# Well-being and attention in medical scientists

Submission date	Recruitment status	Prospectively registered
27/11/2023	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
01/12/2023	Completed	Results
Last Edited	Condition category	Individual participant data
17/12/2024	Other	[X] Record updated in last year

### Plain English summary of protocol

Background and study aims

Medical scientists have dual commitments to clinical care and research efforts. Such commitments can create hectic and stressful work schedules, which may impact well-being and cognition. This study tests the hypothesis that mindfulness coaching plus wearable-based lifestyle monitoring can benefit such individuals.

#### Who can participate?

Physician Scientists affiliated with the UC San Diego Medical Scientist Training Program aged between 20 and 45 years old

#### What does the study involve?

The study involved mindfulness coaching plus wearable-based lifestyle monitoring. Mindfulness coaching will be provided in small group sessions led by two psychological health experts as coaches. The coaches lead three sessions (each lasting 1.5 hours) spread over one academic quarter for each small group. The group meetings will be held virtually to maximize accessibility and comfort for the participants and will be scheduled per the participants' mutually preferred times. In each session, the coaches will conduct mindfulness exercises and lead an interactive discussion on health, well-being, resilience, leadership training, and self-compassion. Different mindfulness tools and perspectives will be suggested and discussed by the facilitators and by the participants themselves to address work-life challenges. The participants will be encouraged to share their sources of support (e.g., exploring sources of strength, stress management techniques, and spending time with loved ones). Focused mindfulness practice will be offered and taught at each session. At the end of each session, participants will be encouraged to prepare to share a tool or thought that was discussed at the next session. Intervention group participants will each be provided with a Garmin Vivosmart® 4 (a wearable device) to passively monitor and receive feedback on lifestyle factors (physical activity, heart rate, sleep quality and stress) via an app coupled to the Garmin for the duration of the intervention.

What are the possible benefits and risks of participating? Benefits to participants may include improvements in well-being and/or cognition. This is a minimal-risk study.

Where is the study run from?
The University of California San Diego School of Medicine (USA)

When is the study starting and how long is it expected to run for? January 2021 to December 2022

Who is funding the study?

National Institutes of Health (Grant: NIH/NIGMS 1T32GM121318-01) (USA)

Who is the main contact?

Jyoti Mishra, jymishra@ucsd.edu (USA)

### Contact information

### Type(s)

Public, Scientific, Principal Investigator

#### Contact name

Dr Jyoti Mishra

#### **ORCID ID**

http://orcid.org/0000-0001-6612-4557

#### Contact details

9500 Gilman Drive MC0875 La Jolla United States of America 92037 +1 8582322855 jymishra@ucsd.edu

### Additional identifiers

### **EudraCT/CTIS** number

Nil known

#### IRAS number

### ClinicalTrials.gov number

Nil known

### Secondary identifying numbers

Nil known

## Study information

#### Scientific Title

Mindfulness coaching with digital lifestyle monitoring enhances selective attention in medical scientists

#### Acronym

WellMSTP

### **Study objectives**

Mindfulness coaching with digital lifestyle monitoring will enhance well-being and cognition and medical scientists

### Ethics approval required

Ethics approval required

### Ethics approval(s)

Approved 29/01/2021, University of California San Diego Institutional Review Board (9500 Gilman Drive, La Jolla, 92037, United States of America; +1 858-246-4777; hrpp@ucsd.edu), ref: 180140

### Study design

Interventional

#### Primary study design

Interventional

### Secondary study design

Randomised controlled trial

#### Study setting(s)

Internet/virtual, University/medical school/dental school

### Study type(s)

Prevention

### Participant information sheet

See study outputs table

#### Health condition(s) or problem(s) studied

Well-being

#### **Interventions**

Mindfulness Coaching

Participants will be randomized using a simple randomization method into the intervention or a waitlist control group. The intervention is a digital approach (delivered on Zoom) with group mindfulness sessions led by two psychological health faculty members as coaches. For these sessions, participants will be divided into sub-groups by career stage: pre-clinical medical scientist training program (MSTP) students, MSTP graduate students, or clinician/junior faculty. The coaches will lead three sessions (each 1.5 hours) spread over one academic quarter for each sub-group. The group meetings will be held virtually to maximize accessibility and comfort for the participants and will be scheduled per the participants' mutually preferred times. In each session, the coaches will conduct mindfulness exercises and lead an interactive discussion on health, well-being, resilience, leadership training, and self-compassion. Participants will be asked to turn on their videos for the duration of the sessions and to share personal opinions and life experiences. In these sessions, the primary goal for the coaches will be to listen attentively and promote well-being. They will also aim to validate and acknowledge challenges that participants face as physician-scientists in training or as early career faculty. Different mindfulness tools and perspectives will be suggested and discussed by the facilitators and by the participants

themselves to address work-life challenges. The participants will be encouraged to share their sources of support (e.g., exploring sources of strength, stress management techniques, and spending time with loved ones). Focused mindfulness practice will be offered and taught at each session. The participants will also discuss potential changes to the physician-scientist environment that would facilitate well-being if implemented. At the end of each session, participants will be encouraged to consider a tool or thought that was discussed to share at the next session. Intervention group participants will each be provided with a Garmin Vivosmart® 4 (a wearable device) to passively monitor and receive feedback on lifestyle factors (physical activity, heart rate, sleep quality, and stress) via an app coupled to the Garmin for the duration of the intervention.

### Intervention Type

Behavioural

#### Primary outcome measure

1. Subjective well-being measured using a combination of the Mindful Attention Awareness Scale (MAAS), Short Warwick Edinburgh Mental Well-Being Scale (SWEMWBS), Self-Compassion Scale (SCS), and the Maslach Burnout Inventory (MBI) at baseline and post-intervention 2. Objective cognition measured using the BrainE neurocognitive platform at baseline and post-intervention

### Secondary outcome measures

Neuroplasticity underlying cognition measured using electroencephalography (EEG) recordings synchronized to the BrainE neurocognitive platform assessments at baseline and post-intervention

### Overall study start date

01/01/2021

### Completion date

31/12/2022

## **Eligibility**

### Key inclusion criteria

Physician Scientists affiliated with the UC San Diego Medical Scientist Training Program

### Participant type(s)

Healthy volunteer, Carer

### Age group

Adult

### Lower age limit

20 Years

### Upper age limit

45 Years

Sex

### Target number of participants

43

### Total final enrolment

43

### Key exclusion criteria

Individuals not affiliated with the UCSD Medical Scientist Training Program

#### Date of first enrolment

01/10/2021

#### Date of final enrolment

01/12/2021

### Locations

#### Countries of recruitment

United States of America

# Study participating centre University of California San Diego School of Medicine

9500 Gilman Drive La Jolla United States of America 92037

# Sponsor information

### Organisation

University of California San Diego Medical Center

### Sponsor details

9500 Gilman Drive La Jolla United States of America 92037 +1 858-246-4777 irb@ucsd.edu

### Sponsor type

Hospital/treatment centre

#### Website

https://health.ucsd.edu/

#### **ROR**

https://ror.org/03aw5sn18

# Funder(s)

### Funder type

Government

#### **Funder Name**

National Institutes of Health

#### Alternative Name(s)

Institutos Nacionales de la Salud, US National Institutes of Health, NIH

#### **Funding Body Type**

Government organisation

### **Funding Body Subtype**

National government

#### Location

United States of America

### **Results and Publications**

### Publication and dissemination plan

Planned publication in a high-impact peer-reviewed journal

### Intention to publish date

01/06/2025

### Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study will be stored in a publically available repository, datadryad.org. The type of data stored includes all data outcome variables collected and analysed for the study. The data will be freely available for download post-publication of the research. Consent was obtained from participants for de-identified data sharing. The data will be de-identified and do not have any personal health information for participants. The data are de-identified with no ethical or legal restrictions.

### IPD sharing plan summary

Stored in publicly available repository

Study outputs

Output type Details Date created Date added Peer reviewed? Patient-facing?

Participant information sheet 22/03/2016 28/11/2023 No Yes