



Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its COVID-19-related research that is available on the COVID-19 resource centre - including this research content - immediately available in PubMed Central and other publicly funded repositories, such as the WHO COVID database with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source. These permissions are granted for free by Elsevier for as long as the COVID-19 resource centre remains active.



Contents lists available at ScienceDirect

Journal of Affective Disorders

journal homepage: www.elsevier.com/locate/jad

Short communication

Moderators of a resiliency group intervention for frontline clinicians during the COVID-19 pandemic

Louisa G. Sylvia^{a,b,c,*}, Nevita George^{a,c}, Dustin J. Rabideau^{a,b}, Joanna M. Streck^{a,b}, Evan Albury^{a,c}, Daniel L. Hall^{a,b,d,e}, Christina M. Luberto^{a,b,d,e}, Helen R. Mizrach^{a,e}, Giselle K. Perez^{a,b,d}, Sydney Crute^{a,e}, Darshan H. Mehta^{a,b,d}, Mary Susan Convery^{a,b,d}, Sara E. Looby^{a,b}, Gregory Fricchione^{a,b,d}, Maurizio Fava^{a,b}, Sabine Wilhelm^{a,b}, Elyse R Park^{a,b,d,e}

^a Massachusetts General Hospital, Boston, MA, USA^b Harvard Medical School, Boston, MA, USA^c Dauten Family Center for Bipolar Treatment Innovation, Boston, MA, USA^d Benson-Henry Institute, Boston, MA, USA^e Benson-Henry Institute for Mind Body Medicine, Boston, MA, USA

ARTICLE INFO

Keywords:

Frontline clinicians
COVID-19 pandemic
Stress coping
Resiliency program

ABSTRACT

Background: To mitigate the psychological burdens of COVID-19 for frontline clinicians (FCs), we adapted an existing evidence-based resiliency program, Stress Management and Resilience Training Relaxation Response Program (SMART-3RP), for FCs. This analysis explores moderators of stress coping to determine which subgroups of FCs benefited most from SMART-3RP.

Methods: 102 FCs from Mass General Brigham hospitals engaged in the adapted SMART-3RP. Assessments were completed at group entry (Week 0) and completion (Week 4). The primary outcome was stress coping, and we examined 15 possible baseline moderators. We fit linear mixed effects regression models and assessed potential baseline moderators using a likelihood ratio test. We report model-based estimates and confidence intervals for each moderator-by-time interaction (i.e., differential effect), where positive/negative values indicate more/less improvement in average perceived stress coping.

Results: Stress coping improved from Week 0 to Week 4 (mean improvement [95% CI] = 0.9 [0.6 to 1.2]). FCs with higher anxiety (differential effect [95% CI] = 0.3 [0.1 to 0.4]), depression (0.4 [0.2 to 0.6]), and loneliness (0.4 [0.1 to 0.6]), but lower levels of mindfulness (CAMS-R_{focus}: 1.0 [0.4 to 1.6]; CAMS-R_{accept}: 1.3 [0.7 to 2.0]) and self-compassion (0.4, [0.1 to 0.8]) at baseline experienced greater benefits in perceived stress coping from the SMART-3RP. Baseline health uncertainty along with sociodemographic and work characteristics did not moderate stress coping.

Discussion: Results highlight particular sub-populations of FCs that may benefit more from a stress management intervention, especially during emergency responses (e.g., COVID-19 pandemic).

1. Background

The novel coronavirus (COVID-19) created a workforce of frontline clinicians (FCs) who treated and cared for COVID-19 patients and subsequently experienced a myriad of stressors (Lu et al., 2020). This led to the development of stress-related disorders for FCs, such as anxiety and depression (Pappa et al., 2020). To support FCs at the onset of the COVID-19 pandemic, we adapted an evidence-based resiliency program,

the Stress Management and Resilience Training Relaxation Response Program (SMART-3RP) to build relaxation and mindfulness techniques among FCs (Mehta et al., 2016; Park et al., 2020). This program was adapted to help FCs face stressful circumstances specific to the pandemic such as health uncertainty, changes in their work environment, and personal challenges.

We recently reported that this adapted SMART-3RP program for FCs decreased emotional distress and increased perceived stress coping,

* Corresponding author: 50 Staniford St, Suite 580, Boston, MA 02114

E-mail address: lsylvia2@mgh.harvard.edu (L.G. Sylvia).

<https://doi.org/10.1016/j.jad.2021.06.036>

Received 1 January 2021; Received in revised form 5 April 2021; Accepted 19 June 2021

Available online 24 June 2021

0165-0327/© 2021 Elsevier B.V. All rights reserved.

defined as individual's ability to cope with stressors in their life (Park et al., 2020). In the present investigation, which is a secondary analysis of our earlier study, we explored potential moderators of perceived stress coping to determine which characteristics of FCs were associated with differential improvement in perceived stress coping from SMART-3RP. We focused on perceived stress coping as prior studies among FCs have highlighted that increases in perceived stress coping is associated with improvements in overall psychological health (Dunkley et al., 2017; Thimm et al., 2018; Rettie and Daniels, 2020), thus perceived stress coping is a proxy for global improvements in well-being. Identifying characteristics that moderate the effect of the SMART-3RP on improvements in perceived stress coping could be used to target FCs who will benefit the most from a stress management intervention, especially during emergency responses like the COVID-19 pandemic.

2. Methods

2.1. Subjects and procedure

Participants were English-speaking adult (>18 years) clinicians (i.e., responsible for patient care) in the Massachusetts General Brigham (MGB) healthcare system. There were no other exclusion criteria to recruit a generalizable sample of clinicians. FCs were grouped by the following specialties: physicians, nurses, physical therapists, occupational therapists, respiratory therapists, speech language pathologists, advanced practice providers, and mental health clinicians. Inclusion criteria required FCs to be over 18 years old and from either Boston Hope or Massachusetts General Hospital. Participants were excluded if they did not speak English. The study was conducted in accordance with ethical standards of the World Medical Association and was approved by our Institutional Review Board.

FCs (Age: $M=45$, $SD=12.2$, 92.1% female, 83.3% white) were recruited through hospital-wide emails and departmental announcements. Once registered (via an online website), participants completed a 25-item survey at group entry (Week 0) and group completion (Week 4) through REDCap (Research Electronic Data Capture), a secure, HIPAA compliant web-based application. Groups were conducted via zoom, an MGB-approved and HIPAA-compliant video conferencing platform. Participants were assigned to a group based on their specialty area, and each group met twice per week for 60 min over 4 weeks (i.e., 8 sessions in total). Groups were co-led by psychologists, physicians, social workers, and/or nurses employed at MGB and trained in delivering the SMART-3RP.

2.2. Measures

All assessments were administered pre- and post-group, but focused on baseline moderators.

2.3. Primary outcome

Stress Coping (Stress Coping) (Park et al., 2020). The perceived stress coping question, created by author (EP), is a single item measure ("How able have you been to cope with the stress in your life?") rated on a 10-point scale (i.e., 0=not at all) to 10=very well).

2.4. Potential baseline moderators

Sociodemographic variables. Age, provider specialty, gender identity, and race/ethnicity.

Work Variables. Self-report of a recent change in work hours (i.e., increased, decreased or stayed the same) and work characteristics (i.e., work setting, clinical role, patient population, use of telehealth to conduct clinical care).

Measure of Current Status (Carver, 2006) (MOCS-A). We used two items: coping response ("I am confident about being able to choose the

best coping response for hard situations") and degree of emotional thoughts ("I can come up with emotionally balanced thoughts even during negative times"). Items were rated on a 5-point scale. The MOCS-A has demonstrated reliable psychometric properties in previous studies (Antoni et al., 2006).

Cognitive and Affective Mindfulness Scale- Revised (Feldman et al., 2007) (CAMS-R). We used two items: acceptance ("able to accept thoughts and feelings") and mindfulness ("focus on the present moment"). Items were rated on a 4-point scale. The CAMS-R has good psychometric properties and focuses on aspects of mindfulness most clearly distinct from worry and rumination (Feldman et al., 2007).

Patient Health Questionnaire-4 (PHQ-4) (Kroenke et al., 2009). The PHQ-4 assesses anxiety with two items ("feeling nervous, anxious or on edge" and "not being able to stop or control worrying?") and depression with two items ("feeling down, depressed, or hopeless?" and "little interest or pleasure in doing things?"). We summed the two items to have a total score for anxiety and depression (range=0 to 6). This scale has good internal reliability, construct validity, and factorial validity along with reliable criterion, construct, and procedural validity of its two subscales, the PHQ-2 and GAD-2 (Kroenke et al., 2009).

Self-compassion Scale (Neff, 2003) (SCS). The SCS assessed self-compassion via one item ("When times are really difficult, I am tough on myself") rated on a 5-point Likert scale. Higher scores indicated less self-compassion. The SCS has good test-retest reliability as well as other psychometric properties (Neff, 2003).

University of California, Los Angeles (UCLA) Loneliness Scale (Russell et al., 1978). The UCLA Loneliness Scale assessed loneliness/isolation with two items ("I feel completely alone" and "I feel isolated from others"). Questions were rated on a 3-point scale (Not at All=0, Several Days=1, More than Half the Days=2, Nearly Every Day=3) and has demonstrated highly reliable psychometric properties, both in terms of internal consistency and test-retest reliability (Russell et al., 1978).

Health Uncertainty Item (Rogers et al., 2016). The Health Uncertainty Item is a single item measure that was used to assess how uncertain participants were about their health during COVID-19. For this study, we adapted the language on the scale to say, "my health" rather than "cancer recurrence".

Personal Strengths (Yanez et al., 2011). The Personal Strengths questions were taken from the Current Experiences Scale and assessed how confident participants were in their ability to cope with stress: ("I am confident about being able to choose the best coping response or hard situations") and ("I can come up with emotionally balanced thoughts even during negative times").

3. Statistical analysis

We examined six categorical baseline moderators (gender, race, clinical specialty, work hours in the past month, CAMS-R acceptance, CAMS-R mindfulness) and nine continuous moderators (i.e., anxiety and depression per the PHQ-4, SCS, ULCA, health uncertainty, MOCS-A coping response and emotional thoughts, age, number of individuals in one's household).

We calculated pairwise mean differences (post – pre) in perceived stress coping and corresponding 95% confidence intervals (CIs) for all participants and separately for each subgroup. Subgroups based on continuous moderators (e.g., age quartiles) were created for illustrative purposes only. Positive/negative pairwise mean difference values correspond to improvement/worsening in average perceived stress coping post-intervention. We then fit linear mixed effects regression models with random individual intercepts to incorporate all-available baseline and Week 4 outcomes into our analysis. Models were fit with and without an interaction term between the potential moderator and a post-intervention indicator (i.e., a moderator-by-time interaction) and assessed using a likelihood ratio test. We report likelihood-based estimates and CIs for the interaction terms. For simplicity, we assumed a

linear relationship between continuous moderators and mean change in perceived stress coping. Due to the exploratory nature of these analyses, no adjustments for multiple comparisons were made.

4. Results

Perceived stress coping significantly improved from pre to post-intervention (model-based mean improvement [95% CI] = 0.9 [0.6 to 1.2]) (Park et al., 2020). Pairwise mean differences and 95% CIs overall

and for each subgroup are displayed in Figure 1; complete model-based results are presented in the Table 1. Estimates and CIs in the Table 1 correspond to differential effects between subgroups (for categorical variables) or per one-unit increase (for continuous variables) (e.g. the difference in average improvement among males vs. females or per one-point increase on the PHQ-4 anxiety subscale). We found that baseline levels of acceptance and mindfulness (both CAMS-R items), anxiety and depression (PHQ-4), self-compassion (SCS), loneliness (UCLA), and coping response (MOCS-A) (the p-value was 0.08 for the

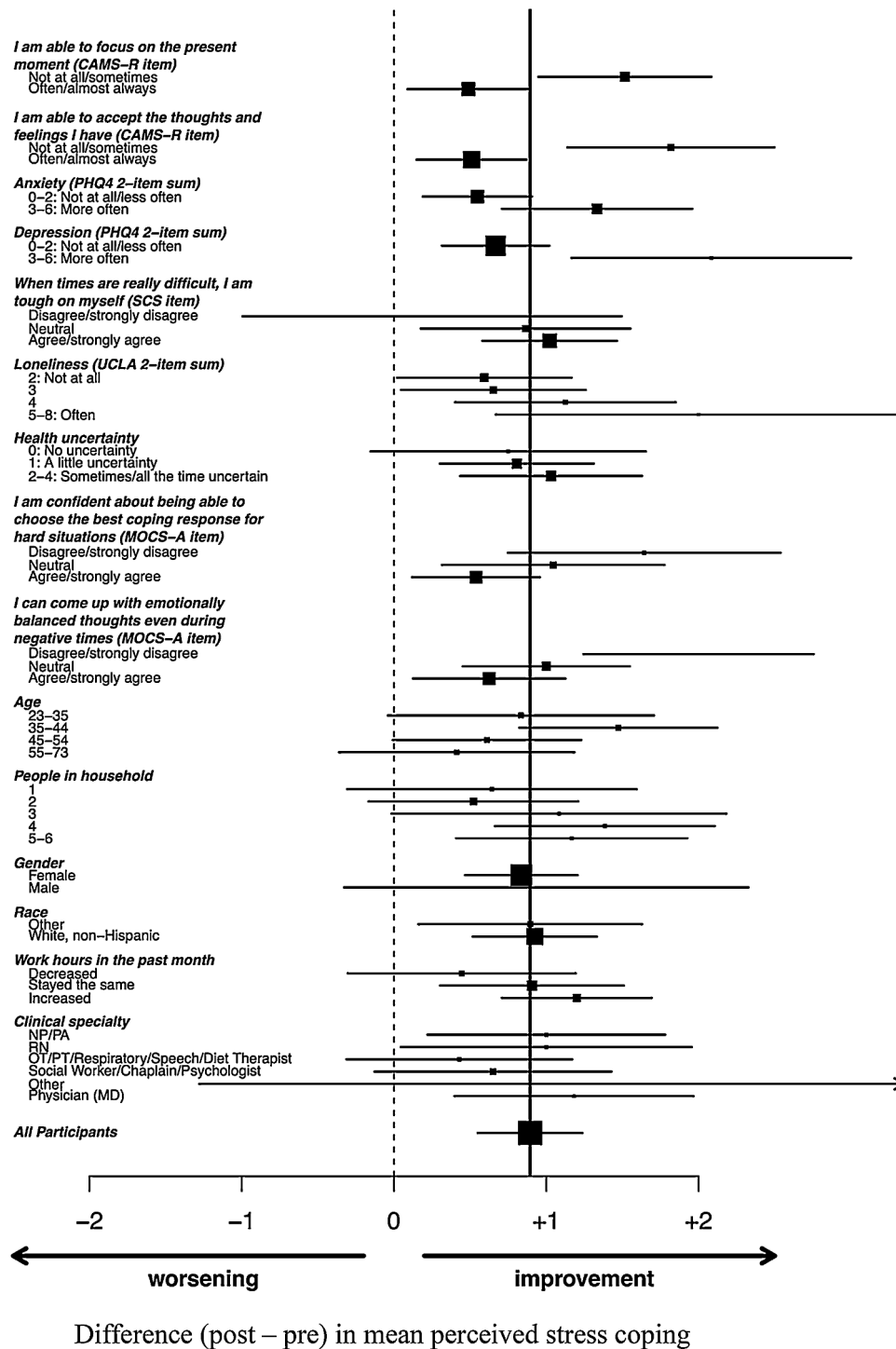


Fig. 1. Forest plot of the differential effect of each moderator. We present pairwise mean differences (solid squares) and 95% confidence intervals (horizontal lines). Larger/smaller squares corresponds with larger/smaller sample sizes. Vertical lines indicate no differential effect (dotted) and observed pairwise differential effect among all participants (solid).

Table 1
Results based on mixed-effects regression models for each potential moderator.

Variable*	Subgroup	Differential effect**	95% CI	p-value
Age	(per 10-year increase)	-0.10	[-0.37, 0.17]	0.46
People in household		0.19	[-0.04, 0.43]	0.10
Gender	Male	-0.04	[-1.27, 1.17]	0.94
	Female	Reference		
Race	Other†	0.12	[-0.63, 0.88]	0.75
	White, non-Hispanic	Reference		
Work hours in the past month	Decreased	-0.67	[-1.53, 0.19]	0.31
	Stayed the same	-0.26	[-1.00, 0.48]	
	Increased	Reference		
Clinical specialty	NP/PA	-0.27	[-1.42, 0.88]	0.53
	RN	-0.26	[-1.40, 0.87]	
	OT/PT/Respiratory/Speech/Diet Therapist	-0.85	[-1.98, 0.27]	
	Social Worker/Chaplain/Psychologist	-0.69	[-1.75, 0.35]	
	Other‡	0.18	[-1.41, 1.72]	
	Physician (MD)	Reference		
Focus on present moment	Not at all/sometimes	0.99	[0.36, 1.61]	<0.01
	Often/almost always	Reference		
Accept thoughts and feelings	Not at all/sometimes	1.32	[0.68, 1.97]	<0.01
	Often/almost always	Reference		
Anxiety		0.25	[0.08, 0.42]	0.01
Depression		0.42	[0.21, 0.62]	<0.01
Tough on myself		0.41	[0.05, 0.77]	<0.01
Loneliness		0.36	[0.08, 0.63]	0.01
Health uncertainty		0.18	[-0.21, 0.56]	0.37
Ability to choose coping response		-0.45	[-0.83, -0.06]	0.02
Emotionally balanced thoughts		-0.39	[-0.82, 0.05]	0.08

*Focus on present moment: Cognitive and Affective Mindfulness Scale- Revised; Accept thoughts and feelings: Cognitive and Affective Mindfulness Scale-Revised; Anxiety: Patient Health Questionnaire -4; Depression: Patient Health Questionnaire-4; Tough on myself: Self-Compassion Scale; Loneliness: UCLA Loneliness Scale; Ability to choose coping response: Measure of Current Status Questionnaire; Emotionally balanced thoughts: Measure of Current Status Questionnaire

**Differential effect corresponds to the difference in average perceived stress coping improvement for subgroup vs. reference group (for categorical variables) or per one-unit increase in variable (for continuous variables), except as noted for age, which corresponds to a per 10-year increase. Positive/negative differential effect indicates more/less improvement in average perceived stress coping, whereas a differential effect of zero indicates no association between the moderator variable and average perceived stress coping improvement.

†Includes 2 American Indian or Alaska Native, 9 Asian, 4 Black or African American, 7 Hispanic, and 1 Other

‡Includes 4 Technicians/Translators, 3 Advance Practice Clinicians/Midwives/Nurse Anesthetists, and 4 Other.

other MOCS-A item, emotionally balanced thoughts) moderated changes in perceived stress coping. Specifically, individuals who perceived themselves as benefiting more, in terms of perceived stress coping, from the intervention were generally less mindful and self-compassionate, but more anxious, depressed, and lonely at baseline. None of the questions assessing sociodemographic variables, work characteristics, or baseline health uncertainty moderated changes in perceived stress coping.

5. Discussion

We found that FCs who benefited the most from the SMART-3RP program, evidenced by their level of improvement in perceived stress coping, were generally less mindful and self-compassionate and more anxious, depressed, and lonely before initiating the program. Conversely, sociodemographic, work characteristics and baseline health uncertainty did not moderate improvements in perceived stress coping for FCs. These findings build upon the growing literature surrounding the impact of the COVID-19 pandemic on mental health outcomes. Research has found that negative mental health outcomes, such as depression and anxiety, may persist into post-lockdown environments (Woon et al., 2020). Therefore, there is a great need to identify effective prevention and early intervention methods during an emergency response to improve psychological outcomes (Sidi, 2020). As FCs are especially vulnerable to poorer mental health outcomes during the pandemic, future efforts to design sustainable interventions should focus on identifying moderating factors that may strengthen protective mental health mechanisms, such as resiliency and stress coping (Gavin et al., 2020). While our study is one of the first, to our knowledge, to evaluate the moderating factors of stress coping in the early stages of the COVID-19 pandemic, creating frameworks to support mental health should be a priority at this time (Rauch et al., 2020).

These findings had several limitations. First, to reduce survey burden, we only selected single items from empirically validated scales (e.g., Current Experiences Scale). Additionally, the generalizability of study results was limited by the lack of demographic diversity and overall low scores on items suggesting that participants were generally psychologically stable at baseline. Given this latter point, it is possible that the FCs who may need this program most did not register for it voluntarily and therefore, a larger proportion of FCs may have been less psychologically stable than the study accounted for and could have benefited more from this intervention. Strengths of the study includes the diversity in clinician specialty and institutional setting (i.e., hospital affiliation) among the sample as well as the implementation of SMART-3RP during the initial peak of COVID-19. The latter allowed researchers to examine the impact of the intervention to improve perceived stress coping at the height of the pandemic. Overall, our analyses add to the growing body of knowledge on the impact of psychological stressors in the midst of an ongoing pandemic and shed light onto which FCs may benefit most from a stress management intervention during an emergency response.

Declaration of Competing Interest

Dr. Sylvia receives funding from NIH, PCORI, Samsung and Takeda and royalties from New Harbinger.

Dr. Fava receives the following funding:

Research Support:

Abbott Laboratories; Acadia Pharmaceuticals; Alkermes, Inc.; American Cyanamid; Aspect Medical Systems; AstraZeneca; Avanir Pharmaceuticals; AXSOME Therapeutics; BioClinica, Inc; Biohaven; BioResearch; BrainCells Inc.; Bristol-Myers Squibb; CeNeRx BioPharma; Cephalon; Cerecor; Clarus Funds; Clexio Biosciences; Clintara, LLC; Covance; Covidien; Eli Lilly and Company;EnVivo Pharmaceuticals, Inc.; Euthymics Bioscience, Inc.; Forest Pharmaceuticals, Inc.; FORUM Pharmaceuticals; Ganeden Biotech, Inc.; Gentelon, LLC; GlaxoSmithKline; Harvard Clinical Research Institute; Hoffman-LaRoche; Icon

Clinical Research; Indivior; i3 Innovus/Ingenix; Janssen R&D, LLC; Jed Foundation; Johnson & Johnson Pharmaceutical Research & Development; Lichtwer Pharma GmbH; Lorex Pharmaceuticals; Lundbeck Inc.; Marinus Pharmaceuticals; MedAvante; Methylation Sciences Inc; National Alliance for Research on Schizophrenia & Depression (NARSAD); National Center for Complementary and Alternative Medicine (NCCAM); National Coordinating Center for Integrated Medicine (NiiCM); National Institute of Drug Abuse (NIDA); National Institutes of Health; National Institute of Mental Health (NIMH); Neuralstem, Inc.; NeuroRx; Novartis AG; Organon Pharmaceuticals; Otsuka Pharmaceutical Development, Inc.; PamLab, LLC.; Pfizer Inc.; Pharmacia-Upjohn; Pharmaceutical Research Associates, Inc.; Pharmavite® LLC; PharmRx Therapeutics; Photothera; Premiere Research International; Reckitt Benckiser; Roche Pharmaceuticals; RCT Logic, LLC (formerly Clinical Trials Solutions, LLC); Sanofi-Aventis US LLC; Shenox Pharmaceuticals, LLC; Shire; Solvay Pharmaceuticals, Inc.; Stanley Medical Research Institute (SMRI); Synthelabo; Taisho Pharmaceuticals; Takeda Pharmaceuticals; Tal Medical; VistaGen; Wyeth-Ayerst Laboratories

Advisory Board/ Consultant:

Abbott Laboratories; Acadia; Aditum Bio Management Company, LLC; Affectis Pharmaceuticals AG; Alfasigma USA, Inc.; Alkermes, Inc.; Altimate Health Corporation; Amarin Pharma Inc.; Amorsa Therapeutics, Inc.; Angelini S.p.A; Aptinix Inc.; Arbor Pharmaceuticals, LLC; Aspect Medical Systems; AstraZeneca; Auspex Pharmaceuticals; Avani Pharmaceuticals; AXSOME Therapeutics; Bayer AG; Best Practice Project Management, Inc.; Biogen; BioMarin Pharmaceuticals, Inc.; BioXcel Therapeutics; Biovail Corporation; Boehringer Ingelheim; Boston Pharmaceuticals; BrainCells Inc; Bristol-Myers Squibb; Cambridge Science Corporation; CeNeRx BioPharma; Cephalon, Inc.; Cerecor; Clexio Biosciences; Click Therapeutics, Inc; CNS Response, Inc.; Compellis Pharmaceuticals; Cypress Pharmaceutical, Inc.; DiagnoSearch Life Sciences (P) Ltd.; Dainippon Sumitomo Pharma Co. Inc.; Dov Pharmaceuticals, Inc.; Edgemont Pharmaceuticals, Inc.; Eisai Inc.; Eli Lilly and Company; ElMindA; EnVivo Pharmaceuticals, Inc.; Enzymotec LTD; ePharmaSolutions; EPIX Pharmaceuticals, Inc.; Esthismos Research, Inc.; Euthymics Bioscience, Inc.; Evecxia Therapeutics, Inc.; ExpertConnect, LLC; FAAH Research Inc.; Fabre-Kramer Pharmaceuticals, Inc.; Forest Pharmaceuticals, Inc.; Forum Pharmaceuticals; Gate Neurosciences, Inc.; GenOmind, LLC; GlaxoSmithKline; Grunenthal GmbH; Happify; H. Lundbeck A/S; Indivior; i3 Innovus/Ingenix; Intracellular; Janssen Pharmaceutica; Jazz Pharmaceuticals, Inc.; JDS Therapeutics, LLC; Johnson & Johnson Pharmaceutical Research & Development, LLC; Knoll Pharmaceuticals Corp.; Labopharm Inc.; Lorex Pharmaceuticals; Lundbeck Inc.; Marinus Pharmaceuticals; MedAvante, Inc.; Merck & Co., Inc.; MSI Methylation Sciences, Inc.; Naurex, Inc.; Navitor Pharmaceuticals, Inc.; Nestle Health Sciences; Neuralstem, Inc.; Neurocrine Biosciences, Inc.; Neuronetics, Inc.; NextWave Pharmaceuticals; Niraxx Light Therapeutics, Inc; Northwestern University; Novartis AG; Nutrition 21; Opiant Pharmaceuticals; Orexigen Therapeutics, Inc.; Organon Pharmaceuticals; Osmotica; Otsuka Pharmaceuticals; Ovid Therapeutics, Inc.; PamLab, LLC.; Perception Neuroscience; Pfizer Inc.; PharmaStar; Pharmavite® LLC.; PharmRx Therapeutics; Polaris Partners; Praxis Precision Medicines; Precision Human Biolaboratory; Prexa Pharmaceuticals, Inc.; Protagenic Therapeutics, Inc; PPD; PThera, LLC; Purdue Pharma; Puretech Ventures; PsychoGenics; Psylin Neurosciences, Inc.; RCT Logic, LLC (formerly Clinical Trials Solutions, LLC); Relmada Therapeutics, Inc.; Rexahn Pharmaceuticals, Inc.; Ridge Diagnostics, Inc.; Roche; Sanofi-Aventis US LLC.; Sentier Therapeutics; Sepracor Inc.; Servier Laboratories; Schering-Plough Corporation; Shenox Pharmaceuticals, LLC; Solvay Pharmaceuticals, Inc.; Somaxon Pharmaceuticals, Inc.; Somerset Pharmaceuticals, Inc.; Sonde Health; Sunovion Pharmaceuticals; Supernus Pharmaceuticals, Inc.; Synthelabo; Taisho Pharmaceuticals; Takeda Pharmaceutical Company Limited; Tal Medical, Inc.; Tetrigenex; Teva Pharmaceuticals; TransForm Pharmaceuticals, Inc.; Transcept Pharmaceuticals, Inc.; University of Michigan, Department of Psychiatry; Usona Institute, Inc.; Vanda Pharmaceuticals,

Inc.; Versant Venture Management, LLC; VistaGen

Speaking/Publishing:

Adamed, Co; Advanced Meeting Partners; American Psychiatric Association; American Society of Clinical Psychopharmacology; AstraZeneca; Belvoir Media Group; Boehringer Ingelheim GmbH; Bristol-Myers Squibb; Cephalon, Inc.; CME Institute/Physicians Postgraduate Press, Inc.; Eli Lilly and Company; Forest Pharmaceuticals, Inc.; GlaxoSmithKline; Global Medical Education, Inc.; Imedex, LLC; MGH Psychiatry Academy/Primedia; MGH Psychiatry Academy/Reed Elsevier; Novartis AG; Organon Pharmaceuticals; Pfizer Inc.; PharmaStar; United BioSource, Corp.; Wyeth-Ayerst Laboratories.

Stock/Other Financial Options: Equity Holdings: Psy Therapeutics

Royalty/patent, other income:

Patents for Sequential Parallel Comparison Design (SPCD), licensed by MGH to Pharmaceutical Product Development, LLC (PPD) (US_7840419, US_7647235, US_7983936, US_8145504, US_8145505); and patent application for a combination of Ketamine plus Scopolamine in Major Depressive Disorder (MDD), licensed by MGH to Biohaven. Patents for pharmacogenomics of Depression Treatment with Folate (US_9546401, US_9540691).

Copyright for the MGH Cognitive & Physical Functioning Questionnaire (CPFQ), Sexual Functioning Inventory (SFI), Antidepressant Treatment Response Questionnaire (ATRQ), Discontinuation-Emergent Signs & Symptoms (DESS), Symptoms of Depression Questionnaire (SDQ), and SAFER; Lippincott, Williams & Wilkins; Wolters Kluwer; World Scientific Publishing Co. Pte.Ltd.

All disclosures can also be view online at:

<https://mghcme.org/faculty> > Maurizio Fava, MD > View Bio > View Disclosures Here

Dr. Sabine Wilhelm is a presenter for the Massachusetts General Hospital Psychiatry Academy in educational programs supported through independent medical education grants from pharmaceutical companies; she has received royalties from Elsevier Publications, Guilford Publications, New Harbinger Publications, Springer, and Oxford University Press. Dr. Wilhelm has also received speaking honoraria from various academic institutions and foundations, including the International Obsessive Compulsive Disorder Foundation, Tourette Association of America, and Brattleboro Retreat. In addition, she received payment from the Association for Behavioral and Cognitive Therapies for her role as Associate Editor for the Behavior Therapy journal, as well as from John Wiley & Sons, Inc. for her role as Associate Editor on the journal Depression & Anxiety. Dr. Wilhelm has also received honorarium from One-Mind for her role in PsyberGuide Scientific Advisory Board. Dr. Wilhelm has received salary support from Novartis and Koa Health.

Ms. George, Dr. Rabideau, Dr. Streck, Ms. Albury, Dr. Hall, Dr. Luberto, Ms. Mizrach, Dr. Perez, Ms. Crute, Dr. Mehta, Ms. Convery, Dr. Looby, Dr. Fricchione and Dr. Park have no disclosures to report.

Acknowledgements

TBD

We certify that all authors have seen and approved the final version of the manuscript being submitted. This article is based on original work and has not received prior publication or is under consideration for publication elsewhere.

References

- Lu, W, Wang, H., Lin, Y., Li, L., 2020. Psychological status of medical workforce during the COVID-19 pandemic: a cross-sectional study. *Psychiatry Research*.
- Pappa, S, Ntella, V., Giannakas, T., Giannakouli, V.G., Papoutsis, E., Katsaounou, P., 2020. Prevalence of depression, anxiety, and insomnia among healthcare workers during the COVID-19 pandemic: a systematic review and meta-analysis. *Brain Behav. Immun.*
- Mehta, DH PG, Traeger, L, Park, ER, Goldman, RE, Haime, V, et al., 2016. Building resiliency in a palliative care team: a pilot study. *J 2016 Pain Symptom Manag.* 51, 604–608.

- Park, ER, PG, Millstein, RA, Luberto, CM, Traeger, L, Proszynski, J, et al., 2020. A virtual resiliency intervention promoting resiliency for parents of children with learning and attentional disabilities: a randomized pilot trial. *Matern Child Health J.* 24, 39–53.
- Park, ER, Sylvia, L., Streck, J.M., Luberto, C.M., Stanton, A., Perez, G.K., Denninger, J., 2020. Launching a resiliency group program to assist frontline clinicians in meeting the challenges of the COVID-19 pandemic: Results of a hospital-based systems trial. *General Hospital Psychiatry.*
- Dunkley, DM, Lewkowski, M., Lee, I.A., Preacher, K.J., Zuroff, D.C., Berg, J.L., Westreich, R., 2017. Daily stress, coping, and negative and positive affect in depression: Complex trigger and maintenance patterns. *Behavior Therapy* 48 (3), 349–365.
- Thimm, JC, Wang, C.E., Waterloo, K., Eisemann, M., Halvorsen, M., 2018. Coping, thought suppression, and perceived stress in currently depressed, previously depressed, and never depressed individuals. *Clinical Psychol. Psychotherapy* 25 (3), 401–407.
- Rettie, H, Daniels, J., 2020. Coping and tolerance of uncertainty: predictors and mediators of mental health during the COVID-19 pandemic. *American Psychologist.*
- Carver, CS., 2006. Measure of Current Status.
- Antoni, MH, Lechner, S.C., Kazi, A., Wimberly, S.R., Sifre, T., Urcuyo, K.R., Phillips, K., Gluck, S., Carver, C.S., 2006. How stress management improves quality of life after treatment for breast cancer. *J. Consult. Clin. Psychol.* 74, 1143–1152.
- Feldman, G, Heyes, A.K.S., Green, J., Laurenceau, J., 2007. Mindfulness and emotional regulation: The development and initial validation of the cognitive and affective mindfulness scale-revised (CAMS-R). *J. Psychopathology Behav. Assess* 29 (2), 177–190.
- Kroenke, K, Spitzer, R.L., Williams, J.B., Löwe, B., 2009. An ultra-brief screening scale for anxiety and depression: the PHQ-4. *Psychosomatics* 50 (6), 613–621 doi: 10.1176/appi.psy.50.6.613. PMID: 19996233.
- Neff, KD., 2003. The development and validation of a scale to measure self-compassion. *Self Identity* 2 (3), 223–250.
- Russell, D, Peplau, L.A., Ferguson, M.L., 1978. Developing a measure of loneliness. *J. Pers. Assess.* 42, 290–294.
- Rogers, SN, Cross, B., Talwar, C., Lowe, D., Humphris, G., 2016. A single-item screening question for fear of recurrence in head and neck cancer. *Eur. Archives Otorhinolaryngol.* 273 (5), 1235–1242.
- Yanez, BR SA, Hoyt, MA, Tennen, H, Lechner, S., 2011. Understanding perceptions of benefit following adversity: how do distinct assessments of growth relate to coping and adjustment to stressful events? *J. Soc. Clin. Psychol.* 699–721.
- Woon, LS SH, Nik Jaafar, NR, Leong Bin Abdullah, MFI, 2020. Mental health status of university healthcare workers during the COVID-19 pandemic: a post-movement lockdown assessment. *Int. J. Environ. Res. Public Health* 17.
- Sidi, H., 2020. The psychological sequelae during mental health and Covid-19 Pandemic: Learning from the past for today's coping styles. *Med. Health* 15 (1), 1–4. June2020; 15:1-4.
- Gavin, B, Lyne, J., McNicholas, F., 2020. Mental health and the COVID-19 pandemic. *Irish J. Psychol. Med.* 37, 156–158.
- Rauch, SA, Simon, N.M., Rothbaum, B.O., 2020. Rising tide: responding to the mental health impact of the COVID-19 pandemic. *Depress. Anxiety* 37, 505–509.