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VITALITY, RESILIENCE AND THE NEED FOR SUPPORT AMONG HOSPITAL EMPLOYEES DURING THE CONTINUATION OF THE COVID-19 PANDEMIC: A STUDY PROTOCOL

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VITALITY, RESILIENCE AND THE NEED FOR SUPPORT AMONG HOSPITAL EMPLOYEES DURING THE

CONTINUATION OF THE COVID-19 PANDEMIC: A STUDY PROTOCOL

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Abstract

Introduction

The Coronavirus Disease 2019 (COVID-19) pandemic has a significant impact on the physical and mental functioning of healthcare professionals, especially those working on the 'frontline', and other hospital workers. At the onset of the crisis, various interventions were introduced to promote resilience and offer mental support to these professionals. However, it is unknown whether the interventions will meet the needs of professionals as the COVID-19 pandemic continues.

The goal of the intended study is to gain insight in factors that protect the vitality and resilience of hospital employees during the so-called 'second wave' of the COVID-19 pandemic. This paper describes the study protocol.

Methods and analysis

This study applies a mixed-methods design, using both quantitative and qualitative methods of data collection and analysis. The first part of the study (sub-study I) consists of surveys among doctors and nurses in COVID-19 departments and non-COVID-19 departments, and other professionals in the hospital (i.e., managers and homeworkers) in 2020 and 2021. The second part of the study (sub-study II) consist of focus groups and interviews among professionals of the intensive care unit, COVID-19 departments and infection prevention units.

Ethics and dissemination

The research protocol for this study has been approved by the Medical Ethics Committee (MEC-2020-0705). Professionals with vitality experience less work-related stress and can therefore handle

more work in the new and stressful circumstance. In other words, maintaining professionals' vitality and resilience will contribution to healthcare quality. The outcomes of this study will be used to develop and implement interventions to support hospital employees in maintaining their vitality and resilience during and after the COVID-19 pandemic.

Article summary

Strengths and limitations of this study

- As the COVID-19 pandemic continues, it will be necessary for organizations to maintain professionals' vitality and resilience, as more effort is expected from the professionals and they will be confronted with new and stressful circumstances.
- Frontline workers from different departments managers and homeworkers will be compared in contrast to the majority of studies so far which focused exclusively on the needs of healthcare professionals.
- Real life data started during the beginning of 2nd COVID-19 wave, ongoing to autumn 2021.
- The COVID-19 pandemic is the motivation for this study, but may also limit the response rates or procedure of this study, given its unpredictable course.

Key words:

COVID-19, healthcare professionals, mental support, needs assessment, resilience, vitality.

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Introduction

Worldwide, it has been reported that the Coronavirus Disease 2019 (COVID-19) pandemic has a significant impact on the physical and mental functioning of healthcare professionals, especially for those working on the 'frontline' (e.g., intensive care units (ICUs), COVID-19 departments and infection prevention units), and other hospital workers[1-4]. Indeed, also in the Netherlands, the COVID-19 pandemic had an impact on medical professionals. This is critical, as it has been reported that some Dutch medical professionalsweare already overburdened before the COVID-19 pandemic[5, 6].

The need for high-intensity medical treatment rapidly increased during the COVID-19 pandemic, during which the work circumstances became uncertain and stressful[7]. Work circumstances involved the continuous use of personal protective equipment, adapted responsibilities and tasks, moral dilemmas, and the risk of contamination for the healthcare professionals themselves[8]. Interpersonal contact with patients' family members, one of the core features of the professional practice of nurses, was considerable reduced due to visiting limitations in most hospitals [9, 10]. In addition, the work environment also changed for ICU nurses as their teams changed due to help from (former) colleagues and other healthcare professionals. This sudden shift in activities and responsibilities required ICU nurses to have additional competences to maintain high-quality healthcare. Buddies, or support staff from other departments in the hospital, were sometimes confronted with distressing or even shocking events during the first hectic weeks of the pandemic. Professionals of the infection prevention unit had to deal with an enormous workload due to the accumulation of new tasks and changing work processes under enormous time pressure, as well as the social turbulence resulting from the implemented quarantine measures. In the case of a health crisis such as the continuing COVID-19 outbreaks, the health and vitality of the frontline professionals becomes even more critical than in normal circumstances. This is because the higher workload and stress will do a higher appeal on the physical and mental resources of the

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professionals. The COVID-19 pandemic had not only impact on the clinicians of the hospital. The work environment also changed for non-clinical professionals who suddenly had to do all administrative work and communication from home. In addition to this loss of the work environment and direct contact with colleagues, homeworkers might lack a sense of purpose, solidarity and valuable contribution to the crisis situation[11]. Last, the COVID-19 pandemic required great effort from managers[12]. More than ever, they had to deal with logistic and administrative processes in the upscaling of high-intensity care, improving work alliances and the integration of staff in newly formed teams, and in managing the continuous flow of changing information.

Health, vitality and resilience

In previous virus outbreaks, such as the outbreaks of SARS, Ebola and MERS, it became clear that increased stress levels at work in healthcare professionals were associated with fear of contamination, shortage of materials, poor communication between healthcare professionals, unclear work instructions and information, deficient or non-functioning equipment, and inadequate planning among healthcare professionals[13-16]. Experiences from China during the COVID-19 pandemic showed similar results[17-19]. In a European study after COVID-19 on work-related stress reactions among ICU healthcare professionals half of the respondents (50.4%) showed symptoms of anxiety[1]. Early phase evidence on COVID-19 suggested that healthcare professionals experienced mood and sleep disturbances during this outbreak, stressing the need to establish ways to minimize mental health risks and support interventions aiming at pandemic conditions[3]. In the short-term, this work-related stress can cause fatigue, sleep disorders, mistakes and moral distress[20]. Longterm effects of high work pressure include burnout, depression and post-traumatic stress, resulting in dropout due to illness and abandonment of paid employment[21, 22]. A recent study in the Netherlands on burnout rates among intensivists were reported to be moderate (14.8%)[23]. Furthermore, recovery time - regaining strength after an intensive period at work- is associated with physical and mental well-being[24]. A long recovery time is an early indicator of work-related stress

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and exhaustion[25]. In contrast to high workload, stress and recovery time, vitality, resilience and job satisfaction are characteristics of professionals that counterbalance work-related stress[26, 27]. These characteristics strengthen professionals' mental and physical well-being and their retention for work[28-30]. Professionals with vitality are more resistant to work pressure.

Interventions among healthcare professionals during the COVID-19 pandemic

A wide variety of studies have examined interventions to reduce the work-related stress of healthcare professionals during the COVID-19 pandemic. Providing personal protective equipment is the top priority, followed by fulfilling the psychological needs of professionals[31]. To support mental health and promote the vitality of healthcare professionals, various interventions, including buddy systems, peer support, coaching and easily accessible psychological help, were proposed during the COVID-19 period from March to May 2020[7, 32-36]. Other individual interventions, such as telemedicine activities, e-package and self-help books, appear to be promising[37-40]. For example, a hospital in China offered online courses to help medical professionals to deal with psychological problems[41]. Many interventions have taken an individual approach, but system-level changes in healthcare organizations seems to have a wider reach than individual support[42]. A notable omission in the literature is that protective factors are given limited attention: the focus is on the stressors. So there seems to be many possible interventions to support professionals in times of a pandemic, however, it is not clear which intervention matches the needs of the professional most closely. Therefore, we set out to investigate which supportive interventions, system changes and other supportive factors could meet individual needs during and in the aftermath of the COVID-19 pandemic in a large academic hospital in the Netherlands.

Objectives

The overall goal of the study is to gain insight into the risk and protective factors as well as the needs and barriers in the working environment related to the promotion of the vitality and resilience of

employees. Our objective is to assess levels of vitality and resilience, and the need for support or resources among professionals with a focus on professionals working in ICUs, COVID-19 departments and infection prevention units. Furthermore, to gain more insight into the relationship of vitality and resilience with factors such as self-perceived health, stress, burnout, posttraumatic stress, and need for recovery. Based on the results of this study we aim to formulate recommendations for interventions aiming at increased vitality and resilience for healthcare professionals in the organization.

Methods and analysis

Study design

A mixed-methods design, using both quantitative (Sub-study-I) and qualitative methods (Sub-study-II), is applied. Sub-study I is a cross-sectional online survey administered first in October 2020, when the second wave of the COVID-19 pandemic was upcoming and ongoing, followed with measurements in March and September 2021. Sub-study II includes focus group interviews among nurses, doctors, and professionals regarding the ICU, COVID-19 departments and the infection prevention unit during the end of 2020. The Spirit checklist was used to finalize reporting of the study protocol in detail (Supplemental file S1).

Setting

The study setting is a large academic hospital in the Netherlands.

Study population

<u>Sub-study I</u>: The population consists of a random sample based on voluntary participation of four target groups: professionals working at the COVID-19 department, non-COVID-19 departments, managers and homeworkers.

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<u>Sub-study II</u>: The population consists of a non-random sample of professionals working in the ICU, COVID-19 department and the infection prevention unit. Participants are invited and selected in collaboration with the managers of the study population.

The inclusion criteria for the entire study are (i) a minimum age of eighteen years and (ii) sufficient Dutch language proficiency to complete the questionnaires or to discuss the relevant topic.

Patient and Public Involvement

No patient involved.

Study procedures

Sub-study I: Online surveys

Professionals are informed about the study in several ways. The communication strategy is tailored to each target group and supported by the communication department of the organization. A link to the online survey is published on the intranet of the organization, printed QR-codes containing a link to the survey are available at the coffee corners and canteens, announcements are made in the weekly COVID-19 livestream and by team management via personal email. Participation is voluntary and can be performed during working hours.

The online questionnaire starts with information about the study, privacy statements and a consent form for participation. After providing consent to participate, participants are asked to fill out the entire questionnaire, which consists of two parts. The first part is generic for all employees and takes approximately six minutes to complete; it includes questions on demographic information and the main outcomes. The second part consists of additional modules on working conditions and health and has a completion time of approximately seven minutes. Nurses and homeworkers receive another additional module tailored to their specific work environment.

Sub-study II: Focus groups

In total, six focus groups with 6-10 participants that take approximately 60 minutes are conducted. ICU doctors, ICU nurses, microbiologists, hospital hygienists, COVID-19 unit nurses, and COVID19unit doctors (lung specialists and specialists internal medicine) are individually invited to participate in one of the focus groups through consultation with the team managers. These meetings are preferably in-person (to observe non-verbal attitude and facial expressions), but due to the COVID-19 measures and social distancing, it may not be possible for participants to be physically present. In those cases, the focus groups are carried out via video calling technology.

Prior to the meetings, a topic list is created by the research group based on the literature and internal reports on the experiences of professionals. This topic list is used to guide and structure the meeting. The aim of the focus group is to study protective factors that contribute to vitality and resilience during the COVID-19 pandemic. Furthermore, possible interventions to increase vitality and resilience are explored and elaborated upon. Written informed consent is given prior to the meeting, and two experienced researchers guide the meetings. The focus group interviews are recorded and transcribed verbatim.

Measurements

This paragraph lists all measurement instruments included in the questionnaire. The first part consists of measuring instruments addressing demographics, primary outcomes (i.e. vitality, resilience and needs assessment), and several secondary outcomes (i.e. self-perceived health, stress, burnout, posttraumatic stress, and need for recovery). The second part consists of separate modules for homeworkers and nurses with regard to work ability, working conditions, job satisfaction, workprivate balance, exposure to COVID-19 at work, preventive measures for COVID-19 and career perspectives.

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Demographics

Gender, age (divided into five categories), educational level, function at work, work location and professionals' experience (in years) are assessed. Educational level is divided into three levels: low, medium and high educational level. In total, the list of functions includes 23 positions within the academic hospital (e.g., nurse, ICT employee, pharmacist, educator, doctor).

Main outcome measures

Vitality

Vitality is measured with four items from the original 36-item Short Form Health Survey (SF-36) [43]. The total summed score of four items that refer to the past four weeks: "Did you feel full of liveliness?", "Did you have a lot of energy?", "Did you feel worn out?", and "Did you feel tired?". The answers are rated on a six-point scale from 1 (= constantly) to 6 (= never) [44]. Higher scores indicating a better subjective vitality.

Resilience

Resilience (the ability to cope with stress, setbacks or difficulties at work) is measured with six items from the Psychological Capital Questionnaire[45]. The items contain statements such as: "When I have a setback at work, I have a hard time getting back on track and moving on", "If necessary, I can work well without the help of others" and "I can handle difficult moments at work". The six items are scored from 1 (= strong disagreement) to 6 (= strong agreement). Higher values indicate a higher level of resilience.

Needs assessment

Needs are measured with a self-designed scale with four items. Examples of questions are: "In which area would you like to be supported?" and "What would this support look like?" and "What should

be offered or developed?". A predefined list includes 10 individual- and 14 organizational-related answer options, e.g., support for working from home, time management, and work-private balance.

Other outcome measures

Self-perceived health

Self-rated health is assessed with one question ' "In general, how would you say your health is?" Responses range from 1 (= excellent) to 5 (= poor).

Stress

Stress is measured with a numeric rating scale. The stress score, ranging from 0 (= no stress at all) to 100 (= the worst stress imaginable). This scale is used to retrospectively objectify stress before, during and after the first COVID-19 outbreaks. The three item question was "How did you experience the stress before/during/after the COVID-19 crisis on a scale from 0 to 10?"

Burnout

Burnout is measured using five items, that are based on an adapted version of the Utrecht Burnout Scale [46]. The items refer to the current situation such as "I feel emotionally drained from my job" and "I feel completely exhausted from my work". The responses range from 1 (= never) to 7 (= daily).

Posttraumatic stress

Posttraumatic stress is assessed with the Posttraumatic stress disorder (PTSD) Checklist for the DSM-5 (PCL-5) - COVID-19 version with 20 items[47]. This scale consists of 20 items, measuring PTSD symptoms, with scoring options from 0 (= not all) to 4 (= extremely) and was adapted to the COVID-19 situation. A score of 33 or higher is preceived indicative for PTSD.

Need for recovery

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Work fatigue and the risk of psychological symptoms are measured using the Dutch questionnaire on the Experience and Evaluation of Work (Dutch abbreviation: VBBA)[48, 49]. The need for recovery scale consists of eleven dichotomous items (yes/no), representing short-term effects of a working day[24, 50, 51]. The score of the need for recovery scale ranges from 0 to 100 and is calculated as the sum of points (1 = yes, 0 = no) divided by the number of questions answered, multiplied by 100. Higher scores indicate a higher need for recovery, which is unfavourable.

Work ability

Work ability is measured with the Work Ability Index (WAI)[52]. This widely used index measures self-assessed work ability and consists of seven items. Because the sub-items of the WAI can also be used as a simple indicator for work ability[53], three of the seven items are used: current work ability (one item), and work ability in relation to physical and mental job demands (two items). A total WAI score (range 2–20) is obtained by adding the weight scores of these individual items[54].

Working conditions

Aspects of working conditions measured in this study are: job autonomy, emotional job demands, social support and physical working conditions.

Job autonomy is measured with six items on a three point scale (no; yes, sometimes; yes, regularly). Five items, i.e., those about making decisions, having to find solutions, and being able to take time off, are based on the Job Content Questionnaire[55, 56]. One item on autonomy related to working time based on the Netherlands Working Conditions Survey, is also included in the questionnaire[57].

Emotional job demands is evaluated with four items. Three items are derived from the Copenhagen Psychosocial Questionnaire and assess whether the work leads to emotionally difficult situations, the emotional demands of the job, and emotional involvement in work. An additional item is "Is your

job more emotionally demanding because of COVID-19?". All items are measured on a four-point scale (never to always)[58].

Social support is defined as whether colleagues and supervisors are willing to help and listen to work-related problems and is assessed using four items from COPSOQ[58]. Social support is measured on four-point Likert scales 1 (= almost never) to five (= always).

Physical working conditions are measured with one self-designed question and assesses whether a worker received more or less physically demanding work due to COVID-19 measures. This scale has three answer options (no; yes, sometimes; yes, regularly).

Job satisfaction

Job satisfaction is measured with one item: "Altogether, how satisfied are you with your work?" The responses range from 1 (very dissatisfied) to 5 (very satisfied).

Work-private life balance

Work-private life balance is measured with two questions on the mutual interference between work and home life. The questions are adopted from the Netherlands Working Conditions Survey [57], but were originally constructed by Fox and Dwyer (1999) [59]. Both questions have four answer options ranging from 1 (= no, never) to 4 (= very often).

Exposure to COVID-19 at work

Professionals are asked to what extent they might have been exposed to COVID-19 at the worksite. These questions are derived from the Netherlands Working Conditions Survey COVID-19 [60], based on questionnaires developed within the OMEGAnetwork[61]. Participants are aked if they work with patients, the average number of patients they worked with during a typical working day in the last

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week, and if these patients were suspected to have or had been diagnosed with COVID-19. Additionally, participants evaluate their work on a regular basis with colleagues, the number of colleagues they work with and if they share tools or surfaces with them.

Preventive measures for COVID-19

The questions on preventive measures in the workplace are from the Netherlands Working Conditions Survey COVID-19 [60] and consist of one general question and five more specific questions. The general question assesses the general measures at the department level with regard to the COVID-19 pandemic, with answer options such as homeworking, adjustment of working hours, general preventive measures in the workplace, mandatory inclusion or withdrawal of leave. The specific questions are about the possibility of keeping a 1.5 metre distance between colleagues and/or patients, the availability of personal protective equipment, the usage of personal protective equipment and the application of general hygiene measures. The responses to these five questions are never, sometimes, often and always. This module will not be applied to professionals working from home.

Career perspective

Three items on career perspective are derived from the Netherlands Working Conditions Survey COVID-19[60] and adjusted to fit the study population working in the hospital. These items include the motivation to work in the healthcare sector in the future (responses: less, equal, and more), the intention to change jobs within the health care sector and the intention to change jobs outside the healthcare sector with responses ranging from 1 (= certainly not) to 5 (=certainly yes).

Outcome measures for pre-defined groups or professions

Nurse questionnaire

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The Practice Environment Scale of the Nursing Work Index is the most widely used measure to gauge the state of nursing practice environments[62, 63]. It is the only measure recommended by several organizations promoting quality healthcare. The 15-item questionnaire uses responses ranging from 1 (= strongly disagree) to 4 (= totally agree). This module will be applied to nurses only.

Homeworkers

A total of eight items are specifically tailored to homeworkers. Two items refer to the number of hours in a week people work from home and how many hours a day they work on a screen (e.g., laptop and tablet). One item is focused on the availability of ergonomic work equipment at home (a desk or table with a comfortable working height, a chair that can be adjusted to one's body measurements, a separate display, and a separate computer mouse). The need for other furniture is assessed with one item "Do you need additional materials for a good home workplace?". The answer options yes or no. Moreover, regarding rest breaks – outside the lunch break – are asked "Do you take (short) breaks on a working day, except for a lunch break?". This question includes the following response options: 1 (= yes, regularly), 2 (= yes, sometimes) and 3 (= no). The last three items are about concentration while at home and include the following statements: "Do you have trouble concentrating while working?", "Do you struggle to keep your attention while you work?" and "Do you have difficulty with the reduced social contact with colleagues?" Answers range from 1 (= never) to 4 (- always).

Data handling and statistical analyses

<u>Sub study I</u>: survey data are anonymously collected using Limesurvey (Version 2.06 Its Build 160524) and exported to a secure SPSS database (©IBM SPSS Statistics for Windows, Version 25.0. Armonk, NY: IBM Corp) for analysis. All principal investigators have access to the final study dataset. Data will be stored for fifteen years.

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First, the data are cleaned and checked for missing data. The descriptive statistics are presented as numbers and percentages with a normal distribution around the mean (with standard deviation) for dichotomous variables and a non-normal distribution around the median (with interquartile range) for continuous variables. Data for different subgroups (professionals in COVID-19 departments, non-COVID-19 departments, executives and homeworkers) are analyzed with the Mann-Whitney test or t-tests. Linear and logistics regression analyses are preformed to investigate the associations between risk factors and the main outcomes (vitality and resilience). Statistical significance will be defined as p < .05.

Sub study II: Focus group interview data are audiotaped and transcribed verbatim. Thematic analysis is conducted the principles of thematic content analysis [64]. Two researchers read the transcripts. Each of them develops a structured analysis framework that consists of preliminary themes and codes. After that, they compare their frameworks to reach consensus. Next, one researcher codes the transcripts line by line according to this framework in the software programme NVivo12[®]. The coder uses memos for comments during coding. When coding is finished and the code 'other' is used, the two researchers discuss these codes and rename them into a new or existing codename best reflecting the contents of the otherwise uncategorised text fragment. After coding is finished, the cohesion and inter-relations between codes are analysed by the two researchers. The principal investigators have access to these data, and data will be stored for fifteen years.

Ethics and dissemination

The study is approved by the Medical Ethics Committee of the Erasmus MC (MEC-2020-0705) and conducted according to the principles of the Declaration of Helsinki (64th WMA General Assembly, Fortaleza, Brazil, October 2013) and in accordance with the Medical Research Involving Human Subjects Act. The study complies with the Netherlands Code of Conduct for Scientific Practice from the Association of Universities in the Netherlands. Protocol modifications will be communicated and

to the Medical Ethics Committee by protocol amendment. Participants will be informed about the study both orally and by letter. Consent for participation will be given by written informed consent. Participants can leave the study at any time for any reason if they wish to do so without any consequences. The withdrawal will be registered for informative purpose.

The consequences of the COVID-19 crisis on the mental health and working conditions of healthcare professionals have been recognized worldwide[65]. By using a mixed-methods approach, we aim to gain an overview of vitality, resilience and health (e.g. stress and burnout) among healthcare professionals, as well as the risk factors associated with these outcomes. This is an urgent and rushed study because we want to use the results against the same health crisis that we are investigating. Based on this study, directions for future interventions during the COVID-19 pandemic and thereafter can be provided to improve the vitality and resilience of professionals in the hospital, and therewith support their employability.

Strengths and limitations

The first strength is the mixed-methods design, consisting of qualitative and quantitative methods . Second, we compare different departments and distinguish executives and homeworkers. The majority of studies so far focused exclusively on the needs of healthcare professionals without considering other hospital employees such as supportive staff, researchers and managers. Third, real life data gathering during start of 2nd wave.

The COVID-19 pandemic was the motivation for this research, but may also have limited the procedure of this study, given its unpredictable course. During the writing of this protocol paper, the second wave of COVID-19 had already started in The Netherlands. Therefore, a lower response rate is not unexpected. The second limitation is the cross-sectional design of the study, which makes it impossible to draw causal conclusions from this report.

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Data dissemination

Public access to the study protocol, study details, participant-level dataset, and statistical code can be acquired from the corresponding author. The results will be disseminated to healthcare professionals, health services authorities and the public via presentations at national and international meetings and published in peer-reviewed journals. A lay summary of the results will be written and shared with all professionals of the organization.

Study status

The study is currently ongoing with data recruitment.

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Declarations

Authors contributions

MvM, TK, JB and LK jointly designed the study, raised funding and established the development of the study protocol. MvM, MV, TK, KOH and LK prepared the study materials. MvM, MV and LK gathered the data of both sub-studies and produced the first draft of the article outline together with KOH and TK. All authors (MvM, MV, AP, TK, JB, WH, KOH, LK) contributed substantially to the concept of the study, the analyses of literature, critically revised the content of the manuscript, have read and approved the final version.

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Competing interest statement

The authors declare no conflicts of interest.

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Availability of data and materials

Anonymized data gathered and analysed during the current study are not publicly available due to legal and ethical restriction. These can be requested from the corresponding author as well as text

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and photo material of the developed intervention. Materials described in the manuscript, including all relevant raw data, will be freely available at a reasonable request to any scientist wishing to use them for non-commercial purposes.

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1 2 3 4 5 6			STANDARD PROTOCOL ITEMS: RECOMMENDATIONS FOR INTERVENTIONAL TRIALS					
7 8	SPIRIT 2013 Checklist: Recommended items to address in a clinical trial protocol and related documents*							
9 10 11	Section/item	ltem No	Description	Addressed on page number				
12 13 14								
15 16	Title	1	Descriptive title identifying the study design, population, interventions, and, if applicable, trial acronym	1				
17 18	Trial registration	2a	Trial identifier and registry name. If not yet registered, name of intended registry	NA				
19 20		2b	All items from the World Health Organization Trial Registration Data Set	NA				
21 22 23 24	Protocol version	3	Date and version identifier	3				
	Funding	4	Sources and types of financial, material, and other support	2				
25 26	Roles and responsibilities	5a	Names, affiliations, and roles of protocol contributors	1,2 and 28				
27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46		5b	Name and contact information for the trial sponsor	2, corresponding author				
		5c	Role of study sponsor and funders, if any, in study design; collection, management, agalysis, and interpretation of data; writing of the report; and the decision to submit the report for publication, including whether they will have ultimate authority over any of these activities	28				
		5d	Composition, roles, and responsibilities of the coordinating centre, steering committee, endpoint adjudication committee, data management team, and other individuals or groups over seeing the trial, if applicable (see Item 21a for data monitoring committee)	NA				
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			BMJ Open Den		
1 2	Introduction		2021-6		
3 4 5	Background and rationale	6a	Description of research question and justification for undertaking the trial, including sugnmary of relevant studies (published and unpublished) examining benefits and harms for each intervention	6-8	
6 7		6b	Explanation for choice of comparators	NA	
8 9	Objectives	7	Specific objectives or hypotheses	8	
10 11 12 13	Trial design	8	Description of trial design including type of trial (eg, parallel group, crossover, factorial single group), allocation ratio, and framework (eg, superiority, equivalence, noninferiority, explorator \overline{y})	NA	
14 15	Methods: Participants, interventions, and outcomes				
16 17 18	Study setting	9	Description of study settings (eg, community clinic, academic hospital) and list of courtings where data will be collected. Reference to where list of study sites can be obtained	9	
19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46	Eligibility criteria	10	Inclusion and exclusion criteria for participants. If applicable, eligibility criteria for study centres and individuals who will perform the interventions (eg, surgeons, psychotherapists)	9	
	Interventions	11a	Interventions for each group with sufficient detail to allow replication, including how and when they will be administered	NA	
		11b	Criteria for discontinuing or modifying allocated interventions for a given trial participa f t (eg, drug dose change in response to harms, participant request, or improving/worsening disease) $\frac{2}{2}$	NA	
		11c	Strategies to improve adherence to intervention protocols, and any procedures for m_{2}^{N} itoring adherence (eg, drug tablet return, laboratory tests)	NA	
		11d	Relevant concomitant care and interventions that are permitted or prohibited during the trial	NA	
	Outcomes	12	Primary, secondary, and other outcomes, including the specific measurement variable (eg, systolic blood pressure), analysis metric (eg, change from baseline, final value, time to event), method of aggregation (eg, median, proportion), and time point for each outcome. Explanation of the clinical relevance of chosen efficacy and harm outcomes is strongly recommended	11-17	
	Participant timeline	13	Time schedule of enrolment, interventions (including any run-ins and washouts), assessments, and visits for participants. A schematic diagram is highly recommended (see Figure)	9, NA	
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1 2 3 4 5	Sample size	14	Estimated number of participants needed to achieve study objectives and how it was \vec{k} etermined, including clinical and statistical assumptions supporting any sample size calculations	9, 10 NA
	Recruitment	15	Strategies for achieving adequate participant enrolment to reach target sample size	9, 10
6 7	Methods: Assignment of interventions (for controlled trials)			
8 9 10 11 12 13 14 15 16 17 18 19	Allocation:		nterventions (for controlled trials)	
	Sequence generation	16a	Method of generating the allocation sequence (eg, computer-generated random numbers), and list of any factors for stratification. To reduce predictability of a random sequence, details of any planned restriction (eg, blocking) should be provided in a separate document that is unavailable to those who enrol participants or assign interventions	NA
	Allocation concealment mechanism	16b	Mechanism of implementing the allocation sequence (eg, central telephone; sequentially numbered, opaque, sealed envelopes), describing any steps to conceal the sequence until interventions are assigned	NA
20 21 22 23	Implementation	16c	Who will generate the allocation sequence, who will enrol participants, and who will assign participants to interventions	NA
23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46	Blinding (masking)	17a	Who will be blinded after assignment to interventions (eg, trial participants, care providers, outcome assessors, data analysts), and how	NA
		17b	If blinded, circumstances under which unblinding is permissible, and procedure for recealing a participant's allocated intervention during the trial	NA
	Methods: Data collection, management, and analysis			
	Data collection methods	18a	Plans for assessment and collection of outcome, baseline, and other trial data, including any related processes to promote data quality (eg, duplicate measurements, training of assessors) and a description of study instruments (eg, questionnaires, laboratory tests) along with their reliability and adality, if known. Reference to where data collection forms can be found, if not in the protocol	17-18
		18b	Plans to promote participant retention and complete follow-up, including list of any outcome data to be collected for participants who discontinue or deviate from intervention protocols	NA
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1 2 3 4 5 6 7 8 9 10 11 12 13	Data management	19	Plans for data entry, coding, security, and storage, including any related processes to promote data quality (eg, double data entry; range checks for data values). Reference to where details of data management procedures can be found, if not in the protocol	17-18
	Statistical methods	20a	Statistical methods for analysing primary and secondary outcomes. Reference to where other details of the statistical analysis plan can be found, if not in the protocol $\overset{\circ}{o}_{o}^{\circ}$	17-18
		20b	Methods for any additional analyses (eg, subgroup and adjusted analyses)	17-18
		20c	Definition of analysis population relating to protocol non-adherence (eg, as randomised analysis), and any statistical methods to handle missing data (eg, multiple imputation)	17-18
14 15	Methods: Monitoring		statistical methods to handle missing data (eg, multiple imputation)	
16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42	Data monitoring	21a	Composition of data monitoring committee (DMC); summary of its role and reporting structure; statement of whether it is independent from the sponsor and competing interests; and reference to where further details about its charter can be found, if not in the protocol. Alternatively, an explanation of why a DMC is not needed	NA
		21b	Description of any interim analyses and stopping guidelines, including who will have access to these interim results and make the final decision to terminate the trial	NA
	Harms	22	Plans for collecting, assessing, reporting, and managing solicited and spontaneously reported adverse events and other unintended effects of trial interventions or trial conduct $\overset{g}{}_{\geq}$	NA
	Auditing	23	Frequency and procedures for auditing trial conduct, if any, and whether the process will be independent from investigators and the sponsor	NA
	Ethics and dissemination		by gree	
	Research ethics approval	24	Plans for seeking research ethics committee/institutional review board (REC/IRB) ap	4 and 18
	Protocol amendments	25	Plans for communicating important protocol modifications (eg, changes to eligibility creating important protocol modifications (eg, changes to eligibility creating eria, outcomes, analyses) to relevant parties (eg, investigators, REC/IRBs, trial participants, trial regiseries, journals, regulators)	18
43 44 45			For peer review only - http://bmjopen.bmj.com/site/about/guidelines.xhtml	

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1 2	Consent or assent	26a	Who will obtain informed consent or assent from potential trial participants or authorised surrogates, and how (see Item 32)	18
3 4 5 6		26b	Additional consent provisions for collection and use of participant data and biological generic in ancillary studies, if applicable	NA
7 8 9	Confidentiality	27	How personal information about potential and enrolled participants will be collected, spared, and maintained in order to protect confidentiality before, during, and after the trial	18
10 11 12	Declaration of interests	28	Financial and other competing interests for principal investigators for the overall trial and each study site	28
13 14 15	Access to data	29	Statement of who will have access to the final trial dataset, and disclosure of contract a a greements that limit such access for investigators	18
16 17 18	Ancillary and post- trial care	30	Provisions, if any, for ancillary and post-trial care, and for compensation to those who suffer harm from trial participation	NA
19 20 21 22 23	Dissemination policy	31a	Plans for investigators and sponsor to communicate trial results to participants, healtheare professionals, the public, and other relevant groups (eg, via publication, reporting in results databases, or other data sharing arrangements), including any publication restrictions	19
24 25		31b	Authorship eligibility guidelines and any intended use of professional writers	28
26 27 28		31c	Plans, if any, for granting public access to the full protocol, participant-level dataset, and statistical code	28
29 30	Appendices		23, 2	
31 32 33	Informed consent materials	32	Model consent form and other related documentation given to participants and authorායිed surrogates ද පු	Correspondent author (in Dutch)
34 35 36	Biological specimens	33	Plans for collection, laboratory evaluation, and storage of biological specimens for genetic or molecular analysis in the current trial and for future use in ancillary studies, if applicable	NA
37 38 39 40 41	Amendments to the p	rotocol	that this checklist be read in conjunction with the SPIRIT 2013 Explanation & Elaboratien for important clarifica should be tracked and dated. The SPIRIT checklist is copyrighted by the SPIRIT Group under the Creative Co NoDerivs 3.0 Unported" license.	
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Keywords:	COVID-19, Health policy < HEALTH SERVICES ADMINISTRATION & MANAGEMENT, Human resource management < HEALTH SERVICES ADMINISTRATION & MANAGEMENT

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VITALITY, RESILIENCE AND THE NEED FOR SUPPORT AMONG HOSPITAL EMPLOYEES DURING THE

COVID-19 PANDEMIC: STUDY PROTOCOL OF A MIXED METHODS STUDY

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Abstract

Introduction

The Coronavirus Disease 2019 (Covid-19) pandemic has had a significant impact on the physical and mental functioning of healthcare professionals, especially those working on the 'frontline', and other hospital workers. At the onset of the crisis, various interventions were introduced to promote resilience and offer mental support to these professionals. However, it is unknown whether the interventions will meet the needs of professionals as the Covid-19 pandemic continues.

The goal of this exploratory study is to gain insight in factors that protect the vitality and resilience of Dutch hospital employees during the so-called 'second wave' of the CovidD-19 pandemic. This paper describes the study protocol.

Methods and analysis

This exploratory study applies a mixed-methods design, using both quantitative and qualitative methods of data collection and analysis. The first part of the study (sub-study I) consists of surveys among doctors and nurses in Covid-19 departments and non-Covid-19 departments, and other professionals in the hospital (i.e., managers and homeworkers) in 2020 and 2021. The second part of the study (sub-study II) consists of focus groups and interviews among professionals of the intensive care unit, Covid-19 departments and infection prevention units.

Ethics and dissemination

The research protocol for this study has been approved by the Medical Ethics Committee (MEC-2020-0705). The outcomes of this study will be used to develop and implement interventions to support hospital employees maintaining their vitality and resilience during and after the Covid-19 pandemic. Employees with vitality experience less work-related stress and make a positive contribution to healthcare quality.

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Article summary

Strengths and limitations of this study

- A mixed-methods design will be applied which strengthens the insights on vitality, resilience and the need for support among hospital employees.
- Insight in vitality, resilience and need for support of frontline workers from different departments will be investigated, as well as managers and homeworkers who will be compared in contrast to the majority of studies so far, which focused mainly on the needs of healthcare professionals such as nurses and doctors.
- Real life data gathering started during the beginning of 2nd Covid-19 wave, ongoing to autumn 2021.
- The Covid-19 pandemic is the motivation for this study, but may also limit the response rates or generalizability of this study, given its unpredictable course.

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Key words:

Covid-19, healthcare professionals, mental health, needs assessment, need for support, resilience,

vitality.

Introduction

Worldwide, it has been reported that the Coronavirus Disease 2019 (Covid-19) pandemic had a significant impact on the physical and mental functioning of healthcare professionals, especially for those working on the 'frontline' (e.g., intensive care units (ICUs), Covid-19 departments and infection prevention units)[1-4]. Indeed, also in the Netherlands, the Covid-19 pandemic had an impact on healthcare workers. This is critical, as it has been reported that some Dutch medical professionals were already overburdened before the pandemic[5, 6].

The need for high-intensity medical treatment of patients rapidly increased during the Covid-19 pandemic, during which the work circumstances became uncertain and stressful[7]. These work circumstances included the continuous use of personal protective equipment, adapted responsibilities and tasks, moral dilemmas, and the risk of contamination for the healthcare professionals themselves[8]. Interpersonal contact with patients' family members, one of the core features of the professional practice of nurses, was considerable reduced due to visiting limitations in most hospitals[9, 10]. In addition, the work environment also changed for ICU nurses as their teams changed due to the practical help from (former) colleagues and other healthcare professionals. This sudden shift in activities and responsibilities required ICU nurses to have additional competences maintaining high-quality healthcare. Buddies, or support staff from other departments in the hospital, were sometimes confronted with distressing or even shocking events during the first hectic weeks of the pandemic. Professionals of the infection prevention unit had to deal with an enormous workload due to the accumulation of new tasks and changing work processes, as well as the social turbulence resulting from the implemented quarantine measures. In the case of a health crisis such as the Covid-19 pandemic, the health and vitality of the frontline professionals became even more critical. Because a higher workload and stress could have a higher appeal on the physical and mental resources of the professionals. However, the Covid-19 pandemic not only had impact on the clinicians of the hospital. The work environment also changed for non-

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clinical professionals who suddenly had to work and communicate from home. In addition to this, homeworkers might lack a sense of purpose, solidarity and valuable contribution to the crisis situation[11]. Last, the Covid-19 pandemic required great effort from managers[12]. More than ever, they had to deal with logistic and administrative processes in the upscaling of high-intensity care, improving work alliances and the integration of staff in newly formed teams, and in managing the continuous flow of changing information.

Health, vitality and resilience

In previous virus outbreaks, such as the outbreaks of SARS, Ebola and MERS, it became clear that increased stress levels at work in healthcare professionals were associated with fear of contamination; shortages of materials; poor communication between healthcare professionals; unclear work instructions and information; deficient or non-functioning equipment; and inadequate planning among healthcare professionals[13-16]. Experiences from China during the Covid-19 pandemic showed similar results[17-19]. In a European study on work-related stress reactions among ICU healthcare professionals, half of the respondents (50.4%) showed symptoms of anxiety after the first wave of Covid-19[1]. Early phase evidence on Covid-19 suggested that healthcare professionals experienced mood and sleep disturbances during the outbreaks, stressing the need to establish ways to minimize mental health risks and support interventions aiming at pandemic conditions[3]. In the short-term, this work-related stress can cause fatigue, sleep disorders, mistakes and moral distress[20]. Long-term effects of high work pressure include burnout, depression and post-traumatic stress, resulting in dropout due to illness and abandonment of paid employment[21, 22]. A recent Dutch study among intensivists reported a moderate risk for burnout (14.8%)[23]. Furthermore, recovery time - regaining strength after an intensive period at work- has been associated with physical and mental well-being[24], as a long recovery time is an early indicator of work-related stress and exhaustion[25]. In contrast to high workload, stress and less recovery time, vitality, resilience and job satisfaction were describes as characteristics of professionals that

counterbalance work-related stress[26, 27]. These characteristics could strengthen professionals' mental and physical well-being and their retention for work[28-30]. Therefore, professionals with a high level of vitality and resilience seemed more resistant to work pressure.

Interventions among healthcare professionals during the Covid-19 pandemic

A wide variety of studies have examined interventions to reduce the work-related stress of healthcare professionals during the Covid-19 pandemic. Providing personal protective equipment is the top priority, followed by fulfilling the psychological needs of professionals[31]. To support mental health and promote the vitality of healthcare professionals, various interventions, including buddy systems, peer support, coaching and easily accessible psychological help, were proposed during the first months of Covid-19 wave [7, 32-36]. Other individual interventions, such as telemedicine activities, e-package and self-help books, appeared promising[37-40]. For example, a hospital in China offered online courses to help medical professionals to deal with psychological problems[41]. Many interventions have taken an individual approach, but system-level changes in healthcare organizations seemed to have a wider reach than individual support[42]. A notable omission in the literature is that protective factors were given limited attention: the focus is on the stressors. Many possible interventions were likely to support professionals in times of a pandemic, however, it is not clear which intervention matches the needs of the professional most closely. Therefore, a study was set up to investigate which supportive interventions, system changes and other supportive factors could meet individual needs during and in the aftermath of the Covid-19 pandemic in a large academic hospital in the Netherlands.

Objectives

The overall goal of the explorative study is to gain insight into the risk and protective factors as well as the needs and barriers in the working environment related to the promotion of the vitality and resilience of employees. Our objective is to assess levels of vitality and resilience, and the need for

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support or resources among professionals with a focus on professionals working in ICUs, Covid-19 departments, homeworkers and infection prevention units. Furthermore, to gain more insight into the relationship of vitality and resilience with factors such as self-perceived health, stress, burnout, posttraumatic stress, and need for recovery. The aim of the current paper is to describe the protocol of this explanatory mixed-methods study.

Methods and analysis

Study design

A mixed-methods design, using both quantitative (Sub-study-I) and qualitative methods (Sub-study-II), is applied. Sub-study I is a cross-sectional online survey administered first in October 2020, when the second wave of the Covid-19 pandemic was upcoming and ongoing, followed with measurements in March and September 2021. Sub-study II includes focus group interviews among nurses, doctors, and professionals regarding the ICU, Covid-19 departments and the infection prevention unit during the end of 2020.

Setting

The study setting is a large academic hospital in the Netherlands.

Study population

<u>Sub-study I</u>: The population consists of a random sample drawn based on voluntary participation of four target groups: professionals working at the Covid-19 department, non-Covid-19 departments, managers and homeworkers. A convenience sample has been used to monitor the health of the hospital workers , as was also done in comparable studies performed during the COVID-19 pandemic [43, 44]. We estimated the sample size of the consecutive quantitative measurements as 25% of the healthcare workers in the four target groups. Several organisational strategies will be followed to stimulate participation and reach the threshold of the aimed response rates.

<u>Sub-study II</u>: The population for the focus groups are the frontline workers. Maximum variation sampling is used, with respect to the type of frontline departments (ICU, COVID-19 departments, infection prevention unit) and occupational groups (physicians, nurses and infection prevention experts), resulting in six focus groups.

The inclusion criteria for the entire study are (i) a minimum age of eighteen years and (ii) sufficient Dutch language proficiency to complete the questionnaires or to discuss the relevant topic.

Patient and Public Involvement

No patient involved.

Study procedures

Sub-study I: Online survey

Hospital employees are informed about the study in several ways. The communication strategy is tailored to each target group and supported by the communication department of the organization. A link to the online survey is published on the intranet of the organization, printed QR-codes containing a link to the survey are available at the coffee corners and canteens, announcements are made in the weekly Covid-19 livestream and by team management via personal email. Participation is voluntary and can be performed during working hours.

The online questionnaire starts with information about the study, privacy statements and an informed consent form for participation. After providing consent, participants are asked to fill out the entire questionnaire, which consists of two parts. The first part is generic for all employees and takes approximately six minutes to complete; it includes questions on demographic information and the main outcomes. The second part consists of additional modules on working conditions and health and takes approximately seven minutes. Nurses and homeworkers receive an additional module tailored to their specific work environment.

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Sub-study II: Focus groups

In total, six focus groups with 6-10 participants that take approximately 60 minutes are conducted. ICU doctors, ICU nurses, microbiologists, hospital hygienists, Covid-19 unit nurses, and Covid19-unit doctors (lung specialists and specialists internal medicine) are individually invited to participate in one of the focus groups through consultation with the team managers. These meetings are preferably in-person (to observe non-verbal attitude and facial expressions), but due to the Covid-19 measures and social distancing, it may not be possible for participants to be physically present. In those cases, the focus groups are carried out via video calling technology.

Prior to the meetings, a topic list is created by the research group based on the literature and internal reports on the experiences of professionals. This topic list is used to guide and structure the meeting. The aim of the focus group is to study protective factors that contribute to vitality and resilience during the Covid-19 pandemic. Furthermore, possible interventions to increase vitality and resilience are explored and elaborated upon. Written informed consent is given prior to the meeting, and two experienced researchers guide the meetings. The focus group interviews are recorded and transcribed verbatim.

Measurements

This paragraph lists all measurement instruments included in the questionnaire. The first part consists of measuring instruments addressing demographics, primary outcomes (i.e. vitality, resilience and needs assessment), and several secondary outcomes (i.e. self-perceived health, stress, burnout, posttraumatic stress, and need for recovery). The second part consists of separate modules for homeworkers and nurses with regard to work ability, working conditions, job satisfaction, workprivate balance, exposure to Covid-19 at work, preventive measures for Covid-19 and career perspectives.

Demographics

Gender, age, educational level, job titles, work location and professionals' experience (in years) are assessed. Educational level is divided into three levels: low, medium and high educational level. In total, the list of job titles includes 23 positions within the academic hospital (e.g., nurse, ICT employee, pharmacist, educator, researcher).

Main outcome measures

Vitality

Vitality is measured with four items from the original 36-item Short Form Health Survey (SF-36) [45]. The total summed score of four items that refer to the past four weeks: "Did you feel full of liveliness?", "Did you have a lot of energy?", "Did you feel worn out?", and "Did you feel tired?". The answers are rated on a six-point scale from 1 (= constantly) to 6 (= never) [46]. Higher scores indicating a better subjective vitality.

Resilience

Resilience (the ability to cope with stress, setbacks or difficulties at work) is measured with six items from the Psychological Capital Questionnaire[47]. The items contain statements such as: "When I have a setback at work, I have a hard time getting back on track and moving on", "If necessary, I can work well without the help of others" and "I can handle difficult moments at work". The six items are scored from 1 (= strong disagreement) to 6 (= strong agreement). Higher values indicate a higher level of resilience.

Needs assessment

Needs are measured with a self-designed scale with four items. Examples of questions are: "In which area would you like to be supported?" and "What would this support look like?" and "What should

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be offered or developed?". A predefined list includes 10 individual- and 14 organizational-related answer options, e.g., support for working from home, time management, and work-private balance.

Other outcome measures

Self-perceived health

Self-rated health is assessed with one question: "In general, how would you say your health is?" Answer options from 1 (= excellent) to 5 (= poor).

Stress

Stress is measured with a numeric rating scale. The stress score, ranging from 0 (= no stress at all) to 100 (= the worst stress imaginable). This scale is used to retrospectively objectify stress before, during and after the first COVID-19 outbreaks. The three item question was "How did you experience the stress before/during/after the COVID-19 crisis on a scale from 0 to 10?"

Burnout

Burnout is measured using five items, that are based on an adapted version of the Utrecht Burnout Scale [48]. The items refer to the current situation such as "I feel emotionally drained from my job" and "I feel completely exhausted from my work". The answer options from 1 (= never) to 7 (= daily).

Posttraumatic stress

Posttraumatic stress is assessed with the Posttraumatic stress disorder (PTSD) Checklist for the DSM-5 (PCL-5) - Covid-19 version with 20 items[49]. This scale consists of 20 items, measuring PTSD symptoms, with scoring options from 0 (= not all) to 4 (= extremely) and was adapted to the Covid-19 situation. A score of 33 or higher is preceived indicative for PTSD.

Need for recovery

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Work fatigue and the risk of psychological symptoms are measured using the Dutch questionnaire on the Experience and Evaluation of Work (Dutch abbreviation: VBBA)[50, 51]. The need for recovery scale consists of eleven dichotomous items (yes/no), representing short-term effects of a working day[24, 52, 53]. The score of the need for recovery scale ranges from 0 to 100 and is calculated as the sum of points (1 = yes, 0 = no) divided by the number of questions answered, multiplied by 100. Higher scores indicate a higher need for recovery, which is unfavourable.

Work ability

Work ability is measured with the Work Ability Index (WAI)[54]. This widely used index measures self-assessed work ability and consists of seven items. Because the sub-items of the WAI can also be used as a simple indicator for work ability[55], three of the seven items are used: current work ability (one item), and work ability in relation to physical and mental job demands (two items). A total WAI score (range 2–20) is obtained by adding the weight scores of these individual items[56].

Working conditions

Aspects of work load in the current study are: job autonomy, emotional job demands, social support and physical working conditions.

Job autonomy is measured with six items on a three point scale (no; yes, sometimes; yes, regularly). Five items, i.e., those about making decisions, having to find solutions, and being able to take time off, are based on the Job Content Questionnaire[57, 58]. One item on autonomy related to working time based on the Netherlands Working Conditions Survey, is also included in the questionnaire[59].

Emotional job demands is evaluated with four items. Three items are derived from the Copenhagen Psychosocial Questionnaire and assess whether the work leads to emotionally difficult situations, the emotional demands of the job, and emotional involvement in work. An additional item is "Is your

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job more emotionally demanding because of Covid-19?". All items are measured on a four-point scale (never to always)[60].

Social support is defined as whether colleagues and supervisors are willing to help and listen to work-related problems and is assessed using four items from COPSOQ[60]. Social support is measured on four-point Likert scales 1 (= almost never) to five (= always).

Physical work load are measured with one self-designed question and assesses whether a worker received more or less physically demanding work due to Covid-19 measures. This scale has three answer options (no; yes, sometimes; yes, regularly).

Job satisfaction

Job satisfaction is measured with one item: "Altogether, how satisfied are you with your work?" The answer options range from 1 (very dissatisfied) to 5 (very satisfied).

Work-private life balance

Work-private life balance is measured with two questions on the mutual interference between work and home life. The questions are adopted from the Netherlands Working Conditions Survey [59], but were originally constructed by Fox and Dwyer (1999) [61]. Both questions have four answer options ranging from 1 (= no, never) to 4 (= very often).

Exposure to COVID-19 at work

Professionals are asked to what extent they might have been exposed to Covid-19 at the worksite. These questions are derived from the Netherlands Working Conditions Survey Covid-19 [62], based on questionnaires developed within the OMEGAnetwork[63]. Participants are asked if they work with patients, the average number of patients they work with during a typical working day in the last

week, and if these patients arre suspected to have or had been diagnosed with Covid-19. Additionally, participants are asked if and with how many workers they work on a regular basis with colleagues, and if they share tools or surfaces with their colleagues.

Preventive measures for Covid-19

The five questions on preventive measures with regard to Covid-19 are derived from the Netherlands Working Conditions Survey Covid-19 [62]. One general question assesses the general measures taken at the department level with regard to the Covid-19 pandemic, with answer options such as homeworking, adjustment of working hours, general preventive measures in the workplace, mandatory inclusion or withdrawal of leave. The specific questions on preventive measures include the possibility of keeping a 1.5 metre distance between colleagues and/or patients, the availability of personal protective equipment, the usage of personal protective equipment and the application of general hygiene measures. The responses to these five questions are never, sometimes, often and always. This module will not be applied to homeworkers.

Career perspective

Three items on career perspective are derived from the Netherlands Working Conditions Survey Covid-19[62] and adjusted to fit the study population working in the hospital. These items include the motivation to work in the healthcare sector in the future (responses: less, equal, and more), the intention to change jobs within the health care sector and the intention to change jobs outside the healthcare sector with responses ranging from 1 (= certainly not) to 5 (=certainly yes).

Outcome measures for pre-defined groups or professions

Nurse questionnaire

The Practice Environment Scale of the Nursing Work Index is the most widely used measure to gauge the state of nursing practice environments[64, 65]. It is the only measure recommended by several

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organizations promoting quality healthcare. The 15-item questionnaire uses responses ranging from 1 (= strongly disagree) to 4 (= totally agree). This module will be applied to nurses only.

Homeworkers

A total of eight items are specifically tailored to homeworkers. Two items refer to the number of hours in a week people work from home and how many hours a day they work on a screen (e.g., laptop and tablet). One item is focused on the availability of ergonomic work equipment at home (a desk or table with a comfortable working height, a chair that can be adjusted to one's body measurements, a separate display, and a separate computer mouse). The need for other furniture is assessed with one item "Do you need additional materials for a good home workplace?". Moreover, participants are asked if they take (short) breaks on a working day, except for a lunch break?". This question includes the following answer options: 1 (= yes, regularly), 2 (= yes, sometimes) and 3 (= no). The last three items are about concentration while at home and include the following statements: "Do you have trouble concentrating while working?", "Do you struggle to keep your attention while you work?" and "Do you have difficulty with the reduced social contact with colleagues?" Answer options range from 1 (= never) to 4 (- always).

Data handling and statistical analyses

<u>Sub study I</u>: survey data are anonymously collected using Limesurvey (Version 2.06 Its Build 160524) and exported to a secure SPSS database (©IBM SPSS Statistics for Windows, Version 25.0. Armonk, NY: IBM Corp) for analysis. All principal investigators have access to the final study dataset. Data will be stored for fifteen years.

First, the data are cleaned and checked for missing data. The descriptive statistics are presented as numbers and percentages for dichotomous variables and mean and standard deviation for for continuous variables. Data for different subgroups (professionals in Covid-19 departments, non-Covid-19 departments, managers and homeworkers) are analyzed with the Mann-Whitney test or t-

tests. Linear and logistics regression analyses are preformed to investigate the associations between risk factors and the main outcomes (vitality and resilience). Statistical significance will be defined as p < .05.

<u>Sub study II:</u> Focus groups data will be analyzed by means of thematic content analysis [66]. This method organizes and describes the data set in rich detail and investigates patterns of response or meaning within the data set. We take an inductive approach to identify possible themes. Once a satisfactory thematic map is established, the themes are examined to identify the 'essence' of what each individual theme is about and to understand how they are interrelated in relation to our research question. To achieve this, the following steps will be taken:

Focus group interview data are audiotaped and transcribed verbatim. [66]Two researchers will read the transcripts in detail. Each of them starts with developing a structured analysis framework that consists of preliminary codes and themes. They make use of mind-maps and tables to organize the data. After that, they compare their frameworks to reach consensus. Next, one researcher codes the transcripts line by line according to this framework in the software programme NVivo12[®]. The coder uses memos for comments during coding. When coding is finished and the code 'other' is used, the two researchers discuss these codes and rename them into a new or existing codename best reflecting the contents of the otherwise uncategorised text fragment. During and after coding, the two researchers review and check the themes for internal homogeneity and external heterogeneity. Finally, the two researchers analyse the cohesion and inter-relations between themes to come to a coherent account and accompanying narrative of the data. The principal investigators have access to these data, which will be stored for fifteen years.

Ethics and dissemination

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The study is approved by the Medical Ethics Committee of the Erasmus MC (MEC-2020-0705). It will beconducted according to the principles of the Declaration of Helsinki (64th WMA General Assembly, Fortaleza, Brazil, October 2013) and in accordance with the Medical Research Involving Human Subjects Act. The study complies with the Netherlands Code of Conduct for Scientific Practice from the Association of Universities in the Netherlands. Protocol modifications will be communicated and to the Medical Ethics Committee by protocol amendment. Participants will be informed about the study both orally and by letter. Consent for participation will be given by written informed consent. Participants can leave the study at any time for any reason if they wish to do so without any consequences. The withdrawal will be registered for informative purpose.

Discussion

The consequences of the Covid-19 crisis on the mental health and working conditions of healthcare professionals have been recognized worldwide[67]. Hospital employees with vitality experience less work-related stress and can therefore handle more work in the new and stressful circumstance. In other words, maintaining professionals' vitality and resilience will contribution to healthcare quality. By using a mixed-methods approach, we aim to gain an overview of vitality, resilience and health (e.g., stress and burnout) among healthcare professionals, as well as the risk factors associated with these outcomes. The covid-19 pandemic has put an extra focus on the impact of work-related stress and how to deal with its causes and consequences. Even though the pandemic entails a specific surge of specific patients, and as such may hamper generalizability, we believe that the outcomes of this study will add to the body of knowledge on how best to deal with the work related stress experienced by healthcare workers worldwide.

This is an urgent and rushed study because we wanted to use the results against the same health crisis that we are investigating. Based on this study, directions for future interventions during the

Covid-19 pandemic and thereafter could provide raised levels of vitality and resilience of professionals in the hospital, and therewith support their employability in the long run. *Strengths and limitations*

The first strength is the mixed-methods design, consisting of qualitative and quantitative methods which provide a more in-depth insight in the need for support in the exploratory study and therewith details information to develop interventions. . Second, we compare different departments and distinguish healthcare workers, managers, and homeworkers. The majority of studies so far focused exclusively on the needs of healthcare professionals without considering other hospital employees such as supportive staff, researchers and managers.

The Covid-19 pandemic was the motivation for this research, but may also have limited the procedure of this study, given its unpredictable course. During the writing of this protocol paper, the second wave of Covid-19 had already started in The Netherlands. Therefore, a lower response rate is not unexpected from the frontline healthcare workers. The second limitation is the cross-sectional design of the study, which makes it impossible to draw causal conclusions from this report and to investigate the lont-term effects.

Data dissemination

Public access to the study protocol, study details, participant-level dataset, and statistical code can be acquired from the corresponding author. The results will be disseminated to healthcare professionals, health services authorities and the public via presentations at national and international meetings and published in peer-reviewed journals. A lay summary of the results will be written and shared with all professionals of the organization.

Study status

The study is currently ongoing with data recruitment.

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Declarations

Authors contributions

MvM, TK, JB and LK jointly designed the study, raised funding and established the development of the study protocol. MvM, MV, TK, KOH and LK prepared the study materials. MvM, MV and LK gathered the data of both sub-studies and produced the first draft of the article outline together with KOH and TK. All authors (MvM, MV, AP, TK, JB, WH, KOH, LK) contributed substantially to the concept of the study, the analyses of literature, critically revised the content of the manuscript, have read and approved the final version.

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Competing interests

The authors declare no conflicts of interest.

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Availability of data and materials

Anonymized data gathered and analysed during the current study are not publicly available due to legal and ethical restriction. These can be requested from the corresponding author as well as text and photo material of the developed intervention. Materials described in the manuscript, including all relevant raw data, will be freely available at a reasonable request to any scientist wishing to use them for non-commercial purposes.