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Mindfulness-based programme for residents: study protocol of a randomised controlled trial

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ABSTRACT

Introduction Residency is a stressful phase associated with high prevalence of mental distress. Besides impaired personal health, mental distress in residents has an impact on the quality of patient care and produces economic costs. Therefore, there is demand for interventions that improve resident physicians’ mental health. The aim of the present study is to examine the effects of a mindfulness-based intervention that has been tailored to residents’ needs. Specifically, mindfulness has been supplemented by a focus on the concept of Muße.

Methods and analysis This study applies a randomised controlled multimethod design. Residents assigned to the intervention group will participate in an 8-week mindfulness course followed by a 4-month maintenance phase, whereas residents assigned to the control group will read text-based information about mindfulness on a weekly basis for the duration of 8 weeks. The intervention is focussed on a transfer of learnt techniques into the daily routine and is targeted to promote residents’ self-care as well as on building empathic relationships. Participants will be assessed before, directly after the intervention, after the maintenance phase as well as at follow-up 6 months after the intervention group completes the intervention. Assessments will consist of self-report measures, physiological data, qualitative interviews, third-party reports as well as implicit and projective measures and will focus on both psychopathology and salutogenesis. The primary outcome will be burnout. Data will be analysed using linear mixed modelling.

Ethics and dissemination The study was approved by the ethics committee of the Medical Center - University of Freiburg and is funded by the German Research Foundation as part of the interdisciplinary Collaborative Research Center ‘SFB Muße 1015’. The results of this study will be published in scientific journals and disseminated through the study’s website, and conferences.

Trial registration number DRKS00014015.

INTRODUCTION

Several studies show that hospitals are a challenging workplace as characterised by hierarchical structures, work density, time pressure, documentation duties and increasing focus on economic aspects. Within this system it is recognised that residency is an outstandingly stressful phase due to long working hours combined with high workload, great responsibility, not feeling competent, lack of supervision, unclear hierarchies, lack of work-life balance and emotional strain.

Accordingly, prevalence of mental distress is high in residents. Several large-scale studies report rates of burnout between 20% and 60%, rates of depression between 20% and 50% and rates of anxiety between 30% and 40%. Furthermore, residents with burnout have been shown to have increased rates of suicidality and substance abuse. However, stress at the workplace hospital and mental distress in physicians not only have consequences for residents themselves. That is, mental distress in physicians is associated with decreased patient safety and increased treatment errors. At the same time, stress is associated with decreased empathy and compassion while empathy has been shown to be an important factor in

Strengths and limitations of this study

► This study investigates a mindfulness-based intervention tailored to the particular needs and circumstances of resident physicians.
► The study investigates a long-term intervention (8-week course, 4-month maintenance phase) using a longitudinal study design with four assessment points over the course of 1 year.
► We use a multimethod assessment including self-reports, physiological data, qualitative interviews, third-party reports as well as implicit and projective measures.
► If results are positive, this study will provide evidence for the effectiveness of an intervention through which residents can learn how to cope with stress and improve their mental well-being.
► The lack of a third group that receives standard mindfulness-based stress reduction (MBSR) will not allow for determining whether a mindfulness course tailored to residents’ needs is superior to standard MBSR.
treatment success and patient satisfaction.25 Furthermore, mental distress in physicians is associated with increased economic costs due to early retirement, reduced clinical hours and increased physician turnover.26–28 In sum, mental distress in residents comes at a price on residents’ personal health, patient care as well as economic costs.

As a result, there is great demand for interventions that prevent mental distress and improve residents’ well-being. Mindfulness-based interventions (MBI) are a promising approach to reduce mental distress and promote residents’ well-being. Mindfulness is defined as a state of awareness that is characterised by focussing one’s attention on the present moment with an attitude of openness and acceptance.29 A large body of literature demonstrates the beneficial effects of mindfulness interventions on reducing stress, depression and anxiety while increasing well-being and quality of life.30 31 Accordingly, several studies show similar results for MBIs in physicians.32–38 In addition to improving physicians’ mental health and well-being, these studies also show that MBIs improve the quality of treatment.39–42 Consistent with these findings, preliminary studies reported beneficial effects of MBIs in residents.43–45 For instance, Rosdahl and others43 found that a single mindfulness session decreased perceived stress and burnout. Similarly, Lases and others44 and Ireland and others45 reported significant decreases in perceived stress and burnout after 5 or 10 sessions of mindfulness. However, by contrast Goldhanen and others,46 Verweij and others,47 and Taylor and others48 did not find such an association. These initial studies are promising although inconclusive.

Residency represents a special stage within the professional career; therefore a MBI tailored to residents’ needs may be more effective. In support of this, qualitative analyses by Lases and others44 indicated that residents who participated in a mindfulness course would have welcomed a tailoring of the programme to their needs. Furthermore, secular mindfulness interventions are increasingly being criticised for focussing too much on stress reduction, self-optimisation and improved performance rather than self-awareness and equanimity.49 Particularly within the context of clinical training that promotes self-sacrificing attitudes and performance orientation,50 mindfulness interventions are prone to be functionalised to further increase stress tolerance and performance. Consequently, there is a need for MBIs that emphasise a complementary state to daily stress and continuous performance orientation. Therefore, the aim of the present study is to investigate the effects of a MBI that has been tailored to residents’ needs. To prevent the intervention from solely focussing on increasing performance and stress reduction, the main target of the intervention, that is, mindfulness, will be supplemented by a focus on Muße within the present study. The term Muße is well introduced in the German language but cannot be directly translated into English. Muße refers to a state of serenity, equanimity, self-sufficiency and fulfilment in which one feels free of pressure.51 52 In other words, when experiencing Muße individuals feel at ease, content and free. By implementing a focus and goal on the experience of Muße as a state, the aim of the MBI is to enable participants—also at work—to access inner states that are beyond task-orientation, stress and time pressure. At a first glance, Muße and work may appear to contradict each other. However, being present in the moment and one’s inner attitude are key to the experience of Muße regardless of context, and these qualities can be promoted through mindfulness practice. As a result of implementing a focus on Muße, the intervention aims at creating an opposite pole to stress and functioning.

In sum, the aim of this trial is to investigate a Muße-directed mindfulness intervention tailored to residents’ needs consisting of an 8-week course followed by a 4-month maintenance phase. A randomised controlled design will be applied including quantitative, qualitative, implicit, projective, third-party, as well as physiological measures. The primary outcome is change in burnout from baseline to 6 months. The control group will receive a coursebook containing text-based information about mindfulness sent to participants on a weekly basis for the duration of 8 weeks. This will mimic the structure of the intervention group in terms of information and knowledge but not with respect to guided practice of mindfulness (ie, experience). We hypothesise that the mindfulness intervention compared with the control group will lead to (A) a reduction in psychopathological variables including burnout, perceived stress, depression and anxiety, and (B) will lead to an increase in positive aspects of mental health, including flourishing, satisfaction with life and self-compassion. Further, the aim of the qualitative analysis is to gain understanding of the effects of the MBI on participants’ everyday life.

**METHOD**

**Design**

A randomised, controlled trial applying a mixed-method approach will be conducted. Participants will be randomly assigned to either the intervention group receiving an 8-week mindfulness course followed by a 4-month maintenance phase or to the control group receiving a coursebook containing theoretical information about mindfulness for self-study. Assessments will be held at four time points: baseline (t0, 0 months), post-intervention (t1, 2 months), after the maintenance phase (t2, 6 months) as well as 6 months after the end of the maintenance phase (t3, 12 months).

**Setting**

The study will be run in the south-west part of Germany. Residents will be recruited from a wide range of hospitals including university hospitals, hospitals in private sponsorship, hospitals in urban as well as rural areas. The courses of the mindfulness intervention will take place at the Medical Center of the University of Freiburg,
Germany, while the control group will receive the self-study material about mindfulness via email.

Participants
To be included in the study, participants need to (1) be employed as resident at a hospital, (2) have a work contract with a minimum employment of 40%, (3) have regular contacts with patients, (4) be less than 45 years old, (5) have sufficient German language skills and (6) have given informed consent. Exclusion criteria is not being willing to conform to study requirements.

Recruitment, consent and allocation to interventions
Residents will be recruited by approaching hospitals in south-west Germany and advertising the study to the employed residents, by email, intranet, informing head physicians about the study and by giving short presentations about the trial at the hospitals. Additionally, a webpage and flyers will inform about the trial. Interested residents may sign up using the registration form on the webpage or via email.

Before taking part in the study, residents will be provided with detailed information about the study in written form and will be given the opportunity to ask questions to a trained assessor. After consent has been obtained and the baseline assessment has been completed, participants will be assigned to either the intervention or the control group using a minimisation procedure for stratification. Stratification factors used for minimisation are gender and level of burnout (Copenhagen Burnout Inventory (CBI) values 1 to 2.5=low, 2.6 to 3.5=medium, 3.6 to 6.5=high burnout). The minimisation procedure applies a base probability of allocation of 0.8 and variance as distance measure. Minimisation will be carried out with the software QMinim. Allocation concealment will be ensured, as minimisation will take place after completion of all baseline assessments and will be carried out by a researcher who has no direct contact with participants. The flow of participants from recruitment to the end of the study is displayed in figure 1.

Intervention
Mindfulness-based intervention: Residents allocated to this condition will take part in an 8-week course with weekly sessions lasting 2.5 hours each as well as a full day retreat (day of mindfulness, between Session 5 and 6), followed by a 4-month maintenance phase consisting of three monthly booster sessions lasting 2.5 hours each. Additionally, participants are encouraged to practice at home by completing homework over the course of the 8-week intervention. The courses will be organised in groups of 6 to 14 participants and will be led by three physicians who are certified mindfulness-based stress reduction (MBSR) and mindfulness-based cognitive therapy (MBCT) teachers with extensive experience in teaching mindfulness classes.

The mindfulness intervention is based on the original manual of MBSR by John Kabat-Zinn, has been tailored to residents’ needs and given a focus on Muße. To tailor the programme, we conducted an a priori needs assessment consisting of an in-depth literature search (ie, residency specific stressors, character traits that are more prevalent in residents, content and culture of medical training) as well as a set of interviews with residents (ie, asking about residency specific stressors and how they experience their work environment). The specific customisation addressed four aspects: First, we identified and integrated resident specific topics such as mindful communication with patients or dealing with stressors of residency (ie, high responsibility while not feeling competent enough, lack of supervision, unclear hierarchies or high workload, fear of making mistakes, etc). This also includes the capability to establish an empathic and compassionate relationship with patients despite time pressure. Table 1 gives an overview of the topics covered during the course. Second, a result of the literature search was that within the context of medical education, a MBI might run the danger of solely being used for self-optimisation and to function better under stress. Therefore, mindfulness is supplemented by a focus on Muße that is incompatible with self-optimisation. The word Muße is well established in the German language. Thus, the aim is to activate the cognitive concept of Muße and create an association between states of Muße and practicing mindfulness meditation.
experience of Muße and participants are encouraged to practice mindfulness in a spirit of Muße. A third adaptation is the inclusion of a large number of transfer exercises with the aim of enabling participants to get into a state of mindful awareness during everyday life (informal practice) and not only during formal meditation practice. Example of such transfer exercises are, feeling one’s feet while walking down the hospital corridor, feeling one’s body during regular hand disinfection or taking a short break observing one’s breath before entering a patient’s room. Fourth, additional information and scientific background to course elements is given to residents. Such an approach complies with the recommendations of the American Medical Association, British Medical Association and Canadian Medical Association for mindfulness interventions addressing physicians.

On the structural level, every course session consists of five elements: (1) an educational input, (2) mindfulness exercises (ie, sitting meditation, body scan, mindful yoga or mindful walking), (3) a reflection of one’s practice, (4) transfer exercises such as role plays or group exercises and (5) home assignments.

The maintenance phase consists of three monthly booster sessions of 2.5 hours each during which exercises learnt and topics treated in the course are refreshed. Additionally, the sessions of the maintenance phase allow for space for participants to reflect and discuss their experiences and difficulties in practicing and applying mindfulness in their work context as well as in everyday life.

Control group: Residents assigned to the control group will receive a coursebook on mindfulness containing the same written material about mindfulness practice and the respective background that participants of the intervention group will receive. This text-based material consists of a detailed description and definition of mindfulness, studies on mindfulness, stress, coping with stress, acceptance, self-care and meaning in medicine as well as inspiring stories and poems about mindfulness. The chapters of the coursebook will be sent to participants on a weekly basis via email for the duration of 8 weeks, in parallel to the mindfulness course. By this choice of control group, we can compare experience-based learning with text-based learning.

Outcome measures

The aim of the present study is to investigate the effects of the adapted mindfulness intervention on both stress and associated pathology and positive mental health. The primary outcome (ie, burnout) as well as secondary outcomes will be assessed at four points of time: at baseline (t0, 0 months), after the intervention (t1, 2 months), after the maintenance phase (t2, 6 months), as well as 6 months after completion of the maintenance phase (t3, 12 months). Assessment methods used include standardised self-report questionnaires, implicit tests,

| Session | Topic | Content |
|---------|-------|---------|
| 1       | Mindfulness | Exploring residents’ needs and expectations; introducing mindfulness as a mode of being, as contrasted to a mode of doing and performing during everyday life. |
| 2       | Dealing with barriers and subjective time perception | Discussing ways to deal with barriers to mindfulness practice; exploring mindfulness anchors in daily routine (eg, mindful walking along hospital corridors; mindful stop before entering a patient’s room; mindful hand disinfection); introducing mindfulness to experience slower passage of time and to mitigate the feeling of time pressure. |
| 3       | Dis-identification | Coping with painful emotions, thoughts and physical sensations; raising awareness of the process of constructing reality through ones experiences; connecting with the inner-observer and exploring dis-identification to learn to non-identify with the self and to reduce reactivity towards them. |
| 4       | Stress | Discussing resident specific stressors; psychoeducation on physiological and psychological processes of stress; exploring how to cope with stress using mindfulness. |
| 5       | Acceptance | Learning acceptance of oneself as well as the given reality of experiences, events and working conditions. Exploring the difference between acceptance and resignation/fatalism and the importance of acceptance for self-care. |
| 6       | Mindfulness in patient contact | Using mindfulness in therapeutic interactions; building a compassionate communication atmosphere with patients, even in moments of time pressure. Learning to listen mindfully and exploring the benefits of letting patients complete voicing their agenda of concerns. |
| 7       | Self-care | Discussing why self-care is especially relevant to resident physicians and its connection to quality of care; exploring ways to take care of oneself in daily routine. |
| 8       | Enhancing meaning in work and mindfulness as part of life | Exploring what is meaningful in professional life and how meaning can be enhanced. Reinforcing mindfulness as part of everyday life and daily medical practice. |
projective tests, physiological and behavioural markers, qualitative measures and third-party reports. Additionally, at baseline, demographics will be assessed as well as participants’ previous experience with meditation. Similarly, at t3, major life events that might influence the results will be assessed as well as participants’ sick leave days over the past year.

**Primary outcome**
The primary outcome is change in burnout between t0 and t2, and it will be assessed using the German version of the CBI.58 59 The CBI is a 19-item self-report questionnaire that yields three subscales for personal burnout, work-related burnout and client-related burnout. Participants will rate to what extent they experience certain symptoms such as feeling tired or exhausted on a 5-point Likert scale from 0 (never/rarely) to 100 (very often). The CBI has been shown to have high internal consistency with a Cronbach’s alpha between 0.85 and 0.87.58 59

**Secondary outcomes**

**Self-report questionnaires**
We differentiate between measures pertaining to psychopathology and measures pertaining to salutogenesis as well as additional variables including errors at work, self-reported mindfulness, physician empathy and perception of time.

The measures pertaining to psychopathology include perceived stress, psychopathology and mental distress, job strain, depression and anxiety. Perceived stress will be assessed by the German version of the Perceived Stress Scale (PSS).60 61 The PSS consists of 10 items that require participants to rate how often they experienced certain aspects of stress during the preceding month using a 5-point Likert scale from 0 (never/rarely) to 100 (very often). The scale has been shown to have good internal consistency with a Cronbach’s alpha of 0.83 to 0.86.60 61

Psychopathology and mental distress will be assessed using the German version of the General Health Questionnaire (GHQ-12)62 63 an instrument commonly used as a screening instrument for psychological disorders. Participants rate how often they experienced certain symptoms such as being able to concentrate or thinking of oneself as worthless during the past weeks on a 4-point Likert scale from 0 (not at all) to 3 (much more than usual). The GHQ-12 has been shown to give good internal consistency with a Cronbach’s alpha of 0.91.63

Depression and anxiety will be assessed with the German translation of the Patient Health Questionnaire (PHQ-4) that consists of four screening items for depression and anxiety.64 65 Participants will rate how often they experienced certain symptoms over the past 2 weeks on a 4-point Likert scale from 0 (not at all) to 3 (almost every day). Both the PHQ-4 and the GHQ-12 have been shown to have high validity and internal consistency with a Cronbach’s alpha between 0.86 and 0.9162 63 and between 0.82 and 0.85.64 65

Job strain will be measured using the German version of the Irritation Questionnaire.66 This scale asks participants to rate eight statements about emotional and cognitive strain on a 7-point Likert scale from 1 (not at all) to 7 (very much). Internal consistency has been shown to be good with a Cronbach’s alpha between 0.85 and 0.93.66

Secondary outcomes focussing on positive mental health (ie, salutogenesis) include well-being, self-compassion, self-esteem, self-efficacy as well as the feeling of being loved. Well-being comprises both hedonic and eudaimonic aspects.67 Hedonic aspects include positive affect and satisfaction with life. Positive affect will be assessed using the Self-Assessment Manikin (SAM).68 The SAM requires participants to indicate their emotional state via five abstract faces depicting a semantic differential from negative (frowning face, coded as 1) to positive emotions (happy face, coded as 5).

Satisfaction with life will be assessed with the one item short scale L-169 that asks participants to give a global rating of how satisfied they are with their life on an 11-point Likert scale from 0 (not satisfied at all) to 10 (absolutely satisfied). The L-1 has been shown to have good test-retest reliability as well as high convergent validity with other, multi-item scales that measure satisfaction with life.69

Eudaimonic well-being can be defined as positive functioning in daily life70 and will be assessed using the German translation of the Flourishing Scale (FS).70 71 The FS requires participants to rate eight statements about the experience of certain aspects of well-being in their lives on a 7-point Likert scale ranging from 7 (strongly agree) to 1 (strongly disagree). The FS has been shown to have good internal consistency with a Cronbach’s alpha between 0.86 and 0.87.70 71

Well-being specifically related to the job will be assessed with the German translation of the Thriving at Work Scale (TS)72 that measures positive functioning at the workplace. The TS consists of ten statements that are rated on a 5-point Likert scale from 1 (not true at all) to 5 (absolutely true) and has been shown to have good internal consistency with a Cronbach’s alpha between 0.90 and 0.93.72

Job satisfaction (ie, a person’s cognitive and affective evaluations of their work) will be assessed by the Faces Scale.73 This scale consists of seven abstract faces representing a semantic differential from negative (frowning face, coded as 1) to positive (happy face, coded as 7).

Self-compassion can be defined as being kind and understanding towards oneself74 and will be measured with the German translation of the short form of the Self-Compassion Scale (SCS).75 76 The short form of the SCS consists of 12 statements that are rated on a 5-point Likert scale from 1 (almost never) to 5 (almost always). The scale has been shown to have a good internal consistency with a Cronbach’s alpha of 0.86.75 76

Global self-esteem will be measured with the Single-Item Self-Esteem Scale (SISE),77 consisting of rating one statement (‘I have high self-esteem’) on a 5-point scale from 1 (not very true for me) to 5 (very true for me). A
validation study demonstrated good test-retest reliability and good convergent validity with other longer measures of self-esteem. The SISE has been translated to German by the authors.

Self-efficacy or the belief in one’s ability to succeed will be assessed using the Self-Efficacy Questionnaire. This self-report scale consists of three items that are scored on a 5-point Likert scale from 1 (not true at all) to 5 (very true). The scale has good internal consistency with a McDonald’s omega of 0.81.

The feeling of being loved by others and the feeling of loving oneself will be assessed with the Feeling Loved Questionnaire. This questionnaire consists of one Yes/No question (ie, ‘Do you feel loved?’) followed by an item assessing the continuous rating (ie, ‘How loved do you feel?’) for both dimensions. The scale has good convergent and discriminant validity. The feeling loved questionnaire was translated to German by the authors.

Self-reported mindfulness will be assessed using the Freiburg Mindfulness Inventory (FMI). The FMI is a self-report instrument consisting of 14 statements about specific aspects of mindfulness experienced in everyday life. Participants indicate how often they experienced these aspects during the past week on a 4-point Likert scale from 0 (rarely) to 3 (almost always). The questionnaire has good internal consistency with a Cronbach’s alpha of 0.86.

Subjective perception of time will be measured by an abridged version of the Time Perception Questionnaire that requires participants to rate statements about how time is perceived on a 5-point Likert scale from strongly disagree to strongly agree. To generate an abridged version, the author of the original questionnaire selected five items, which he considered to be the most relevant.

Physician empathy refers to the ability of understanding the patient’s perspective and being able to communicate this. Physician empathy will be assessed using a shortened version of the Jefferson Scale of Physician Empathy. The original scale consists of 20 statements that are answered on a 7-point scale from 1 (strongly disagree) to 7 (strongly agree). However, in the present study we chose six items that have been shown to have the highest factor loadings in two validation studies with 193 and 704 physicians, respectively. We used the German translation mentioned in.

The experience of Muße will be measured using a self-constructed questionnaire. We believe that like satisfaction with life, Muße is a state that is directly accessible and can thus be measured by directly asking participants about respective experiences. Thus, in a first step, participants will indicate how often they experienced Muße during the past 4 weeks. Answers are given on a 6-point Likert scale from never to very often. In a second step, participants will report how often they experienced Muße within specific situations (ie, at work, in their leisure time, at home, away from their home, when being alone, during quiet situations, when being in action) during the past 4 weeks. Answers are given on the same 6-point Likert scale as in the first part.

Errors at work are assessed with a self-report questionnaire that distinguishes two types of errors: errors due to lack of experience and errors due to lack of time. Participants will indicate how often certain errors had occurred giving their ratings on a 5-point Likert scale from 1 (never occurs) to 5 (occurs often). The questionnaire has satisfactory internal consistency with a Cronbach’s alpha between 0.62 and 0.64 for each subscale. The six questions were translated into German by the authors.

Physiological and behavioural outcome measures

In addition to self-report measures, stress will be assessed by measuring participants’ cortisol level from a hair sample. The advantage of measuring cortisol in a hair sample compared with blood or saliva is that it provides an aggregated long-term measure. Specifically, a hair sample of 1 cm length and 3 mm in diameter will be taken close to the posterior vertex. This allows for evaluating the cumulative cortisol level of approximately the last 4 weeks. The hair samples will be sent to the laboratory of Professor Dr Kirschbaum at the Technical University of Dresden, Germany, to determine the level of cortisol.

Moreover, affect and stress will be assessed on a behavioural level by recording participants’ mouse movement and keyboard use in the online questionnaire. This exploratory measure is based on preliminary results supporting the rationale that emotional states influence sensorimotor and cognitive processes causing measurable changes in our interaction with computer input devices such as the keyboard and mouse usage.

Keyboard and mouse usage behaviour will be captured by participants’ web browser using a JavaScript script embedded in the online questionnaire. The script will track the mouse cursor’s x/y coordinates on the screen as well as each keyboard key press and key release at millisecond precision.

Implicit measures

In contrast to measuring attitude and emotional state using self-report questionnaires, implicit measures do not require introspection and thus provide a different level of tapping the effects of the intervention.

Attitude towards the job will be assessed using the single category implicit association test (SC-IAT). In the SC-IAT, words are displayed in the centre of the screen one at a time together with categorising words for ‘positive’ and ‘negative’ as well as the category ‘job as physician’ on the sides of the screen. Participants need to categorise the words that appear in the centre of the screen. In one block the categorising labels for ‘positive’ and ‘job as physician’ share one key and in a second block labels for ‘negative’ and ‘job as physician’ share a key. Reaction times will be recorded. Shorter reaction times when ‘job as physician’ is paired with positive in contrast to when it is paired with negative indicate a positive attitude towards the job and vice versa.
Furthermore, attitude towards the job will be assessed using the affect misattribution procedure (AMP). In the AMP, participants are presented with briefly appearing pictorial primes referring to the job as physician followed by neutral Chinese pictograms. Participants are instructed to rate how visually pleasant the Chinese pictograms are, giving their responses as fast as possible. It is assumed that the ratings of the Chinese pictograms reflect participants’ attitude towards their job as physician.

Projective measures
Similar to implicit measures, projective tests do not require introspection. Emotional state will be assessed with the Implicit Positive and Negative Affect Test (IPANAT). In the IPANAT participants are presented with six artificial non-sense words (SAFME, VIKES, TUNBA, TALEP, BELNI and SUKOV) and are asked to rate to what extent these words express positive emotional states (happy, cheerful and energetic) and negative emotional states (tense, inhibited and helpless). The positivity or negativity of the ratings reflects participants trait affect.

Additionally, emotional state will be assessed by the word fragment test (WFT). The WFT requires participants to complete a series of word fragments. These fragments are constructed in a way that they can be completed into at least two words that are either associated with positive or negative affect. For example, the word fragment ‘ange_’ can be completed into ‘angel’ or ‘anger’. This test is based on the assumption that depending on the current affective state, either positive or negative words are more accessible and are thus more likely to be generated. Consequently, the proportion of positive and negative words that are generated by participants allows for inferring their affect.

Self-esteem will be assessed using the name-liking task. The name-liking task is a single-item measure that requires participants to rate how much they like their full name. The name-liking task has been derived from the affective or negative affect. For example, the word fragment test (WFT). The WFT requires participants to complete a series of word fragments. These fragments are constructed in a way that they can be completed into at least two words that are either associated with positive or negative affect. For example, the word fragment ‘ange_’ can be completed into ‘angel’ or ‘anger’. This test is based on the assumption that depending on the current affective state, either positive or negative words are more accessible and are thus more likely to be generated. Consequently, the proportion of positive and negative words that are generated by participants allows for inferring their affect.

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descriptions of the course sessions in the instructor’s guide. Moreover, treatment fidelity will be assessed by videotaping two random course sessions that will be rated by two independent raters using the Mindfulness-Based Interventions Teaching Assessment Criteria.97

Procedure
Residents will enrol for the study by email or via the study’s webpage. Eligible residents will be invited to a first on-site appointment, where they will receive further information about the study and will be asked to give their informed consent. After baseline assessment, participants will be assigned to one of the two groups using the minimisation procedure described above. Participants will receive educational points for participating in the mindfulness course but no other incentives will be given. Participants will be invited to take part in the assessments via email. Assessment at each measurement point consists of an online questionnaire98 as well as an on-site appointment. The on-site appointment consists of the collection of the hair sample and the GAS at t0 and t1, while the online questionnaire contains the self-report questionnaires, the implicit and projective tests as well as the recording of mouse and key use. Furthermore, participants of the first three courses and respective control group (ie, 84 residents) will be invited to take part in a qualitative interview, which will also be held on-site. Theoretical sampling is not possible because this study is fully anonymised, meaning that we are unable to link the names of participants to their identifying code and data. However, retrospectively we will be able to determine whether the subsample of residents that took part in the interviews covers a range of typical attributes that are of interest from a theoretical perspective such as experienced stress level and experienced effect of the intervention. Together with the criteria of theoretical saturation, this will determine whether we will invite additional participants to the interview or not. To promote participant retention for the follow-up assessments, participants will be contacted and reminded by email to take part in the assessments.

Procedures to reduce bias
Due to the nature of the study, neither the participants nor the trainers running the mindfulness courses can be completely blinded. However, to minimise bias, the self-report questionnaires as well as the implicit assessments will be conducted using an online survey. Furthermore, all on-site assessments (ie, taking the hair samples) that are carried out after participants have been assigned to a group will be conducted by assessors that will be blind to group allocation. Additionally, participants will be asked not to tell the assessor which group they have been allocated to. An exception to this procedure will be the qualitative interviews that are held with a subsample of participants, and will be carried out by a researcher who will not carry out other assessments.

Data protection
Regarding data protection, all data will be stored under an anonymous code that can be linked to data only by the participants themselves. An exception to this is the short interval between baseline assessment and allocation to group during which a temporary list linking name and anonymous code will be maintained, which allows for communicating to participants to which group they have been allocated. After allocation has taken place, this temporary list will be destroyed.

The hair samples will be sent to a laboratory in Dresden, Germany, (Technical University Dresden, Germany) that will destroy the hair samples after the cortisol level has been obtained.

There will be no formal data monitoring committee since there is no known risk for the intervention and the trial includes no patients. To ensure quality of data, all data that have not been collected via the online survey will undergo a double data entry.

Calculation of sample size
Previous studies that applied a similar design found effects of a mindfulness intervention on mental health variables of around d=0.45.30 31 Accordingly, to find a significant group difference at t2, assuming a power of 1-beta=0.80 and an alpha level of 0.05 (one-tailed) the sample size would have to be at least 62 participants per group. Adding customary drop-out rate of 30 per cent results in a planned sample of 89 participants per group or 178 participants in total. Estimation of sample size was conducted using the software G*Power.99 For the purpose of sample size calculation we used a simple group comparison; however, in the actual analysis we will use a linear mixed model, which by nature has higher power.

Planned analysis
Quantitative data
The analyses will be conducted on an intention-to-treat principle. The primary outcome will be analysed using linear mixed modelling comparing the respective outcome measure between participants of the intervention group and participants of the control group. Participants will be analysed as randomised. In case of drop-out and non-adherence, this subgroup’s baseline values in the primary outcome will be compared with the other participants. We expect that these groups do not differ significantly. Furthermore, the pattern of missing values will be analysed. By conducting linear mixed models, missing data will be dealt with by using maximum likelihood to obtain estimates of missing parameters.

Effects of the intervention will be examined by looking at the interaction of time and group by entering group as fixed factor and assuming a random intercept within the linear mixed model. Additionally, time will be entered as a random effect, that is, allowing for random slopes if this would improve model fit. The hierarchical structure of the data due to the organisation of participants into different courses, taught by different teachers will be

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taken into account by specifying course as a separate level. Similar models will be calculated for secondary outcomes.

**Qualitative data**

Audio recordings of the interviews will be transcribed applying the transcription method by Dresing and Pehl. These transcripts will be analysed using thematic analysis, following the six-step process proposed by Braun and Clarke. In a first step, we will familiarise ourselves with the material by repeatedly reading the transcripts to gain an understanding of the scope and depth of the interviews. In a second step, initial codes will be generated by identifying meaningful groups within the interview data. In a third step, the generated codes will be organised into overarching themes. In a fourth step, these themes will be reviewed by reassessing whether the themes adequately represent the codes and the data as a whole. In a fifth step, the identified themes will be labelled and described in detail with the aim of capturing the essence of each theme. The sixth and last step consists of the final analysis, choosing of representative text excerpts and writing down the results. Steps one to five will each be done individually by two researchers, followed by a comparison and discussion of the results in groups with at least three researchers. Data transcription, processing and analysis will be carried out using the software MAXQDA.

**Patient and public involvement**

The intervention aims at improving resident physicians’ mental health and does not address patients directly. However, there is evidence that physicians’ mental health fosters the quality of care they deliver. Thus, there may be an indirect benefit for patients. Furthermore, patients are involved in the study directly within the third party reports. Specifically, three patients of every resident will be asked to complete a short questionnaire about the respective resident.

There is a study webpage informing the public about the study.

**Ethics approval**

The planned study was approved by the Institutional Ethics Committee of the Albert-Ludwigs University of Freiburg (Reference number: 361/16) and is funded by the German Research Foundation as part of the interdisciplinary collaborative research center ‘SFB Muße 1015’. The trial was registered at the German Clinical Trials Register of Clinical Studies (drks.de). The collection of data started in September 2018. Changes to the study protocol will be reported to the ethics committee and will be updated in the trial registry of the DRKS. Only participants that have given their written informed consent will be included in the study.

**Dissemination plans**

The results of this study will be published in scientific journals and disseminated through the study’s website. Further, participants will receive the results of the hair cortisol analysis after completion of the final assessment at t3.

**DISCUSSION**

Residents show high prevalence of mental distress including burnout, perceived stress, depression, and anxiety. This comes at a price of both inferior patient treatment and increased economic costs. That is why there is a need for interventions that address this problem. The aim of the present work is to examine the effects of a mindfulness intervention that has been tailored to residents’ needs. If results are positive, this study will demonstrate evidence of an intervention through which residents can learn how to cope with stress and improve their mental health. This could further imply that such programmes should become an integral part of occupational health management models in hospitals since there is a lack of programmes in which residents can learn such competencies. Dealing with stressors at work, distress, time pressure and the downsides of economy-driven healthcare systems are crucial competences that have not been part of the formal curriculum so far.

However, such a MBI may run the risk of being functionalised to make residents more stress-resistant in order to engage in either more self-sacrificing or self-optimising behaviour at work. By supplementing mindfulness with a focus on Muße, we aim to make mindfulness practice a counter-pole to stress and functioning that addresses the mental health of residents. The integration of the concept of Muße into the MBI follows a recently emerging line of second-generation MBIs that are characterised by the combination of mindfulness with other principles such as ethical awareness or compassion.

The present study will have several limitations. First, the lack of a third group that receives standard MBSR will not allow for determining whether a mindfulness course tailored to residents’ needs is superior to a standard mindfulness programme. Therefore, if this trial is successful, further studies should include such a comparison in their design. Second, the control group will receive text-based information about mindfulness. It is intended that participants allocated to this group will read on a weekly basis for the duration of 8 weeks, paralleling the structure of the intervention group in terms of information. However, we are not able to control for whether the material will actually be read and comprehended with the same intensity than in the course. Third, participants will be a self-selected sample of residents instead of a random sample. Yet, the strength of this study is its ecological validity by assessing the effectiveness of the mindfulness intervention in those residents who actually seek such an intervention. Furthermore, the likely effects of such an intervention depend on the motivation of the participants and such an intervention cannot be administered passively like in a drug trial. Fourth, external evaluations might be biased because residents may tend to give the questionnaires to patients where treatment was successful and colleagues
and supervisors where the relationship is positive. However, this does not present a threat to internal validity as this practice will be present in both the intervention and the control group. Fifth, the mindfulness courses will be held at one hospital only, which might limit the generalisability of the results to this specific context. However, residents will be recruited from a wide range of hospitals including university hospitals, clinics in private as well as public ownership, church-funded hospitals, as well as hospitals in both urban and rural areas. As a result, we expect a diverse sample that generalises to the hospital landscape in Germany. Last, a part of the measurements will consist of self-report scales that are known to be susceptible to participants’ expectation of intervention effects, vagueness in evaluating their mental state and/or social desirability. For this reasons, self-report scales are supplemented by other means of assessment that are less susceptible to these pitfalls including hair cortisol and implicit measurements.

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Contributors SS and ASG developed the initial study concept and obtained funding. All authors contributed to the design of the study and were involved in the development of the adapted mindfulness intervention. VMA drafted the study protocol while JCF, ASG and SS made important revisions. All authors have read and approved the manuscript.

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