Baseline depressive symptoms, personal control, and concern moderate the effects of preoperative psychological interventions: the randomized controlled PSY-HEART trial

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Abstract This study examined whether baseline (3–14 days pre-surgery) levels of (i) depressive or (ii) anxiety symptoms and (iii) illness beliefs moderate the effects of additional preoperative interventions before coronary artery bypass graft surgery on (i) depressive or (ii) anxiety symptoms and (iii) illness beliefs 1 day before surgery, 1 week and 6 months after surgery. In the PSY-HEART trial, 115 patients were assessed. They were randomized into one of three groups: 1. receiving standard medical care only (SMC), additional psychological interventions: 2. aiming to optimize patients’ expectations (EXPECT), or 3. focusing on emotional support. Patients with a higher baseline level of depressive symptoms receiving a preoperative psychological intervention indicated lower depressive symptoms 6 months after surgery compared to SMC. EXPECT increased personal control and concern levels in patients with low baseline personal control/concern 1 day before surgery. Preoperative psychological interventions can improve psychological outcomes in heart surgery patients. Baseline status may moderate these effects. The study has been approved by the medical ethics committee of the Philipps University of Marburg and has been pre-registered at www.clinicaltrials.gov (NCT01407055) on August 1, 2011.

Keywords Expectation · Depression · Personal control · Concern · Placebo · Preoperative psychological intervention · CABG · PSY-HEART

Introduction

Cardiovascular diseases are associated with a high risk of mortality, disability, a decreased quality of life, and increased costs for the healthcare system (Murray & Lopez, 2013; Virani et al., 2020). Coronary artery bypass grafting (CABG) is an established treatment option for patients with advanced coronary artery disease that has been thoroughly studied over several decades (Hawkes et al., 2006). However, it is still unknown why a substantial number of patients faces problems in the recovery process and does not benefit as much from the surgery as surgeons would predict (Blumenthal et al., 2003; Burg et al., 2003; Hawkes & Mortensen, 2006; Hawkes et al., 2006; Salzman, Euteneuer, et al., 2020; Salzmann, Salzmann-Djufri, et al., 2020). Patients’ recovery after surgery is not explained by medical factors alone; recovery seems to be a multidimensional phenomenon in which physical, psychological, and social factors play important roles as well (Auer et al., 2016; Hawkes et al., 2006; Sadeghi et al., 2017). Growing evidence suggests the importance of psychological preparation for improving post-surgery physical outcomes and psychological outcomes (i.e., quality of life, disability, pain, morbidity, length of hospital...
Changing illness beliefs has enhanced health outcomes in several studies with cardiac patients (Davies et al., 2008; Juergens et al., 2010; Keogh et al., 2011; Petrie et al., 2002, 2012). Research may benefit from focusing more on patient beliefs and expectations, especially about personal control, in exploring the recovery process. Higher scores of preoperative perceived control have been shown to predict postoperative quality of life and lower levels of depression in CABG patients (Kidd et al., 2016). Nonetheless, little is known about the question of which preoperative psychological intervention can influence what kind of illness beliefs and who will benefit from such an intervention specifically (Kidd et al., 2016).

Besides the cognitive responses to a perceived health threat, the CSM highlights the importance of emotional factors in coping with a disease, e.g., illness beliefs such as concern or emotions (Leventhal & Cameron, 1987; Leventhal et al., 1992). Other emotional factors such as depression or anxiety are also highly relevant in cardiac surgery patients: Depression is highly prevalent in patients undergoing CABG (Blumenthal et al., 2003; Head et al., 2013; Poole et al., 2014, 2017; Tully et al., 2009; Young et al., 2019). 20–40% of CABG surgery patients are affected by depression (Blumenthal et al., 2003; Connerney et al., 2001; Young et al., 2019). Depressed patients undergoing CABG surgery report a lower health-related quality of life, have a higher postoperative rate of depression, a higher risk of rehospitalization and death, and stay longer in hospital after the surgical procedure independent from medical factors compared to non-depressed patients (Auer et al., 2017; Blumenthal et al., 2003; Connerney et al., 2001; Contrada et al., 2004; Mallik et al., 2005; Morone et al., 2010; Oxlad et al., 2006; Rollman et al., 2009; Rumsfeld et al., 2004; Timberlake et al., 1997). Dunkel et al. (2011) suggest that patients with higher depression levels might benefit most from additional psychological intervention. Similarly, preoperative anxiety seems to be associated with a negative postoperative course, yet fewer is known about this relationship, especially in CABG patients (Arthur et al., 2000; Heilmann et al., 2016; Lamarche et al., 1998; Székely et al., 2007). Since preoperative anxiety and depressive symptoms seem to be important predictors of the postoperative recovery process, psychological interventions targeting these symptoms could improve postoperative physical and psychological outcomes (i.e., such as depression or anxiety). Heilmann et al. (2016) reported that a preoperative intervention reduced the preoperative and postoperative state anxiety compared to a control group. However, it is mainly unknown how specific interventions can be tailored to the individual needs of CABG patients to reduce patients anxiety and depression levels.
The PSY-HEART trial indicated that receiving a preoperative psychological intervention aiming to optimize patients’ expectations (EXPECT) led to reduced illness-related disability as the primary outcome (Rief et al., 2017). Several positive effects on secondary outcomes were also found: For instance, the EXPECT intervention indicated increased physical and mental quality of life 6 months after CABG surgery and fewer days of hospitalization in comparison to standard medical care (SMC) only (for further information, see Auer et al., 2017; Rief et al., 2017). For depression (another secondary outcome), a non-significant trend was found in favor of EXPECT and SUPPORT (an attention control group) receiving the same amount of time and attention by the psychologist but without working specifically on expectations) compared to SMC only, when assessing baseline and follow-up scores 6 months after surgery. However, it is still unclear who benefitted the most from the preoperative psychological interventions in the PSY-HEART trial and when these interventions seemed to work regarding patients’ depressive and anxiety symptoms as well as illness-beliefs. A meta-analysis of Bower et al. (2013) indicated that patients who had higher levels of depression at baseline showed greater treatment effects than patients with lower levels of depression at baseline. For anxiety, baseline anxiety was the most frequently examined moderator of the effectiveness of psychological and psychoeducational interventions for anxiety in a meta-analysis (Moreno-Peral et al., 2020). Since patients’ depressive and anxiety symptoms and patients’ illness beliefs (i.e., perceived personal control or concern) are considered important outcome predictors, especially in heart surgery patients (Salzmann, Euteneuer, et al., 2020; Salzmann, Salzmann-Djufri, et al., 2020), a more thorough understanding of the (psychological) intervention effects over time regarding these psychological factors is crucial. To better understand how, when, and for whom the preoperative psychological interventions (EXPECT: optimizing expectation group; SUPPORT: emotional support/attention control group) seemed to improve depressive and anxiety symptoms, and illness beliefs in the PSY-HEART trial (Rief et al., 2017), this secondary exploratory analysis examined whether (i) baseline scores of depressive symptoms moderated the effects of the preoperative psychological interventions on depressive symptoms in heart surgery patients 1 day before surgery, 4 to 6 days after surgery and 6 months after surgery, whether (ii) baseline anxiety symptom scores moderated the effects of the preoperative psychological interventions on anxiety symptoms in heart surgery patients 1 day before surgery, 4 to 6 days after surgery and 6 months after surgery, and whether (iii) baseline scores of the illness beliefs (i.e. perceived personal control or concern) moderated the effects of the preoperative psychological interventions on illness beliefs (i.e. perceived personal control or concern) in heart surgery patients 1 day before surgery, 4 to 6 days after surgery and 6 months after surgery.

Methods

Study design

The study is part of the randomized controlled clinical PSY-HEART trial (Rief et al., 2017) (see Laferton et al., 2013, for the study protocol). The PSY-HEART trial was approved by the medical ethics committee of the Philipps University of Marburg and was pre-registered at ClinicalTrials.gov (Identifier: NCT01407055). It examined the effects of preoperative psychological interventions on postoperative physical and psychological outcomes in heart surgery patients (CABG or CABG plus heart valve surgery). Participants were randomly assigned to one of three groups: (1) receiving the standard medical treatment (Standard Medical Care—SMC); (2) receiving SMC and additional psychological treatment focusing on optimizing patients’ expectations (EXPECT); (3) receiving SMC and additional psychological treatment providing attention and emotional support by a psychologist (SUPPORT). Patients were assessed at four measurement time points: 3–14 days pre-surgery (T0, baseline), 1 day pre-surgery [after the psychological intervention, but before surgery (T1)], 6–8 days (“1 week”) post-surgery (T2), approximately 6 months post-surgery (T3, follow-up).

The data collection took place from April 2011 to May 2015 in the Department of Cardiovascular Surgery, Philipps University of Marburg, Germany.

Participants

Participants have been recruited from the waiting list of the Heart Surgery center. If a CABG surgery (with or without heart valve surgery) was planned, the patients were contacted before hospital admission. Interested patients were invited to a first appointment. Thereby they were informed and gave written informed consent.

Inclusion criteria were age between 18 and 80 years, speaking and understanding German fluently, being able to give informed consent. Exclusion criteria were a serious comorbid psychiatric or physical (non-cardiac) condition (e.g. acute psychosis, dementia) that hampers the participation at baseline or will do most likely within 6 months until study completion at follow-up. Further exclusion criteria were previous cardiac surgeries and participation in a different research program.

In Fig. 1, the CONSORT flow-chart shows the recruitment and the count of participated patients for each measurement time point (Rief et al., 2017).
Enrollment

124 Randomized

Allocation

44 Standard Medical Care (SMC)
39 SMC + Supportive Intervention (SUPPORT)
39 SMC + Expectation Manipulation Intervention (EXPECT)

Baseline

Lost to baseline (T0) n = 0

Post psychological intervention

Lost to post-intervention (T1) n = 0

Post-surgery

Lost to post-surgery (T2) n = 1
Deceased after CABG (n = 1)

6 months after surgery

Lost to follow-up (T3) n = 1
Deceased after CABG (n = 1)

Analysis

Numbers analysed (after exclusion of outliers and consideration of missings): 41
Available baseline values:
HADS: Depression, Anxiety: n = 39
B-IPQ: Consequences, Identity, Concern, Emotional response: n = 40
B-IPQ: Timeline, Treatment Control: n = 38
B-IPQ: Personal Control, Understanding: n = 39

Numbers analysed (after exclusion of outliers and consideration of missings): 37
Available baseline values:
HADS: Depression: n = 32
HADS: Anxiety: n = 33
B-IPQ: Consequences, Timeline, Treatment Control, Identity, Concern, Emotional response: n = 37
B-IPQ: Personal Control, Understanding: n = 36

Numbers analysed (after exclusion of outliers and consideration of missings): 37
Available baseline values:
HADS: Depression, Anxiety: n = 36
B-IPQ: Consequences: n = 36
B-IPQ: Timeline: n = 33
B-IPQ: Personal Control, Identity, Concern, Understanding, Emotional response: n = 37
B-IPQ: Treatment Control: n = 35

249 Assessed for eligibility

125 Excluded
Not meeting inclusion criteria (n = 24)
Declined to participate due to lack of interest or difficulties to travel to study appointments (n = 72)
Deceased (n = 2)
Other reasons (n = 27)
The calculated sample size for the primary analysis was $N = 180$ (time by group interaction, $f = 0.2$, $\alpha = 0.05$, $1 - \beta = 0.8$) (Rief et al., 2017). Recruitment goal was not achieved and the trial ended with $N = 124$ participants, this reflects a post hoc power of $85\%$ ($f > 0.15/d > 0.30/\text{number needed to treat} < 6$, $\alpha = 0.05$). After excluding 9 patients (for details, please see statistics), a sample size of $N = 115$ resulted. A post hoc power analysis with a sample size of $N = 115$ resulted in a post hoc power of $81\%$ ($f > 0.15/d > 0.30/\text{number needed to treat} < 6$, $\alpha = 0.05$). Since the Helsinki recommendation implies that trials investigating innovative interventions should not be oversized, this was considered adequate.

**Procedure**

After having been contacted and consenting to participate, patients were informed about the study both orally and in writing at least three to 14 days before surgery. Before participation patients had to sign the informed consent.

At baseline measurement, psychologists performed the Structured Clinical Interview for DSM-IV (First et al., 1996) to screen for psychiatric comorbidities. Medical information as the EuroScore were taken out of the patient’s files. Afterwards, patients completed questionnaires, and blood samples were taken. More information about the questionnaires can be found in the Study Protocol (Laferton et al., 2013; Rief et al., 2017). For more information about the blood samples, please refer to Salzmann, Euteneuer, et al. (2020).

Patients were then randomized into one of the three intervention groups following a permuted block randomization in WINPEPI (block size of 9) (Abramson, 2011). Strata were age ($\leq 65$ years and more than 65 years) and the New Heart Association class (NYHA, I/II and III/IV). The psychologists opened one closed envelope for each patient when a patient had finished the baseline assessments to avoid bias. Patients and the study team were aware of the treatment allocation, while surgeons and all other routine care personal were blinded. Generally, patients randomized to one of the two intervention groups had their first session with the psychologist subsequently.

**Psychological interventions**

Patients were randomized into one of three groups: either the SMC alone (control group) or one of two additional preoperative psychological interventions [EXPECT (IG) or SUPPORT (attention control group)]. Both psychological interventions lasted at least around 140 min (one individual session á 50 min, two phone calls á 20 min and another individual session á 50 min). They both were conducted by three clinical psychologists (2 male, 1 female) with advanced cognitive behavioral therapy skills. The manual-based interventions had a high treatment fidelity (Laferton et al., 2016). EXPECT focused on positive and realistic expectations regarding the patients’ disease, surgery-benefits, and recovery process. The EXPECT intervention is based on the Common Sense Model described in the introduction, with expectations as an inherent component of illness beliefs (Cameron & Leventhal, 2003). The EXPECT intervention focuses on several expectation facets. These expectations are summarized in the integrative model of expectations in patients undergoing medical treatment (Laferton et al., 2017). In particular, the intervention aimed to encourage the patients in developing a realistic understanding of disease and positive outcome expectations (separated in behavior-related expectations as self-efficacy and behavior outcome expectation and treatment-related expectations as structural and process expectations). Patients received psychoeducation about the CABG-procedure (structural expectations), planned when they will be able to return to which positive activities (process expectations), how they can influence controllable risk factors (behavior outcome expectation), and cope with handling side effects of the surgery (self-efficacy). A better understanding of patients’ disease was achieved, and false assumptions were corrected. In the end, patients imagined a positive scene after long-term recovery to strengthen their outcome expectations. Detailed information about the intervention, work sheets and examples of patients’ thoughts can be found elsewhere (Salzmann et al., 2018).

By comparing the intervention group EXPECT and the SMC control group, it would still be unknown if the outcome effects are due to the specific content of the EXPECT intervention (working on patients’ expectations) or are the result of the unspecific intervention ‘ingredients’ such as establishing a therapeutic relationship, providing emotional support, and paying attention to the patient. Therefore, an active control group controlling for these unspecific effects (attention control group) was included. The recommended procedure for trials of psychological interventions includes realizing an attention control group in which patients receive the same amount of time and attention as the patients in the intervention group (Guidi et al., 2018). For this reason, the attention control group SUPPORT was implemented in the study. In the SUPPORT intervention, patients received the same amount of time, attention, and emotional support from the therapists as did the patients in the EXPECT group, but without specifically targeting expectations. Since attention and
emotional support can lead to positive outcome effects by itself (Guidi et al., 2018), and the primary analysis of the PSY-HEART trial indicated some beneficial effects of the SUPPORT intervention compared to the SMC group (Rief et al., 2017), the attention control group SUPPORT could also be seen as another psychological intervention group. In sum, both preoperative psychological intervention groups provided the patients with emotional support, while only the EXPECT intervention specifically targeted patients’ expectations and illness beliefs which have been described as relevant mediators in the CSM.

**Outcome criteria**

The primary outcome of the main trial was illness-related disability measured with an adapted version of the Pain Disability index (PDI) (Tait et al., 1990). Secondary outcomes were quality of life (Short Form 12, SF-12) (Ware et al., 1996), anxiety and depressive symptoms (Hospital Anxiety and Depression Scale, HADS) (Zigmond & Snaith, 1983), illness beliefs (Brief Illness Perception Questionnaire, B-IPQ) (Broadbent et al., 2006), subjective ability to work and the increase of metabolic equivalents of physical activity after surgery (International Physical Activity Questionnaire, IPAQ) (Craig et al., 2003). Results of these and other outcomes are published elsewhere (Rief et al., 2017). As this study examines further results and moderators for change in depression, anxiety and illness perceptions, these variables will be described in more detail.

The HADS examines anxiety and depression in patients with (psycho)somatic conditions. Each subscale has seven items that are scored at a 4-Item-Likert-Scale (0 to 3). Each scale was evaluated on its own and a sum score of the general psychological distress was analyzed (Zigmond & Snaith, 1983). Higher values mean a higher score of anxiety, depression or general psychological distress.

The B-IPQ surveys the cognitive and emotional representations of illness (Broadbent et al., 2006). It is composed of 8 closed items (range from 0 to 10). One additional question asks for possible reasons for the disease. Each item is evaluated individually. Higher scores reflect an increase of the respective dimension. Item 1–5 measure cognitive illness representations (consequences, timeline, personal control, treatment control, and identity). Item 6 and 8 quantify emotional representations (concern and emotions). Item 7 assesses illness comprehensibility. Item 9 is an open question (three most important causal factors in their illness).

**Statistics**

As can be seen in Fig. 1, 9 patients were excluded from the ITT-sample for statistical analysis (violation of design requirements: 1; did not require CABG surgery: 1; resigned from study before baseline assessment: 1; multivariate outliers: 6; for details, please see Rief et al., 2017). Box-plots were screened to identify outliers (> 3 interquartile ranges). Only one outlier was identified for a T1-score of B-IPQ-Understanding, it was excluded before analysis. The following baseline scores were available: HADS – Depression: \( n = 107 \), HADS – Anxiety: \( n = 108 \), BIPQ – Identity, concern and emotional response: \( n = 114 \), B-IPQ – Consequences: \( n = 113 \), B-IPQ – Personal control and understanding: \( n = 112 \), B-IPQ – Treatment control: \( n = 110 \), B-IPQ – Timeline: \( n = 108 \).

To examine the psychological interventions effects as well as the potential moderation effects of baseline anxiety, depressive symptoms and illness beliefs on the intervention effects, multilevel models were used. Each outcome variable was explored in a separate model. Fixed effects were calculated for group (EXPECT, SMC, SUPPORT), time (1-day pre-surgery, 1-week post-surgery, 6 months post-surgery), baseline scores of the respective outcomes, group*time*baseline-scores and all other lower interaction terms (especially group*time-interactions). The analyses were adjusted for baseline differences, as the baseline scores were considered as a covariate.

In each model, a random intercept was included to allow for interindividual effects. First, two-way-interactions were examined (group*time) and in case of significant two-way-interactions post-hoc simple slope analyses were evaluated (\( \alpha \leq 0.05 \)). Second, three-way-interactions were examined (group*time*baseline score of outcome variable). If significant or three-way-interactions were found, for continuous moderators simple-slope follow-up analyses were evaluated (Preacher et al., 2006). By doing so the significance of the intervention effects for conditional values of the moderator [for low (− 1 SD), average (mean) and high (+ 1 SD) baseline scores] were calculated.

Due to the small sample size missing data were imputed using restricted maximum likelihood (REML) methods. Covariance structures were theoretically assumed in a first step and then empirically verified by goodness of fit using the Akaike information criterion (AIC) (Hox et al., 2018). For all statistical analyses SPSS 26 was used (IBM Corp., 2019). Due to the explored character of the article, we did not correct for multiple testing.

**Results**

**Baseline characteristics**

Table 1 shows the baseline characteristics as published in Rief et al. (2017), supplemented with the baseline levels of
the outcome criteria of the analyses in this article. At baseline the groups did not differ significantly.

### Intervention effects over time

Two-way interactions were assessed to test for intervention effects over time (Table 2). Significant group by time-interactions were indicated for consequences ($p = 0.028$), personal control ($p = 0.028$), identity ($p = 0.044$) and concern ($p = 0.030$). No significant two-way interactions were observed for depression ($p = 0.371$), anxiety ($p = 0.583$), HADS sum score ($p = 0.800$), timeline ($p = 0.588$), treatment control ($p = 0.165$), understanding ($p = 0.150$) and emotional response ($p = 0.335$).

The significant group by time-interaction of perceived personal control implied an intervention effect over time ($p = 0.028$, Table 2). Post-hoc simple slope analyses indicated that patients receiving EXPECT or SUPPORT showed significant higher personal control values 1 day before surgery compared to SMC (EXPECT vs. SMC: $p < 0.001$, SUPPORT vs. SMC: $p = 0.045$). There were no statistically significant differences at other measurement timepoints for personal control ($p \geq 0.127$).

The significant group by time-interaction of perceived consequences ($p = 0.028$, Table 2), perceived identity ($p = 0.044$, Table 2) and perceived

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**Table 1** Demographical, medical and psychological characteristics at baseline of patients receiving Standard Medical Care (SMC), Supportive Intervention (SUPPORT) or Expectation Manipulation Intervention (EXPECT) (Rief et al., 2017)

| Note | SMC | SUPPORT | EXPECT |
|------|-----|---------|--------|
| **SMC** | **SUPPORT** | **EXPECT** |
| **SUMMARY** | | | |
| Age in years, $M (SD)$ | 67.07 (8.9) | 64.62 (8.1) | 65.76 (7.8) |
| Sex, male, n (%) | 36 (87.8) | 30 (81) | 32 (86.5) |
| Education, high school, n (%) | 7 (17.1) | 10 (27) | 10 (27) |
| Marital status, married, n (%) | 33 (80.5) | 34 (91.9) | 31 (83.8) |
| BMI, $M (SD)$ (MD = 3) | 29.67 (5.2) | 29.5 (6.6) | 29.03 (5.01) |
| Smoking status, n (%) | 6 (14.6) | 2 (5.4) | 6 (16.2) |
| EuroSCORE II, $M (SD)$ (MD = 11) | 1.53 (0.8) | 1.47 (0.8) | 1.25 (0.8) |
| NYHA, n (%) (MD = 10) | | | |
| I | 1 (2.4) | 1 (2.7) | 0 (0) |
| II | 9 (22.0) | 11 (29.7) | 12 (32.4) |
| III | 28 (68.3) | 20 (54.1) | 17 (45.9) |
| IV | 1 (2.4) | 2 (5.4) | 3 (8.1) |
| LVEF n, (%) (MD = 9) | | | |
| $\geq 50$ | 23 (48.8) | 19 (51.4) | 30 (78.4) |
| 49–30 | 13 (31.7) | 13 (35.1) | 4 (10.8) |
| < 30 | 2 (4.9) | 2 (5.4) | 0 (0) |
| Previous myocardial infarction, n (%) (MD = 5) | 9 (23.1) | 6 (17.1) | 6 (16.7) |
| Combined surgery, n (%) | 6 (14.6) | 6 (16.2) | 3 (8.1) |
| Anxiety, $M (SD)$ (MD = 7) | 4.03 (3.0) | 4.55 (3.2) | 5.17 (4.0) |
| Depression, $M (SD)$ (MD = 8) | 4.59 (3.1) | 4.0 (3.1) | 5.11 (4.0) |
| Consequences, $M (SD)$ (MD = 2) | 5.00 (2.96) | 5.00 (3.38) | 5.33 (3.03) |
| Timeline, $M (SD)$ (MD = 7) | 3.03 (3.04) | 3.95 (3.26) | 1.39 (1.35) |
| Personal Control, $M (SD)$ (MD = 3) | 4.64 (2.57) | 4.31 (3.19) | 3.84 (2.88) |
| Treatment Control, $M (SD)$ (MD = 5) | 8.84 (1.42) | 8.65 (1.51) | 9.31 (0.90) |
| Identity, $M (SD)$ (MD = 1) | 4.33 (2.71) | 4.81 (2.94) | 4.86 (2.64) |
| Concern, $M (SD)$ (MD = 1) | 5.95 (3.06) | 6.46 (3.23) | 5.84 (3.71) |
| Understanding, $M (SD)$ (MD = 4) | 6.72 (2.60) | 7.00 (3.31) | 7.30 (2.73) |
| Emotional response, $M (SD)$ (MD = 1) | 4.43 (2.63) | 4.41 (3.29) | 4.70 (3.27) |

*Notes. SMC = Standard Medical Care. SUPPORT = Supportive Intervention. EXPECT = Expectation Manipulation Intervention. MD = missing data. Body Mass Index (BMI). EuroSCORE (European System for Cardiac Operative Risk Evaluation). NYHA (New York Heart Association functional classification). LVEF (Left ventricular ejection fraction). Anxiety and Depression (Hamilton Anxiety and Depression Scale; HADS) range = 0–21. Disability (Pain Disability Index; PDI) range = 0–70. Mental quality of Life (Mental component of the Short-Form Health Survey; SF-12). Physical quality of Life (Physical component of the Short-Form Health Survey; SF-12). Physical activity (International physical activity questionnaire (IPAQ) weighted estimate of total physical activity per week. Cardiac Anxiety (Cardiac Anxiety Questionnaire) range = 0–4. Consequences, Timeline, Personal Control, Treatment Control, Identity, Concern, Understanding, Emotional response (Brief Illness Perception Questionnaire, B-IPQ) range 0–10*
Table 2 Outcome measures at baseline, 1 day before surgery, 1 week after surgery and 6 months after surgery of patients receiving standard medical care (SMC), supportive intervention (SUPPORT) or expectation manipulation intervention (EXPECT) (observed measures) and test statistics for two-way interactions between intervention group and assessment time and also three-way-interactions between intervention group, assessment time and baseline score

|                | SMC M (SD) | SUPPORT M (SD) | EXPECT M (SD) | Test statistic (F scores of two-way interaction terms) | Test statistic (F scores of three-way interaction terms) |
|----------------|------------|----------------|---------------|---------------------------------------------------------|---------------------------------------------------------|
| **HADS**       |            |                |               |                                                         |                                                         |
| Depressive symptoms |          |                |               |                                                         |                                                         |
| Baseline       | 4.54       | 4.54           | 4.54          |                                                         |                                                         |
| 1 day before surgery | 5.00 (0.79) | 4.21 (0.80)    | 4.77 (0.81)   | \(F(4, 163.827) = 1.075, p = .371\)                      |                                                         |
| 1 week after surgery | 5.40 (0.79) | 3.64 (0.83)    | 4.70 (0.79)   | \(F(4, 162.184) = 2.569, p = .040\)                      |                                                         |
| 6 months after surgery | 3.65 (0.78) | 2.12 (0.83)    | 2.48 (0.81)   |                                                         |                                                         |
| Anxiety        |            |                |               |                                                         |                                                         |
| Baseline       | 4.55       | 4.55           | 4.55          |                                                         |                                                         |
| 1 day before surgery | 5.03 (0.44) | 4.72 (0.46)    | 4.98 (0.47)   | \(F(4, 190.192) = 0.714, p = .583\)                      | \(F(4, 181.434) = 1.547, p = .191\)                     |
| 1 week after surgery | 4.18 (0.45) | 3.06 (0.48)    | 3.53 (0.45)   |                                                         |                                                         |
| 6 months after surgery | 3.36 (0.43) | 2.57 (0.47)    | 3.23 (0.45)   |                                                         |                                                         |
| Total score    |            |                |               |                                                         |                                                         |
| Baseline       | 8.94       | 8.94           | 8.94          |                                                         |                                                         |
| 1 day before surgery | 10.01 (0.67) | 8.68 (0.74)    | 9.34 (0.73)   | \(F(4, 133.152) = 0.411, p = .800\)                      | \(F(4, 136.404) = 1.178, p = .323\)                     |
| 1 week after surgery | 9.68 (0.74) | 6.35 (0.84)    | 8.02 (0.76)   |                                                         |                                                         |
| 6 months after surgery | 7.01 (0.68) | 4.42 (0.81)    | 5.20 (0.74)   |                                                         |                                                         |
| **B-IPQ**      |            |                |               |                                                         |                                                         |
| Consequences   |            |                |               |                                                         |                                                         |
| Baseline       | 5.09       | 5.09           | 5.09          |                                                         |                                                         |
| 1 day before surgery | 5.22 (0.51) | 4.94 (0.52)    | 5.64 (0.53)   | \(F(4, 196.677) = 2.791, p = .028\)                      | \(F(4, 194.173) = 2.215, p = .069\)                     |
| 1 week after surgery | 6.97 (0.52) | 6.36 (0.53)    | 6.50 (0.53)   |                                                         |                                                         |
| 6 months after surgery | 3.28 (0.51) | 3.10 (0.53)    | 2.82 (0.53)   |                                                         |                                                         |
| Timeline       |            |                |               |                                                         |                                                         |
| Baseline       | 2.65       | 2.65           | 2.65          |                                                         |                                                         |
| 1 day before surgery | 2.86 (0.29) | 2.44 (0.32)    | 3.11 (0.43)   | \(F(4, 121.741) = 0.708, p = .588\)                      | \(F(4, 121.999) = 0.541, p = .706\)                     |
| 1 week after surgery | 4.82 (0.39) | 4.61 (0.43)    | 3.73 (0.56)   |                                                         |                                                         |
| 6 months after surgery | 4.64 (0.55) | 4.34 (0.62)    | 6.96 (0.94)   |                                                         |                                                         |
| Personal control |            |                |               |                                                         |                                                         |
| Baseline       | 4.41       | 4.41           | 4.41          |                                                         |                                                         |
| 1 day before surgery | 3.83 (2.09) | 4.96 (2.09)    | 5.84 (2.09)   | \(F(4, 127.549) = 2.805, p = .028\)                      | \(F(4, 127.373) = 2.511, p = .045\)                     |
| 1 week after surgery | 5.45 (2.09) | 6.05 (2.10)    | 6.22 (2.09)   |                                                         |                                                         |
| 6 months after surgery | 4.23 (2.10) | 5.02 (2.10)    | 4.71 (2.11)   |                                                         |                                                         |
| Treatment control |            |                |               |                                                         |                                                         |
| Baseline       | 8.98       | 8.98           | 8.98          |                                                         |                                                         |
| 1 day before surgery | 8.83 (0.24) | 8.58 (0.25)    | 8.86 (0.32)   | \(F(4, 117.891) = 1.654, p = .165\)                      | \(F(4, 117.259) = 1.308, p = .271\)                     |
| 1 week after surgery | 8.02 (0.27) | 8.69 (0.28)    | 8.39 (0.29)   |                                                         |                                                         |
| 6 months after surgery | 7.77 (0.38) | 8.77 (0.42)    | 7.37 (0.45)   |                                                         |                                                         |
| Identity       |            |                |               |                                                         |                                                         |
| Baseline       | 4.60       | 4.60           | 4.60          |                                                         |                                                         |
| 1 day before surgery | 3.97 (3.70) | 4.21 (3.70)    | 4.58 (3.70)   | \(F(4, 152.844) = 2.519, p = .044\)                      | \(F(4, 152.141) = 2.216, p = .070\)                     |
| 1 week after surgery | 5.56 (3.71) | 4.44 (3.72)    | 5.53 (3.71)   |                                                         |                                                         |
| 6 months after surgery | 3.22 (3.71) | 2.67 (3.71)    | 2.27 (3.71)   |                                                         |                                                         |
concern ($p = 0.030$, Table 2) implied intervention effects over time. However, no significant group differences were indicated in post-hoc simple slope analyses for all three outcomes (consequences: $p \geq 0.192$, identity: $p \geq 0.060$, concern: $p \geq 0.101$). The results of the significant group by time-interactions are diagrammed in the Supplementary Fig. 5.

Moderation effects of baseline scores

Three-way interactions were assessed to test for moderating effects of the baseline score regarding the intervention effects (Table 2). Significant group by time by baseline scores of the outcome-interactions were indicated for depressive symptoms ($p = 0.040$), personal control ($p = 0.045$), and concern ($p = 0.046$). No significant three-way interactions were observed for anxiety ($p = 0.191$), HADS sum score ($p = 0.323$), consequences ($p = 0.069$), timeline ($p = 0.706$), treatment control ($p = 0.271$), identity ($p = 0.070$), understanding ($p = 0.258$) and emotional response ($p = 0.498$). The model fit statistics are accessible in Table 1 of the Supplementary material. The full Table 2 (continued)

|                  | SMC       | SUPPORT   | EXPECT    | Test statistic (F scores of two-way interaction terms) | Test statistic (F scores of three-way interaction terms) |
|------------------|-----------|-----------|-----------|--------------------------------------------------------|--------------------------------------------------------|
| Concern          |           |           |           | $F(4, 152.853) = 2.763, p = 0.030$                       | $F(4, 150.618) = 2.492, p = 0.046$                      |
| Baseline         | 6.15      | 6.15      | 6.15      |                                                        |                                                        |
| 1 day before surgery | 5.93 (1.01) | 5.71 (1.02) | 6.06 (1.01) |                                                        |                                                        |
| 1 week after surgery | 5.74 (1.04) | 4.62 (1.05) | 4.75 (1.03) |                                                        |                                                        |
| 6 months after surgery | 3.40 (1.01) | 3.23 (1.03) | 3.15 (1.02) |                                                        |                                                        |
| Understanding    |           |           |           | $F(4, 182.219) = 1.706, p = 0.150$                       | $F(4, 181.222) = 1.337, p = 0.258$                      |
| Baseline         | 7.10      | 7.10      | 7.10      |                                                        |                                                        |
| 1 day before surgery | 7.11 (0.54) | 7.74 (0.56) | 8.25 (0.55) |                                                        |                                                        |
| 1 week after surgery | 7.31 (0.54) | 7.60 (0.56) | 7.55 (0.53) |                                                        |                                                        |
| 6 months after surgery | 7.06 (0.54) | 7.99 (0.56) | 7.20 (0.54) |                                                        |                                                        |
| Emotional response|           |           |           | $F(4, 165.681) = 1.149, p = 0.335$                       | $F(4, 162.035) = 0.845, p = 0.498$                      |
| Baseline         | 4.56      | 4.56      | 4.56      |                                                        |                                                        |
| 1 day before surgery | 4.42 (1.19) | 3.99 (1.20) | 4.64 (1.20) |                                                        |                                                        |
| 1 week after surgery | 4.55 (1.22) | 4.44 (1.23) | 4.20 (1.22) |                                                        |                                                        |
| 6 months after surgery | 2.84 (1.19) | 2.50 (1.20) | 2.20 (1.20) |                                                        |                                                        |

Intervention groups: Standard Medical Care (SMC), Supportive Intervention (SUPPORT) or Expectation Manipulation Intervention (EXPECT); Assessment times (adjusted for baseline scores): 1 day pre-surgery, 1 week post-surgery and 6 months after surgery. Statistically significant results are displayed in bold.

![Image](Fig. 2 Post-hoc tests comparing intervention groups. Patients values of depressive symptoms (HADS) for low (−1 SD), average (mean) and high (+1 SD) baseline rates receiving SMC, SUPPORT or EXPECT at baseline, 1 day before surgery, 1 week after surgery and 6 months after surgery. *$p < 0.05$)
results of the analyses were included in the Supplementary material.

**Moderation effects of baseline depressive symptoms**

The baseline score of depressive symptoms (HADS) moderated the intervention effects on depressive symptoms. The significant group by time by baseline depressive symptoms-interaction implied this moderation \( (p = 0.040, \text{Table 2}) \). For all baseline scores of depressive symptoms significant group differences were reported in post-hoc simple slope analyses (see Fig. 2): Patients with high baseline scores of depressive symptoms (+1 SD) receiving SUPPORT or EXPECT showed significant lower scores of depressive symptoms 6 months after surgery compared to the control group (EXPECT vs. SMC: \( p = 0.015 \), SUPPORT vs. SMC: \( p = 0.004 \)). 1 week after surgery patients receiving SUPPORT also showed significant lower scores of depressive symptoms compared to SMC \( (p = 0.009) \), while there were no significant differences for EXPECT vs. SMC \( (p = 0.246) \). No statistically significant differences were observed 1 day before surgery for high baseline scores of depressive symptoms. There were no statistically significant differences between the intervention groups \( (p \geq 0.077) \). Patients with average scores of depressive symptoms at baseline (mean) receiving SUPPORT showed significant lower scores of depressive symptoms 1 week \( (p = 0.007) \) and also 6 months \( (p = 0.020) \) after surgery but not 1 day before surgery \( (p = 0.201) \) compared to SMC, while there were no significant differences for EXPECT vs. SMC at any time \( (1 \text{ day pre-surgery: } p = 0.741, 1 \text{ week post-surgery: } p = 0.243, 6 \text{ months post-surgery: } p = 0.057) \). No significant group differences were found between EXPECT and SUPPORT \( (p \geq 0.105) \). Patients with low baseline scores of depressive symptoms \( (-1 \text{ SD}) \) receiving SUPPORT showed significant lower scores of depressive symptoms 1 day before surgery compared to the control group \( (SMC; p = 0.028) \). No statistically significant differences were observed regarding the other measurement points for low baseline scores of depressive symptoms \( (p \geq 0.363) \). Confidence intervals and further details of post-hoc tests can be found in the Supplementary Table 2.

**Moderation effects of baseline personal control**

Perceived personal control (B-IPQ) at baseline moderated the effects of the preoperative psychological interventions significantly. The significant group by time by baseline level of personal-control-interaction indicated this moderation \( (p = 0.045, \text{Table 2}) \). For all baseline scores (low, average, high) of personal control post-hoc simple slope analyses indicated statistically significant group differences (see Fig. 3): Patients with a low baseline personal control score \( (-1 \text{ SD}) \) receiving EXPECT showed significant higher personal control scores 1 day before surgery compared to SMC and SUPPORT \( (both: p < 0.001) \), while there were no significant differences for SUPPORT vs. SMC \( (p = 0.964) \). No statistically significant differences were observed regarding the other measurement time points for low baseline personal control \( (p \geq 0.127) \). Patients with an average personal control score at baseline \( (mean) \) receiving EXPECT or SUPPORT showed significant higher personal control values 1 day before surgery compared to SMC \( (EXPECT: p < 0.001, SUPPORT: p = 0.045) \). There were no statistically significant differences at other measurement timepoints for average

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**Fig. 3** Post-hoc tests comparing intervention groups. Patients scores of personal control (B-IPQ) for low \((-1 \text{ SD})\), average \((\text{mean})\) and high \((+1 \text{ SD})\) baseline values receiving SMC, SUPPORT or EXPECT at baseline, 1 day before surgery, 1 week after surgery and 6 months after surgery. *\(p < .05\)
baseline personal control ($p \geq 0.127$). No significant group differences were found between EXPECT and SUPPORT ($p \geq 0.118$). Patients with a high baseline personal control score (+1 SD) receiving SUPPORT showed significant higher personal control values 1 day before surgery compared to SMC ($p = 0.005$), while there were no significant differences for EXPECT vs. SMC ($p = 0.284$). No statistically significant differences were observed regarding the other measurement time points for high baseline personal control ($p \geq 0.118$). There were no statistically significant differences between the intervention groups ($p \geq 0.118$). Confidence intervals and further details of post-hoc tests can be found in the Supplementary Table 3.

**Moderation effects of baseline concern**

Baseline levels of perceived concern (B-IPQ) moderated the effects of the preoperative psychological interventions significantly. The significant group by time by baseline level of concern-interaction indicated this moderation ($p = 0.046$, Table 2). For low baseline scores of concern significant group differences were reported in post-hoc simple slope analyses (see Fig. 4): Patients with a low baseline concern score (−1 SD) receiving EXPECT showed significant higher concern scores 1 day before surgery compared to SMC ($p = 0.045$), while there were no significant differences for SUPPORT versus SMC ($p = 0.938$) or SUPPORT vs. EXPECT ($p = 0.052$). No statistically significant differences were observed regarding the other measurement time points for low baseline concern ($p \geq 0.156$). There were no statistically significant differences at any measurement time points for patients with an average (mean) or high (+1 SD) score of concern at baseline ($p \geq 0.101$). Confidence intervals and further details of post-hoc tests can be found in the Supplementary Table 4.

**Additional analysis**

To explore associations between our psychological outcome criteria (HADS and B-IPQ) and physical outcomes, we explored the correlations between our outcomes and the number of patients rehospitalized due to complications and the pump function of the heart (ejection fraction; EF) at 6 months follow-up. For instance, depressive symptom levels and concern 6 months after surgery were positively associated with the amount of patients rehospitalized due to complications ($r_{\text{depressive symptoms}} = 0.214, p = 0.044$; $r_{\text{concern}} = -0.293, p = 0.005$). Rehospitalization scores after surgery were lowest in the EXPECT group (9% vs. 23% in the SUPPORT and 26% in the SMC group); however, this difference was not statistically significant (Rief et al., 2017). All results of the additional analysis can be found in the Supplementary Material Table 5.

**Discussion**

This article aimed to explore whether baseline levels of patients’ depressive/anxiety symptoms and illness beliefs moderated the effects of preoperative psychological interventions (EXPECT/SUPPORT) on these constructs in heart surgery patients to develop a better understanding of whether and when psychological interventions may have an effect on these important outcomes in heart surgery patients.

Baseline levels of depressive symptoms, personal control, and concern seemed to moderate the intervention effects on depressive symptoms, personal control, and concern. Especially for patients with high baseline scores of depressive symptoms, both preoperative psychological interventions led to reduced levels of depressive symptoms 6 months after surgery compared to the control group SMC. Considering the Minimally Clinically Important Difference (MCID) of

![Fig. 4](image-url)  
Fig. 4 Post-hoc tests comparing intervention groups. Patients scores of concern (B-IPQ) for low (-1 SD), average (mean) and high (+1 SD) baseline values receiving SMC, SUPPORT or EXPECT at baseline, 1 day before surgery, 1 week after surgery and 6 months after surgery.* $^p < .05$

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HADS scores for patients with cardiovascular disease that is ≥ 1.7 (Lemay et al., 2019), at least all discovered statistically significant effects of HADS were also clinically meaningful (≥ 1.763), except the discovered effect of SUPPORT and SMC for patients with average baseline levels of depressive symptoms at T3 (1.526). Also, some effects were found for illness beliefs, especially for personal control and concern: For a low personal control baseline-score the EXPECT-intervention led to increased personal control 1 day before surgery compared to SUPPORT and SMC. For patients with low baseline concern scores, the EXPECT-group indicated increased levels of concern 1 day before surgery compared to SMC. These results implicate that both preoperative psychological interventions may be especially relevant for patients with higher baseline depressive symptoms. Patients with lower baseline perceived personal control may benefit from the EXPECT-intervention.

As mentioned above, depression is a risk factor for many negative physical and psychological outcomes such as the higher risk of major adverse cardiac events, mortality, higher levels of medical complications, longer hospital stays, and lower quality of life after CABG surgery (AbuRuz, 2019; Auer et al., 2017; Blumenthal et al., 2003; Burg et al., 2003; Flaherty et al., 2017; McKenzie et al., 2010). Our study also indicated an association between depressive symptoms 6 months after surgery and postoperative complications leading to rehospitalization. Due to the importance of depression for psychological and physical outcomes, some studies tried to reduce depression in patients undergoing CABG surgery (Heilmann et al., 2016; McKenzie et al., 2010; Rollman, Belnap, LeMenager, Mazumdar, Houck et al., 2009; Rollman, Belnap, LeMenager, Mazumdar, Schulberg, & Reynolds III, 2009). The Bypassing the Blues (BtB) trial showed that CABG-patients who were depressed after their surgery profited from a collaborative care program (Rollman & Belnap, 2011). In the BtB-trial, patients participated after their surgery and had at least 1–28 contacts (median = 10) with the care manager in a period of eight months in the intervention groups. Compared to this trial, our results suggest that even a brief preoperative psychological intervention (five contacts) may be an effective way to decrease long-term levels of depressive symptoms and may improve heart surgery outcomes in patients with higher levels of depressive symptoms before surgery.

However, in our trial even most of the patients with ‘high’ baseline depressive symptoms did not reach the cut-off criterion for a clinical diagnosis of a depressive disorder (depression score ≥ 8) (Bjelland et al., 2002). Therefore, future research should examine how high a depressive burden has to be for letting patients benefit from a preoperative psychological intervention before undergoing surgery. Our results may not only be of scientific interest.

They may have important clinical implications: CABG-patients with significant depressive symptoms have almost twice the risk of having a cardiac event in the first 6 months after CABG-procedure (Scheier et al., 1999). In our trial, an association between depressive symptoms 6 months after surgery and complications leading to rehospitalization was found. Therefore, participation in a preoperative psychological intervention and ongoing support after the surgery (e.g., using booster calls) should be offered to these patients to improve psychological and physical outcomes. Given that further studies can replicate these findings, in line with the findings from Rollman and Belnap (2011), CABG patients should be screened for depressive symptoms before undergoing surgery. If depressed CABG-patients have a higher risk for adverse events such as cardiac events or mortality (Blumenthal et al., 2003; Scheier et al., 1999), patients with high levels of depressive symptoms who benefit from the individual surgery preparation would get the necessary assistance, while patients with lower levels (who may not benefit from the preoperative psychological interventions according to our results) would not need to take time for the intervention before and after surgery. Such a tailored treatment would provide every patient with the most profit and the fastest recovery possible. Also, the clinic and the healthcare system save capacities for the patients who need support and costs for less helpful interventions, or hospital stays (if the patients’ recovery is faster and the time patients stay in hospital is shorter) (Auer et al., 2017; Oxlad et al., 2006).

Future reimbursement models will most probably focus more on outcome parameters including quality of life (value based medicine), than on diagnoses and procedures (DRG systematics). Accordingly, interventions that can positively impact the short and long-term outcomes may be worthwhile from medical and economic perspectives. The results for depression also showed that patients with high or average levels of baseline depressive symptoms receiving the SUPPORT intervention benefitted 1 week and 6 months after the CABG surgery. Patients with low levels of baseline depressive symptoms receiving the SUPPORT intervention benefitted 1 day before the CABG surgery. Thus, additional psychosocial support seems to be helpful.

No significant differences between EXPECT (optimizing expectations) and SUPPORT (focusing on emotional support) were found for depressive symptoms. Against the background of the CSM, this may be explained by the fact that depressive symptoms can be characterized as an emotional and interpersonal challenge and a cognitive alteration; patients suffering from depression indicate dysfunctional cognitions and maladaptive information processing (Beck, 1979). Both interventions included at least one placebo mechanism (optimizing expectations, empathic relationship between patient and provider) (Schedlowski et al., 2015) targeting these factors. This may have led to
no significant differences between the intervention groups for depressive symptoms. Further research should focus on the question, which kind of intervention is the most helpful for patients with depressive symptoms to further elucidate crucial mechanisms.

Personal control as one of the illness beliefs seems to play an important part in the effects of preoperative psychological interventions. Previous studies indicated that personal control was most amenable to change compared to other illness belief dimensions (Broadbent et al., 2015; Laferton et al., 2016). In our trial, we found that receiving EXPECT or SUPPORT led to increased levels of personal control 1 day before surgery. Patients seemed to be more convinced to recover from or control their heart disease by their action after both preoperative psychological interventions than SMC patients. This could be due to the fact that patients pay attention to themselves and their heart disease in both interventions and therefore get the idea that their behavior influences their recovery. Furthermore, it is conceivable that patients in the SUPPORT group had the opportunity to self-reflect and may thus have come to the conclusion that their personal behavior may be part of their recovery. Since patients were not blind about their psychological intervention, receiving any kind of preoperative psychological intervention (compared to receiving only SMC) could have contributed to increased personal control levels in both psychological interventions.

The analyses indicated that patients with low perceived personal control levels at baseline benefitted from EXPECT in the short term compared to both other groups. Higher perceived personal control is associated with a higher quality of life and lower levels of depressive symptoms after CABG surgery (Kidd et al., 2016). Therefore, the increase of perceived personal control in our study might help avoid or reduce depressive symptoms in patients and hereby reduce physical outcomes as explained above. By now, only a short-term effect was found. Future studies should examine whether a sustained effect of increased personal control after surgery would have additional positive effects on long-term outcomes. Therefore, it should be examined, if more booster sessions can maintain the increase in perceived personal control for patients with lower baseline levels of personal control.

For patients with average and high baseline levels of personal control, the SUPPORT intervention led to increased levels of personal control 1 day before surgery. This finding may lead to the assumption that validation and emotional support may increase patients’ perceived controllability as someone strengthens the patients’ confidence and trust in their thoughts and preparations.

Regarding patients’ level of concern, a short-term effect was observed for patients with low scores of baseline concern. In the EXPECT group, an increase of concern was observed after the intervention 1 day before surgery. By focusing on psychoeducational aspects in this group, it is not surprising that patients’ worries increased short-dated. This result may explain why no effect was found for anxiety. By focusing on realistic expectations, patients also discussed topics that may have been perceived as concerning.

Some limitations need to be considered when interpreting the results of the study. Patients were only included in the study when they could appear in the study hospital a few days before the planned surgery date. Therefore, only patients with enough interest, time, and the possibility to drive to the hospital (even if some lived far away) were included. These facts may limit the generalizability of the findings. When getting informed, patients received the information that three treatment groups are included in the study and that two of them will receive additional conversations. It is possible that the expectation of receiving “just the standard of care” or “something special, additional” may have affected the outcomes. Focusing on depressive symptoms, most of the patients did not reach the cut-off criterium (depression score ≥ 8) (Bjelland et al., 2002). Therefore, the patients in the trial were not depressed on a high level. Further, due to the explorative character of the analyses conducted, our findings should be interpreted with caution. No correction for multiple testing has been done. Multi-centered confirmatory trials including more patients are needed to confirm the findings, generalize from one study site to the general healthcare systems, investigate further clinical outcome variables, and gain more knowledge, who would benefit from which intervention. It would be important to replicate the findings with a larger sample focusing on physical and psychological symptoms such as hospital stay, mortality, rehospitalization, depression, anxiety, illness beliefs and their associations.

In conclusion, our findings indicate that some patients may benefit from preoperative interventions while others will not. This study indicated that brief preoperative psychological interventions might improve critical psychological outcomes such as depressive symptoms or personal control in some heart surgery patients, but not in all patients. It further indicated that this may especially apply to specific subgroups of patients (i.e., high baseline depressive symptoms, low baseline personal control). Patients’ psychological status at baseline may moderate the effectiveness of psychological interventions. The second important finding is that assessing baseline levels is essential to offer tailored psychological interventions to improve long-term heart surgery outcomes. Gathering patients’ psychological status before undergoing heart surgery and providing psychological interventions if they are indicated (e.g., for patients with high scores of depressive symptoms or low levels of perceived control) would be beneficial. More studies are needed to examine
which patients may benefit from what kind of preoperative psychological intervention at which timepoint and why.

**Author contributions** Study conception, study design, and material preparation were performed by Melike Shedden-Mora, Johannes Laferton, Rainer Moosdorf, and Winfried Rief. Data collection was performed by Johannes Laferton, Stefan Salzmann, Rainer Moosdorf, and Winfried Rief. Analysis was performed by Nicole Horn and Stefan Salzmann. The first draft of the manuscript was written by Nicole Horn, and all authors commented on previous versions of the manuscript. All authors read and approved the final manuscript.

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**Data availability** Original data files and materials are stored at the division of clinical psychology. All authors had full access to the data. Data sets are available, as long as patients’ anonymity and ethical issues are respected.

**Code availability** Not applicable.

**Declarations**

**Conflict of interest** The authors have no conflicts of interest to declare that are relevant to the content of this article.

**Ethical approval** All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards. The study has been approved by the medical ethics committee of the Philipps University of Marburg and has been pre-registered at www.clinicaltrials.gov (NCT01407055) on August 1, 2011.

**Informed consent** Informed consent was obtained from all individual participants included in the study.

**Consent for publication** Not applicable.

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**References**

Abramson, J. H. (2011). Winpepi updated: Computer programs for epidemiologists, and their teaching potential. *Epidemiologic Perspectives & Innovations: EP+I*, 8, 1. https://doi.org/10.1186/1742-5573-8-1

AbuRuz, M. E. (2019). Pre-operative depression predicted longer hospital length of stay among patients undergoing coronary artery bypass graft surgery. *Risk Management and Healthcare Policy*, 12, 75.

Arthur, H. M., Daniels, C., McKeilie, R., Hirsh, J., & Rush, B. (2000). Effect of a preoperative intervention on preoperative and postoperative outcomes in low-risk patients awaiting elective coronary artery bypass graft surgery: A randomized, controlled trial. *Annals of Internal Medicine*, 133, 253–262.

Auer, C. J., Glombiewski, J. A., Doering, B. K., Winkler, A., Laferton, J. A. C., Broadbent, E., & Rief, W. (2016). Patients’ expectations predict surgery outcomes: A meta-analysis. *International Journal of Behavioral Medicine*, 23, 49–62.

Auer, C. J., Laferton, J. A. C., Shedden-Mora, M. C., Salzmann, S., Moosdorf, R., & Rief, W. (2017). Optimizing preoperative expectations leads to a shorter length of hospital stay in CABG patients: Further results of the randomized controlled PSY-HEART trial. *Journal of Psychosomatic Research*, 97, 82–89.

Beck, A. T. (1979). *Cognitive therapy of depression*. Guilford press.

Bjelland, I., Dahl, A. A., Haug, T. T., & Neckelmann, D. (2002). The validity of the Hospital Anxiety and Depression Scale: An updated literature review. *Journal of Psychosomatic Research*, 52, 69–77.

Blumenthal, J. A., Lett, H. S., Babyak, M. A., White, W., Smith, P. K., Mark, D. B., Jones, R., Mathew, J. P., Newman, M. F., & Investigators, N. (2003). Depression as a risk factor for mortality after coronary artery bypass surgery. *The Lancet*, 362, 604–609.

Bower, P., Kontopantelis, E., Sutton, A., Kendrick, T., Richards, D. A., Gilbody, S., Knowles, S., Cuijpers, P., Andersson, G., & Christensen, H. (2013). Influence of initial severity of depression on effectiveness of low intensity interventions: meta-analysis of individual patient data. *BMJ: British Medical Journal (Online)*, 346.

Broadbent, E., Ellis, C. J., Thomas, J., Gamble, G., & Petrie, K. J. (2009). Further development of an illness perception intervention for myocardial infarction patients: A randomized controlled trial. *Journal of Psychosomatic Research*, 67, 17–23.

Broadbent, E., Petrie, K. J., Main, J., & Weinman, J. (2006). The brief illness perception questionnaire. *Journal of Psychosomatic Research*, 60, 631–637.

Broadbent, E., Wilkes, C., Koschwanez, H., Weinman, J., Norton, S., & Petrie, K. J. (2015). A systematic review and meta-analysis of the Brief Illness Perception Questionnaire. *Psychology & Health*, 30, 1361–1385.

Burg, M. M., Benedetto, M. C., Rosenberg, R., & Soufer, R. (2003). Presurgical depression predicts medical morbidity 6 months after coronary artery bypass graft surgery. *Psychosomatic Medicine*, 65, 111–118.

Cameron, L. D., & Leventhal, H. (2003). *The self-regulation of health and illness behaviour*. Routledge.

Connerney, E., Shapiro, P. A., McLaughlin, J. S., Bagiella, E., & Sloan, R. P. (2001). Relation between depression after coronary artery bypass surgery and 12-month outcome: A prospective study. *The Lancet*, 358, 1766–1771.
Condra, R. J., Goyal, T. M., Cather, C., Rafafson, L., Idler, E. L., & Krause, T. J. (2004). Psychosocial factors in outcomes of heart surgery: The impact of religious involvement and depressive symptoms. *Health Psychology, 23*, 227.

Craig, C. L., Marshall, A. L., Sjöström, M., Bauman, A. E., Booth, M. L., Ainsworth, B. E., Pratt, M., Ekelund, U. L., Yngve, A., & Sallis, J. F. (2003). International physical activity questionnaire: 12-country reliability and validity. *Medicine & Science in Sports & Exercise, 35*, 1381–1395.

Davies, M. J., Heller, S., Skinner, T. C., Campbell, M. J., Carey, M. E., Craddock, S., Dallosso, H. M., Daly, H., Doherty, Y., Eaton, S., Fox, C., Oliver, L., Rantell, K., Rayman, G., & Khunti, K. (2008). Effectiveness of the diabetes education and self-management for ongoing and newly diagnosed (DESMOND) programme for people with newly diagnosed type 2 diabetes: Cluster randomised controlled trial. *BMJ, 336*, 491–495.

Dunkel, A., Kendel, F., Lehmkuhl, E., Hetzer, R., & Regitz-Zagrosek, V. (2011). Causal attributions among patients undergoing coronary artery bypass surgery: Gender aspects and relations to depressive symptomatology. *Journal of Behavioral Medicine, 34*, 351–359.

First, M. B., Spitzer, R. L., Gibbon, M., & Williams, J. B. W. (1996). Structured clinical interview for DSM-IV axis I disorders research version (SCID-I). *New York, NY: Biometrics Research, New York State Psychiatric Institute.*

Flaherty, L. B., Wood, T., Cheng, A., & Khan, A. R. (2017). Pre-existing psychological depression confers increased risk of adverse cardiovascular outcomes following cardiac surgery: A systematic review and meta-analysis. *The Journal of Thoracic and Cardiovascular Surgery, 154*, 1578–1586, e1.

Gallagher, R., & McKinley, S. (2009). Anxiety, depression and perceived control in patients having coronary artery bypass grafts. *Journal of Advanced Nursing, 65*, 2386–2396.

Guidi, J., Brakemeier, E.-L., Bockting, C. L. H., Cosci, F., Cuijpers, P., Jarrett, R. B., Linden, M., Marks, I., Peretti, C. S., & Rafanelli, C. (2018). Methodological recommendations for trials of psychological interventions. *Psychotherapy and Psychosomatics, 87*, 276–284.

Hawkes, A. L., & Mortensen, O. S. (2006). Up to one third of individual cardiac patients have a decline in quality of life post-intervention. *Scandinavian Cardiovascular Journal: SCJ, 40*, 214–218.

Hawkes, A. L., Nowak, M., Bidstrup, B., & Speare, R. (2006). Outcomes of coronary artery bypass graft surgery. *Vascular Health and Risk Management, 2*, 477–484.

Head, S. J., Osnabrugge, R. L. J., Howell, N. J., Freemantle, N., Bridge-Gallagher, R., & McKinley, S. (2009). Anxiety, depression and post-coronary artery bypass surgery: The impact of religious involvement and depressive symptoms and quality of life but not health behaviour following coronary artery bypass graft surgery. *Journal of Behavioral Medicine, 32*, 120–127. https://doi.org/10.1007/s10865-015-9677-7

Laforton, J. A. C., Auer, C. J., Shedden-Mora, M. C., Moosdorf, R., & Rief, W. (2016). Optimizing preoperative expectations in cardiac surgery patients is moderated by level of disability: The successful development of a brief psychological intervention. *Psychology, Health & Medicine, 21*, 272–285.

Laforton, J. A. C., Kube, T., Salzmann, S., Auer, C. J., & Shedden-Mora, M. C. (2017). Patients’ expectations regarding medical treatment: A critical review of concepts and their assessment. *Frontiers in Psychology, 8*, 233.

Laforton, J. A. C., Mora, M. S., Auer, C. J., Moosdorf, R., & Rief, W. (2013). Enhancing the efficacy of heart surgery by optimizing patients’ preoperative expectations: Study protocol of a randomised controlled trial. *American Heart Journal, 165*, 1–7.

Lamarche, D., Taddeo, R., & Pepler, C. (1998). The preparation of patients for cardiac surgery. *Clinical Nursing Research, 7*, 390–405.

Lau, R. R., & Hartman, K. A. (1983). Common sense representations of common illnesses. *Health Psychology, 2*, 167.

Lemay, K. R., Tulloch, E. H., Pipe, A. L., & Reed, J. L. (2019). Establishing the minimal clinically important difference for the hospital anxiety and depression scale in patients with cardiovascular disease. *Journal of Cardiopulmonary Rehabilitation and Prevention, 39*, E6–E11.

Leventhal, H., & Cameron, L. D. (1987). Behavioral theories and the problem of compliance. *Patient Education and Counseling, 10*, 117–138.

Leventhal, H., Dieffenbach, M., & Leventhal, E. A. (1992). Illness cognition: Using common sense to understand treatment adherence and affect cognition interactions. *Cognitive Therapy and Research, 16*, 143–163.

Leventhal, H., Leventhal, E. A., & Cameron, L. D. (2001). Representations, proce-dures and affect in illness self-regulation: A perceptual-cognitive model. In S. J. E. Revenson TA (Ed.), *Handbook of health psychology* (pp. 19–48). Lawrence Erlbaum.

Levett, D. Z., & Grimmett, C. (2019). Psychological factors, prehabilitation and surgical outcomes: Evidence and future directions. *Anaesthesia, 74*, 36–42.

Malik, S., Krumholz, H. M., Lin, Z. Q., Kasl, S. V., Mattera, J. A., Roumains, S. A., & Vaccarino, V. (2005). Patients with depressive symptoms have lower health status benefits after coronary artery bypass surgery. *Circulation, 111*, 271–277.

McKenzie, L. H., Simpson, J., & Stewart, M. (2010). A systematic review of pre-operative predictors of post-operative depression and anxiety in individuals who have undergone coronary artery bypass graft surgery. *Psychology, Health & Medicine, 15*, 74–93.

Moreno-Peral, P., Bellón, J. Á., Motrico, E., Campos-Paino, H., Martín-Gómez, C., Ebert, D. D., Buntrock, C., Roca, M., & Conejo-Cerón, S. (2020). Moderators of psychological and psychoeducational interventions for the prevention of anxiety: A systematic review. *Journal of Anxiety Disorders, 76*, 102317.

Morone, N. E., Weiner, D. K., Belnap, B. H., Karp, J. F., Mazumdar, S., Houck, P. R., He, F., & Rollman, B. L. (2010). The impact of pain and depression on post-CABG recovery. *Psychosomatic Medicine, 72*, 620.

Murray, C. J. L., & Lopez, A. D. (2013). Measuring the global burden of disease. *New England Journal of Medicine, 369*, 448–457.

Oxlad, M., Sterberfield, J., Stuklis, R., Edwards, J., & Wade, T. D. (2006). Psychological risk factors for increased post-operative… Springer
length of hospital stay following coronary artery bypass graft surgery. *Journal of Behavioral Medicine*, 29, 179–190.

Parfeni, M., Nistor, I., & Covic, A. (2013). A systematic review regarding the association of illness perception and survival among end-stage renal disease patients. *Nephrology Dialysis Transplantation*, 28, 2407–2414.

Petrie, K. J., Cameron, L. D., Ellis, C. J., Buick, D., & Weinman, J. (2002). Changing illness perceptions after myocardial infarction: An early intervention randomized controlled trial. *Psychosomatic Medicine*, 64, 580–586.

Petrie, K. J., Perry, K., Broadbent, E., & Weinman, J. (2012). A text message programme designed to modify patients’ illness and treatment beliefs improves self-reported adherence to asthma preventer medication. *British Journal of Health Psychology*, 17, 74–84.

Petrie, K. J., Weinman, J., Sharpe, N., & Buckley, J. (1996). Role of patients’ view of their illness in predicting return to work and functioning after myocardial infarction: Longitudinal study. *BMJ*, 312, 1191–1194.

Poole, L., Kidd, T., Leigh, E., Ronaldson, A., Jahangiri, M., & Steptoe, A. (2015). Psychological distress and intensive care unit stay after cardiac surgery: The role of illness concern. *Health Psychology*, 34, 283.

Poole, L., Kidd, T., Ronaldson, A., Jahangiri, M., & Steptoe, A. (2014). The combined association of depression and socioeconomic status with length of post-operative hospital stay following coronary artery bypass graft surgery: Data from a prospective cohort study. *Journal of Psychosomatic Research*, 76, 34–40.

Poole, L., Ronaldson, A., Kidd, T., Leigh, E., Jahangiri, M., & Steptoe, A. (2017). Pre-surgical depression and anxiety and recovery following coronary artery bypass graft surgery. *Journal of Behavioral Medicine*, 40, 249–258.

Preacher, K. J., Curran, P. J., & Bauer, D. J. (2006). Computational tools for probing interactions in multiple linear regression, multi-level modeling, and latent curve analysis. *Journal of Educational and Behavioral Statistics*, 31, 437–448. https://doi.org/10.3102/10769986031004437

Rief, W., Shedden-Mora, M. C., Laferton, J. A. C., Auer, C., Petrie, K. J., Salzmann, S., Schedlowksi, M., & Moosdorff, R. (2017). Preoperative optimization of patient expectations improves long-term outcome in heart surgery patients: Results of the randomized controlled PSY-HEART trial. *BMC Medicine*, 15, 4. https://doi.org/10.1186/s12916-016-0767-3

Rollman, B. L., & Belnap, B. H. (2011). The Bypassing the Blues trial: Collaborative care for post-CABG depression and implications for future research. *Cleveland Clinic Journal of Medicine*, 78, S4-12.

Rollman, B. L., Belnap, B. H., LeMenager, M. S., Mazumdar, S., Schulberg, H. C., & Reynolds, C. F., III. (2009). The Bypassing the Blues treatment protocol: Stepped collaborative care for treating post-CABG depression. *Psychosomatic Medicine*, 71, 217.

Rumsfeld, J. S., Ho, P. M., Magid, D. J., McCarthy, M., Jr., Shroyer, A. L. W., MaWhinney, S., Grover, F. L., & Hammermeister, K. E. (2004). Predictors of health-related quality of life after coronary artery bypass surgery. *The Annals of Thoracic Surgery*, 77, 1508–1513.

Sadeghi, M., Hashemi, M., Sararoudi, R. B., Merasi, M. R., Molaein-ezhad, M., & Shamsolketabi, H. (2017). Demographic and psychological predictors of recovery from coronary artery bypass graft. *Journal of Education and Health Promotion*, 6.

Salzmann, S., Euteneuer, F., Laferton, J. A. C., Shedden-Mora, M. C., Schedlowksi, M., Moosdorff, R., & Rief, W. (2020a). IL-8 and CRP moderate the effects of preoperative psychological interventions on postoperative long-term outcomes 6 months after CABG surgery - The randomized controlled PSY-HEART trial. *Brain, Behavior, and Immunity*.

Salzmann, S., Laferton, J., Auer, C., Shedden-Mora, M. C., Wambach, K., & Rief, W. (2018). Patientenerwartungen optimieren: Beschreibung einer präoperativen Kurzintervention am Beispiel von Patienten vor einer Bypass-Operation. *Verhaltenstherapie*, 28, 157–165.

Salzmann, S., Salzmann-Djufri, M., Wilhelm, M., & Euteneuer, F. (2020). Psychological preparation for cardiac surgery. *Current Cardiology Reports*, 22, 172.

Scheier, M. F., Matthews, K. A., Owens, J. F., Schulz, R., Bridges, M. W., Magovern, G. J., & Carver, C. S. (1999). Optimism and rehospitalization after coronary artery bypass graft surgery. *Archives of Internal Medicine*, 159, 829–835.

Széky, A., Balog, P., Benkő, E., Breuer, T., Székely, J., Kertai, M. D., Horkay, F., Kopp, M. S., & Thayer, J. F. (2007). Anxiety predicts mortality and morbidity after coronary artery and valve surgery— A 4-year follow-up study. *Psychosomatic Medicine*, 69, 625–631.

Tait, R. C., Chibnall, J. T., & Krause, S. (1990). The pain disability index: Psychometric properties. *Pain*, 40, 171–182.

Timberlake, N., Klinger, L., Smith, P., Venn, G., Treasure, T., Harrison, M., & Newman, S. P. (1997). Incidence and patterns of depression following coronary artery bypass graft surgery. *Journal of Psychosomatic Research*, 43, 197–207.

Tully, P. J., Baker, R. A., Turnbull, D. A., Winefield, H. R., & Knight, J. L. (2009). Negative emotions and quality of life six months after cardiac surgery: The dominant role of depression not anxiety symptoms. *Journal of Behavioral Medicine*, 32, 510.

Virani, S. S., Alonso, A., Benjamin, E. J., Bittencourt, M. S., Callaway, C. W., Carson, A. P., Chamberlain, A. M., Chang, A. R., Cheng, S., & Delling, F. N. (2020). Heart disease and stroke statistics—2020 update: A report from the American Heart Association. *Circulation*, E139-E596.

Ware, J. E., Jr., Kosinski, M., & Keller, S. D. (1996). A 12-item short-form health survey: Construction of scales and preliminary tests of reliability and validity. *Medical Care*, 34, 220–233.

Weinman, J., Petrie, K. J., Sharpe, N., & Walker, S. (2000). Causal attributions in patients and spouses following first-time myocardial infarction and subsequent lifestyle changes. *British Journal of Health Psychology*, 5, 263–273.

Wynter-Blyth, V., & Moorthy, K. (2017). Prehabilitation: preparing patients for surgery. *BMJ: British Medical Journal (Online)*, 358.

Young, S., Linden, W., Ignaszewski, A., Con, A., Terhaag, S., & Campbell, T. (2019). Psychosocial and medical predictors of 1-year functional outcome in male and female coronary bypass recipients. *Heart and Mind*, 3, 113.

Zigmond, A. S., & Snaith, R. P. (1983). The hospital anxiety and depression scale. *Acta Psychiatrica Scandinavica*, 67, 361–370.

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