DD-specific interventions that exist in the current literature by including elements of ACT. PLAN D is provided as an add-on therapy for 4 to 6 adolescent and/or young adult outpatients with clinically relevant DD (disorder). The program comprises 8 weekly sessions of 90 min carried Table 1. Elements and techniques of PLAN D

Ρ	Psychoeducation (2 sessions) Aim: information about DD including definition and prevalence; establish an explanatory model (the cogni- tive behavioral model of DD, see Fig. 1), initiate a DD diary Techniques: psychoeducation
L	Lifestyle interventions (1 session) <i>Aim</i> : eliminate maladaptive (safety) behavior and misconceptions about lifestyle behaviors by interactively providing information about lifestyle with a special focus on stress, sleep, substance (ab-)use, social, and physical activity <i>Techniques</i> : psychoeducation, model learning
A	Acceptance and mindfulness training (1 session) Aim: acceptance of DD as well as aversive internal content (e.g., thoughts and feelings) whilst pursuing per- sonal values; learn to be mindful, lessen symptom monitoring, and coping with experiential avoidance Techniques: guided mindfulness training ¹ , relaxation, role plays
N D	New patterns of DD-related cognitions (3 sessions) <i>Aim</i> : identifying and modifying dysfunctional cognitions about DD and demonstrating their impact on emotions and behavior <i>Techniques</i> : cognitive restructuring techniques and evidence gathering

out by 2 therapists. The sessions are accompanied by a "Booklet" with worksheets, such as the DD diary and materials. The sequence of the sessions is structured as follows: beginning with intensive psychoeducation (P) as well as establishing a DD model that can be transferred to every individual case. The diary keeping helps to reveal relevant factors such as situations, emotions, cognitions, and behaviors, which are then discussed in the following lifestyle session (L). The focus of this session lies in correction of false beliefs about DD-related factors, psychoeducation about avoidance behavior, and stress management. In the "A" session, the focus lies in practicing acceptance as a helpful coping strategy for DD-related emotions and training in mindfulness. From this session on, in every following session, mindfulness or meditation will be exercised at the beginning or end of the sessions. The next 3 sessions (N and D) target the reconstruction of DD-specific cognitions. The eighth session is a closing session containing retrospection, summary, feedback, and resource activation in the group. All subjects (or their parents or guardians) had given their written informed consent. The study protocol was approved by the institute's committee on human research.

Study Design

PLAN D was evaluated in a naturalistic setting in the FETZ Bern with data of 2 uncontrolled groups of 4 outpatients. Patients were recruited after the diagnostics if they fulfilled the following inclusion criteria: \geq 13 years and clinically relevant DD disorder according to ICD-10 [25] or DSM-5 [2]. For the recruitment, DD symptoms were assessed in clinical interviews [11, 12]. As an outcome measure, we evaluated the severity and frequency of DD symptoms according to the "Cambridge Depersonalization Scale" (CDS) [26]. The CDS was used as a pre- and postmeasure at baseline before the therapy and 6 months after baseline. The CDSs were handed out in the first session and sent by post 6 months after as a postmeasure. No further clinical evaluation or diagnostic was done at the end of PLAN D as well as 6 months later.

Study Population

The studied sample comprised 8 outpatients (4 males) between 17 and 23 years (M \pm SD: 20.27 \pm 2.01 years). According to ICD-10 [25], all patients fulfilled criteria of a derealization/depersonalization disorder (F48.1), 4 patients additionally had a current depressive episode (F32.1), one had a schizotypal disorder (F21), and one a bipolar disorder (F31.1), and 6 of them further met criteria for a CHR state for psychosis according to the EPA [13]. Three patients with a depressive episode had current antidepressant medication, and 1 patient with a CHR state and schizotypal disorder had an antipsychotic medication. All medication has been established before the start of PLAN D and was stable during the program. Duration of DD was 35.0 months on average (SD: 21.94, Md: 31.0, range: 7-66; 1 patient reported lifetime duration of DD). Seven patients reported enduring DD, and 1 patient reported episodic DD of at least once in a month during a week. Two patients had no treatment history, 5 had psychological outpatient treatment, and one had a psychiatric inpatient treatment before PLAN D.

Results

As shown in Figure 2, at the beginning of PLAN D, all CDS scores were above the cutoff (\geq 65) for clinically relevant DD (M ± SD: 99.14 ± 50.12, Md: 82.0, range: 66–209). At postmeasure, 3 persons (1 male/2 females) showed clinically relevant DD according to the CDS (M ± SD: 74.71 ± 34.99, Md: 59.0, range: 49–149). Although

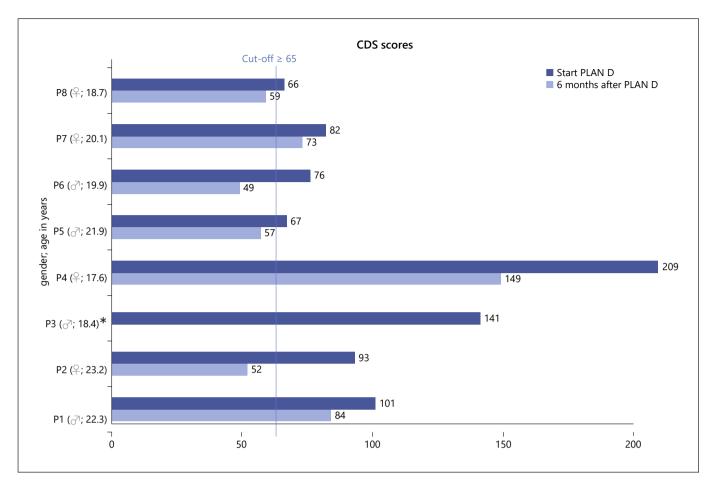


Fig. 2. Pre- and postmeasure scores of the CDS at baseline before the PLAN D group intervention and 6 months after baseline for the sample of 8 participants (P1–P8). P2, 3, and 8 did not continue any therapy after PLAN D; P1 and 4–7 continued further psychological treatment after PLAN D. *Follow-up measurement is missing. CDS, Cambridge Depersonalization Scale.

the effect size of the decrease in DD symptoms was large (Wilcoxon signed rank test: Z = -2.366, p = 0.018, Rosenthal's r = 0.894), according to Figure 2, the mean reduction in the CDS is about 20%, which reflects a minimal improvement in psychopathology. An informal qualitative assessment showed high acceptance and heterogenic, but overall, very good ratings of the different sessions (best rating for sessions "N and D"; lowest rating for session "L," see Table 1).

Discussion

According to an existing CBT-based treatment protocol for DD disorder [6], we conceptualized and implemented a group intervention called PLAN D to face the limited availability of disorder-specific interventions for

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DD. In our first data, we found a reduction in the overall CDS score 6 months after the start of PLAN D with only 3 persons still being over the clinical cutoff. Although we found a statistically strong effect of the intervention (Rosenthal's r > 0.8), due to small sample size, lack of a control group, and no immediate posttreatment measurement (approximately 6 months after baseline), this finding should be interpreted with some caution as the reduction in psychopathology of 20% reflects a minimal improvement. According to our informal quality assessment, the acceptance of the group was very good. Our experiences showed that in the recruiting process some patients have been very open and were immediately dedicated to take part in PLAN D while others were more skeptical and initially reluctant. Consequently, we were confronted with the challenge to carefully explore their reservations and to motivate them to confront,

rather than avoid, specific (social) situations. Therefore, in some cases taking part in PLAN D, group sessions posed a first step in their therapeutic process helping to overcome their (social) withdrawal. As from the first group session on and in the interaction between participants, respectively, we observed a normalizing effect concerning DD experiences. Besides the good compliance, patients showed high interest and involvement especially in the psychoeducation sessions. During these sessions, we carefully elaborate with participants to learn DD is a possible psychological reaction to stress (according to the model in Fig. 1) as well as also to address the distress about DD as a potential psychosis risk symptom. In the feedback session, all participating patients reported to meet other DD-affected persons as very helpful as well as get and give each other support while receiving professional DD-specific intervention. Even though the qualitative ratings of each individual session were relatively high, sessions 5-7 (N and D sessions, see Table 1) were rated highest by participants and in our view highlight the importance of cognitive techniques to cope with DD by overcoming dysfunctional cognitions. In the "lifestyle intervention" session, we discuss existing concerns about lifestyle or behavior and their impact on DD such as sleep, nutrition, or substance use more general and clear up with possible maladaptive beliefs. In this session, participants already demonstrated good knowledge of the topic by sharing their personal experiences with others, which could be the reason for the lowest rating. Besides this, participants found the experience to be positive throughout the course of therapy. However, we are presenting very preliminary data from a naturalistic setting, and some limitations should be noted, such as the lack of a control group and the small sample size. Furthermore, clinical evaluation immediately after completion of the group program PLAN D is not included in the current pilot study, and future studies should endeavor to include such measures. Evaluation of the program as a randomized controlled trial with appropriate control groups would further help to disentangle therapy effects from other possible confounding variables (e.g., other therapies or natural symptomatic reduction) and should be considered in future studies. Nonetheless, PLAN D presents a promising option as an adjunct to existing treatments given in individual settings, as it consists of well-established CBT (based) techniques that can be offered in a time-limited group setting. Furthermore, if our initial findings would be replicated or even improved, it would be considered a cost-effective option

for psychological treatment for DD. Future research should replicate these promising results by addressing the named limitations and continually improve interventions for this highly distressing condition. Early diagnosis and psychoeducation help alleviate DD-associated distress and may promote recovery or prevent a long-term DD history and, consequently, a possible psychotic development. To that end, PLAN D represents a novel, brief intervention and a promising opportunity to positively influence the prognosis of patients suffering from DD.

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Statement of Ethics

All procedures contributing to this work comply with the ethical standards of the relevant national and institutional committees on human experimentation and with the Helsinki Declaration of 1975, as revised in 2013. The Human Research Ethics Committee of the Canton Bern approved the study (ID PB_2016-01991). All participants gave written informed consent, and in minors, parental written informed consent was provided.

Conflict of Interest Statement

All authors declare no conflicts of interest regarding the topic of this article.

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Author Contributions

Rahel Flückiger contributed to conceptualization, project administration, investigation and implementation of the group therapy, methodology, data analysis, and visualization. Stefanie J. Schmidt contributed to conceptualization and supervision. Chantal Michel contributed to project administration and visualization. Jochen Kindler contributed to project administration and supervision. Michael Kaess contributed to project administration, methodology, and supervision. All authors contributed to the preparation of the manuscript.

Data Availability Statement

All data generated or analysed during this study are included in this published article. The qualitative data generated and analysed during the current study are available from the corresponding author on reasonable request.

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