

Coping strategies to prevent or reduce stress and burnout among oncology physicians: a systematic review

Estratégias de enfrentamento para prevenir ou reduzir o estresse e o *burnout* entre médicos oncologistas: uma revisão sistemática

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ABSTRACT

The purpose of this systematic review (SR) was to identify interventions that are effective to prevent or reduce stress and burnout among oncologists. Search was conducted in eight electronic databases and grey literature databases, with no language or time restrictions. Included studies involved medical oncologists and contained interventions to prevent or deal with stress or burnout with outcomes assessment. In two selection phases process, 19 out of 3,020 studies were included. Risk of bias was low for nine studies, moderate for six studies and high for four ones. Certainty of evidence was considered low and very low for the analyzed outcomes. Interventions varied a lot and those which had a significant effect in stress and burnout reduction among oncologists were experience sharing between female doctors in virtual groups, integrative meetings outside the work environment, and team sessions supervised by counselors. Although interventions had variable effects on reducing or preventing burnout and stress, mores studies are needed due to outcomes low evidence.

Keywords: Oncologists; Burnout, Professional; Stress, Psychological; Systematic review; Evaluation of the efficacy-effectiveness of interventions.

RESUMO

O objetivo desta revisão sistemática (RS) foi identificar intervenções eficazes para prevenir ou reduzir o estresse e o burnout entre oncologistas. A busca foi realizada em oito bases de dados eletrônicas e bases de dados de literatura cinzenta, sem restrições de idioma ou tempo. Os estudos incluídos envolveram médicos oncologistas e continham intervenções para prevenir ou lidar com o estresse ou *burnout* com avaliação de resultados. Em duas fases de seleção, 19 dos 3.020 estudos foram incluídos. O risco de viés foi baixo para nove estudos, moderado para seis estudos e alto para quatro. A certeza da evidência foi considerada baixa e muito baixa para os desfechos analisados. As intervenções variaram muito e as que tiveram efeito significativo na redução do estresse e do *burnout* entre os oncologistas foram o compartilhamento de experiências entre médicas em grupos virtuais, reuniões integrativas fora do ambiente de trabalho e sessões de equipe supervisionadas por conselheiros. Embora as intervenções tenham efeitos variáveis na redução ou prevenção de *burnout* e estresse, mais estudos são necessários devido à baixa evidência dos resultados.

Descritores: Oncologistas; Burnout, Profissional; Estresse Psicológico; Revisão sistemática; Avaliação da eficácia-efetividade das intervenções.

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INTRODUCTION

Oncology may be viewed as an extraordinary rewarding specialty.^[1] However, its practice poses many challenges, such as overload from continuous evolution of therapeutic possibilities, increasing bureaucratization and workload due to the increasing incidence of cancer in the population. In addition, oncologists face uncertainties about patients' responses to treatment and have intense involvement with critically ill and/or end-of-life patients, having to handle their own and others' emotions.^[1-3] As a consequence, they are generally under stress and may have burnout.

Burnout is a work-related psychological syndrome characterized by emotional exhaustion, depersonalization and reduced personal accomplishment. It may occur in persons who work with other people when occupational pressures persist over time. Burnout syndrome was first described in the 1970s and has been extensively studied by Dr. Maslach et al. (1986),[4] who developed the Maslach burnout inventory (MBI), a diagnostic method still used in current studies.^[4-6] Its personal and professional consequences include increased incidence of illnesses, car accidents, divorces, obesity, alcoholism, depression, drug use, higher rates of absenteeism, early retirements, difficulty in recruiting and maintaining professionals, decrease in the quality of services and patient satisfaction, increase in medical errors and increase in medical errors and in the number of sues and processes against the doctor.[4]

Data provided by oncologists' professional associations are alarming. Study conducted by the American Society of Clinical Oncology (ASCO) detected that 45% of oncologists had already experienced burnout symptoms. [7] And, a study in 41 European countries by the European Society of Medical Oncology (ESMO) found the percentage of 71% of burnout among oncologists aged up to 40 years. [2]

A prevalence of 58% was found among oncologist who worked in a Brazilian hospital.^[8] The awareness of having burnout syndrome is difficult because many of its signs and symptoms are interpreted as personal failures. When professionals feel unable to manage their conflicts, they may delay looking for help and being diagnosed and treated for burnout.^[9,10]

In 2016, a meta-analysis assessed the effectiveness of various interventions to prevent and reduce physicians' burnout. [9,11] Several approaches proved to be effective, but authors called attention to the need of further studies to establish the best strategies for different realities within the medical profession. [12] We have therefore conducted this systematic review (SR) to answer the following question: "What interventions are effective to prevent or reduce oncologists' signs and symptoms of stress and burnout when compared to oncologists without these interventions?".

METHODS

Protocol and registration

A systematic review protocol was elaborated based on the preferred reporting items for systematic reviews and meta-analyses protocols (PRIS-MA-P),^[13] and registered at the prospective register of systematic reviews (PROSPERO),^[14] available under registration number CRD42019141517. We followed the preferred reporting items for systematic reviews and meta-analyses checklist (PRISMA)^[15] and synthesis without meta-analysis (SWiM) reporting items^[16] for reporting this SR.

Eligibility criteria and search strategy

Based on the acronym PICOS, the participants were oncologist physicians, the interventions included those to prevent and reduce stress and burnout; the comparison (control) were with oncologists who had not undergone these interventions; outcomes included quality of life, and signs and symptoms of stress and burnout; and, the study types included randomized and quasi-randomized clinical trials.

Initially, with assistance of experienced librarians, a search strategy was developed to identify studies that contemplated interventions to deal with stress and burnout among oncologists. The strategy included keywords such as "burnout", "stress", "oncologists", "interventions" and its synonyms. The full search is detailed in Appendix 1.

The search was applied in the following data-bases: CINAHL, Cochrane Library, EMBASE, LILACS, PsycINFO, PubMed, Scopus and Web of Science. In addition, the grey literature was searched on Google Scholar, ProQuest Dissertation and Theses and Open Grey. The reference list of the articles found was carried out manually, and the main authors of the subject were contacted, asking for possibly non included articles to complement the search. Filters for languages and restrictions on the date of publication were not used.

According to pre-elaborated criteria, studies were included if they involved oncologists, either clinical oncologists, radio-oncologists, oncology surgeons, pediatric oncologists or onco-hematologists; and, contained specific data on interventions to prevent or deal with stress or burnout.

Studies were excluded if they: 1) involved only non-medical oncologist professionals or medical students; 2) did not involve interventions to prevent or handle with stress and burnout; 3) had duplicated data from another included study or insufficient data; 4) were conducted in animals; 5) were reviews, letters, books, case report, case series, opinion article, technique articles and guidelines; 6) did not have their complete text available online/published and if the texts were not accessible after three contact attempts in a 15-day period by electronic mail to corresponding authors.



Study selection

Articles found in the databases were organized in the EndNote X9 program. Two reviewers (A.S. and C.W.) selected the articles independently in two phases. In phase-1, the two reviewers read the titles and abstracts applying the eligibility criteria using an online software (Rayyan, Qatar Computing Research Institute). In phase-2, the same reviewers read the full-text, also applying the eligibility criteria. In case of disagreement, in both phases, doubts were resolved by consensus and, if incompatibility remained, a third reviewer (J.M.D.O.) was called.

Data collection process and data items

The first and second reviewers (A.C.S. and C.W.) collected the main information from the selected studies independently. After that, the collected information was cross-checked, and its accuracy confirmed in a consensus meeting. In case of disagreement, conflict was resolved with a final decision by the third reviewer (J.M.D.O).

Criteria for data extraction were determined prior to the review, using a table that included: author, year, participants, country, type of study, primary and secondary objectives, type of intervention, results and outcomes, participants' acceptability, satisfaction of the intervention and intervention effectiveness If data were not found in the article, three contact attempts were tried in a 15-day period by electronic mail to corresponding authors to obtain relevant unpublished information.

Risk of bias in included studies

The risk of bias (RoB) of included studies was evaluated using the Joanna Briggs Institute (JBI) critical appraisal checklist for quasi-experimental,[17] the IBI critical appraisal checklist for analytical cross-sectional studies[18] for descriptive studies, Cochrane risk of bias tool RoB 2.0 for randomized trials and Cochrane risk of bias tool RoB 2.0 for cluster-randomized trials.[19] Judgement was made by two independent reviewers (A.C.S. and C.W.) and decisions about scoring were agreed by both reviewers before critical appraisal assessments. The RoB was characterized by the reviewers as high when the study reached up to 49% of bias, moderate when the study reached 50% to 69%, and low when the study reached more than 70%. All RoB figures and plots were created using robvis.^[20]

Summary measures

Stress levels and burnout symptoms were considered as the main outcome and the analysis was not restricted by any method for measuring or diagnosing them

No restrictions were made on how these outcomes were measured or whether they were obtained by psychologists or by self-assessment.

The extracted data were synthesized in a descriptive manner. Studies including any type of intervention and any number of doctors were accepted.

Synthesis of results

Heterogeneity within studies was evaluated either by the inconsistency index (I²) statistical test or by their clinical, methodological and statistical characteristics.^[21]

Meta-analysis was considered inappropriate due to the included studies heterogeneity in clinical and methodological characteristics. The synthesis of results was also descriptive.

Confidence in cumulative evidence

A summary of the overall confidence in cumulative evidence available by outcomes analyzed was presented using "grading of recommendations assessment, development and evaluation" (GRADE). Summary of Findings (SoF) table was produced using the GRADE online software.^[22]

RESULTS

Study selection

In phase-1, 3,020 citations were identified from electronic databases. After removing duplicated records, a total of 2,067 titles and abstracts were evaluated with the eligibility criteria. Following phase-1, 197 articles entered phase-2. After full-text reading, 19 articles fulfilled the eligibility criteria and were included for qualitative analysis (see on Appendix 2 the exclusions and their reasons). Quantitative analysis was not possible considering heterogeneity. A detailed flowchart of this process is shown in Figure 1.

Study characteristics

The 19 included studies were conducted in thirteen countries. One in Australia, [23,24] Austria, [23] Belgium, [25] Canada, [26,27] Costa Rica, [28] Germany, [23,29] Israel, [30,31] Italy, [32-34] the Netherlands, [35] New Zealand, [23] Switzerland, [23] United Kingdom, [36,37] and five in the United States. [31,38-41]

All included studies evaluated samples with at least one group of medical oncology doctors (fellows, residents, or specialist physicians) and the total of 1,513 individuals were analyzed. They included oncology care providers as specialist physicians (oncology, onco-hematology, pediatric, palliative care),[23-26,28,29,31,33,38] oncology residents,[27,30,40] oncology and onco-hematology fellows,[24,24,28,29,31,34,36-39,41] nurses,[28,31-35] radiotherapy assistants,[24,35] psychologists, and social workers.[31]

A summary of the descriptive characteristics of the various studies can be found in Tables 1 and 2.

In regard to the study design, nine were quasirandomized studies, [26-28,30,33,34,37,40,41] one was crosssectional, [38] and nine were RCT (two cluster-randomized trials [35,39] and seven individually-randomized parallelgroup trial). [23-25,29,31,32,36]



The interventions to prevent and/or reduce stress levels and burnout included mostly class-rooms^[27,31,32,35,36] (in four studies totalizing 15 hours

each), workshops^[23,24,28] (in three studies for approximately 14 hours) and group sessions^[25,29,30,33,34,37,41] (in six studies totalizing 16 hours each).

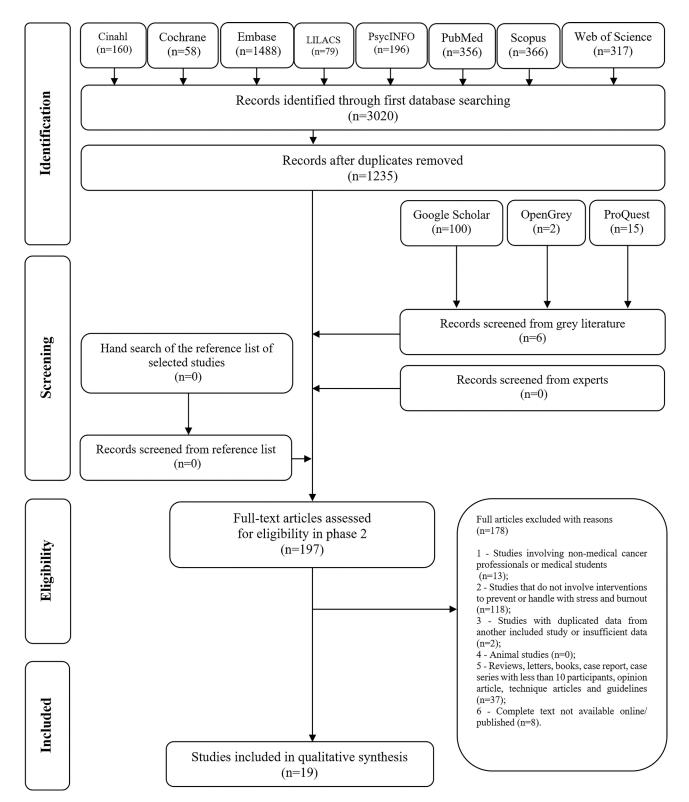


Figure 1. Flow diagram of literature search and selection criteria. (Adapted from PRISMA - Moher et al. (2010):^[15] preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement).

Table 1. Summary of population characteristics of included articles (n=19).

(Country) design Bar-Sela, Lulav- Grinwald Before -after 201230 (Israel) Barzelloni et al, 201432	Participants / Context / Setting	Groups (n/%)	Outcomes measured	Measures
a, Id :nik, oni et j³²				
oni et ₃₂	15 Oncology residents¹ Division of Oncology, Rambam ter Health Care Campus, and Faculty of Medicine, Technion-Israel Institute of Technology	IG Junior: 1° part of residency (<3y) (8/53.3*); Senior: 2° part of residency (>3y) (7/46.7*)	Burnout measures (Emocional Exaustion and Despersonalization), Communication skills and self-awareness	MBI and expectations questionnaire that was completed at the beginning and at the end of the year
(Italy)	34 Physicians and nurses Departments of Medical Oncology the Policlinico Tor Vergata University hospital and John of Procida Salerno	IG (12/35.3*) CG (22/64.7*)	Burnout and Health at T0 and quarterly (T1, T2, T3, T4).	General Health Questionnaire (GHQ) and MBI -
Bragard et al, 2010 ²⁵ (Belgium)	62 Cancer physicians specialists All Belgian specialists working in cancer care were invited	IG (29/46.8*) CG group (33/53.2*)	Burnout, communication skills, contextual variables	MBI; Standardized Breaking Bad News Simulated Interview for assessment of communication skills; Socioprofessional questionnaire and the Job Stress Survey (JSS)
Brown et al, 2014 ²³ (Australian/ New Zealand/ RCT Swiss/ German/ Austrian)	62 Oncologists 21 from 10 Australian/ New Zealand (ANZ) centers and 41 from 10 Swiss/ German/ Austrian (SGA) centers	IG: 1-day workshop group (NR) cg (NR)	Communication behavior, stress and satisfaction	N R
Bui et al, 2021³³ Before -after (Italy)	28 participants among medical doctors, nurses, socio-sanitary assistant, biologists, support, and administrative staff Oncology Department Oncology Units (Recovery Ward and Day Hospital)	Medical doctors (7/23.5%) Nurses (8/29.4%)	Burnout effective, lifestyle and work factors	MBI; B-C Working Fit and Socio demographic questionnaire
Butow et al, 2008 ²⁴ RCT (Australia)	30 medical and radiation oncologists from six tertiary care hospitals in six Australian cities	IG (16/53.3*) CG (14/46.7*)	Doctor behaviors	MBI and Demographic, previous training and current practice assessed at baseline
Clemons et al, 2019 ²⁶ Before -after (Canada)		1 group (13/100*)	Physician, burnout happiness and compliance with the virtues	Abbreviated MBI and Oxford Happiness Questionnaire
Dahn et al, Before -after 2019 ²⁷ (pilot study)	9 participants – 8 residents and 1 staff ter Centre for Radiation Oncology residents. کایا Dalhousie University, Halifax, NS	Resident group (8/88.9%*) Staff group (1/11.1%*)	Burnout, Resiliency, welness	

continue					
Author, Year (Country)	Study design	Participants / Context / Setting	Groups (n/%)	Outcomes measured	Measures
Graff et al, 2018³³ (United States)	Cross Sectional	169 Female oncologist/ hematologist (H/O), pediatric, radiation oncology, surgical specialties, and palliative care from a Facebook closed group	1 group (169/100*)	Burnout, Career satisfaction	12-question online survey using a visual analog scale
Italia et al, 2007³⁴ (Italy)	Before -after	65 Doctors (50.77%*) and nurses of oncology unit in two hospital units of Catania	Group A –16 doctors and 16 nurses of an adult oncology unit (32/49.2*) Group B –17 doctors and 16 nurses of a pediatric oncology unit (33/50.8*)	Burnout	MBI
Kesselheim et al, 2020³³ (United States)	Cluster RCT	19 pediatric hematology-oncology fellowship programs during the 2016-2017 academic year – 100 fellows from All PHO fellowship training programs in the United States	59 intervention and 41 usual training fellows	Pediatric Hematology- Oncology Self- Assessment in Humanism (PHOSAH)	Pediatric Hematology-Oncology Self- Assessment in Humanism (PHOSAH), MBI, Patient-Provider Orientation Scale (PPOS), Empowerment at Work Scale, and a 5-point satisfaction scale
Landaverde et al, 2018 ²⁸ (Costa Rica)	Before -after	Medical oncologists (155 members including hematologists, medical oncologists, pharmacists, laboratory personnel, nurses and secretaries) from medical oncology team at Mexico Hospital, San Jose Costa Rica	N R	Burn out	MBI
Le Blanc et al, 2007 ³⁵ (the Netherlands)	Cluster RCT	664 care providers (physicians, nurses, and radiotherapy assistants) working in direct care for oncology patients.of 29 oncology wards from the Netherlands	IG (260/39.2%) CG (404; 60.8%)	Burnout and Association of burnout and the level of social support, job control, and participation in decision making	MBI and other questionnaires about work situation and well-being
Mache et al, 2017 ²⁹ (Germany)	RCT	80 German-speaking employed junior physicians working in clinic departments of oncology and hematology medicine hospital departments in Germany	IG (39/48.75*) CG (CG) (41/51.25*)	Perceived stress Work-related health Psychosocial skills self- perceived training outcome and training design	Perceived Stress Questionnaire (PSQ), Copenhagen Psychosocial Questionnaire, MBI-emotional exhaustion (EE), and Emotion Regulation Skills Questionnaire-27
Medisauskaite and Kamau, 2019³ ⁶ (United Kingdom)	RCT	91 doctors - 54.9% (50) work in hospitals and 72.5% (66) work >41 h a week (6% oncologists) randomly selected NHS trusts, 9 royal colleges of medicine, and the British Medical Association (BMA)	Trial group 4 – Modules 1-4 / IG (39/*) Trial group 5 – Doctors not assigned to any module / CG (52/*)	Burnout, anxiety, insomnia, grief, alcohol/ drug use, binge eating, physical symptoms, and psychiatric morbidity	MBI, General Anxiety Disorder-7, the 12 items General Health Questionnaire, Texas Revised Inventory of Grief, Patient Health Questionnaire, Alcohol Use Disorder Identification Scale (AUDIT), Commonly Abused Drugs Charts, Insomnia Severity Index, 5 items from the Binge Eating Scale from the Eating Disorder Diagnostic Scale and the Physical Symptom Inventory



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Author, Year (Country)	Study design	Participants / Context / Setting	Groups (n/%)	Outcomes measured	Measures
Moody et al, 2013 ³¹ (United States and Israel)	RCT	47 (21% of oncologist physician and 53% of oncologist nurses) From the Children's Hospital of Montefiore in New York City and the Schneider Children's Hospital in Petach Tivka,Israel	IG (23/48.94*) CG (24/ 51.06*)	Burnout, Depression and perceived stress	MBI, Beck Depression Inventory and Perceived Stress Scale-14
Pathak, Eapen and Zell, 2019⁴⁰ (United States)	Before-after	NR number Oncology trainees from Hematology and Oncology Division of an academic Comprehensive Cancer Care center	The Henrietta Lacks Firm The Jane Wright Firm The Padmini Iyer Firm The Rita Mehta Firm NR numbers	Burnout	MBI
Sekeres et al, 2003 ⁴¹ (United States)	Non- randomized clinical trial	28 first-year hematology-oncology fellows (14 each academic year of 2000-2001 and 2001-2002) From Dana-Farber Cancer Institute/ Brigham and Women's Hospital (DFCI/BWH) and Massachusetts General Hospital (MGH)	Fellows that started at Dana- Farber Cancer Institute/Brigham and Women's Hospital (DFCI/ BWH) – CG and IG (14/50*) (7 each academic year of 2000-2001 and 2001-2002) Fellows that started at Massachusetts General Hospital (MGH) CG and IG(14/50*) (7 each academic year of 2000-2001 and 2001-2002)	Fellows' Fellows' perceptions on: how they related to patients and colleagues "attitudes" questionnaire - attitudes during the course of the first fellowship year	32-item attitudes questionnaire6, scored 1 to 5, at three time points during their first year (within the first week of the start of fellowship; during the sixth month of fellowship, just before the switch-over; and during the final month of the first year of fellowship)
Tjasink and Soosaipillai, 2018³' (England)	Before- after	16 doctors ⁷ (4 participants were from medical oncology, 4 from palliative care, 3 from clinical oncology and 3 from haematology) within a large teaching hospital in Central London	Single group	Burnout	MBI

Abreviattions: CI-confidence interval; d=days; CG=Control Group; IG=Intervention Group; h=hour; min=minutes; mo=month; MBI=Maslach Burnout Inventory; EE=Emotional exhaustion; RPA=Reduced personal accomplishment; Dp=Depersonalization; NR=Not reported; RCT=randomized-controlled trial; SD=Standard Deviation; U=Unclear; w=weeks; y=years; (*) data calculated by the authors.

Of the 17 residents that were participating, 2 decided not to continue after 2 meetings and therefore were excluded.

¹³ physicians completed the baseline scores, 11 completed Maslach/Oxford scores at the end of the study, and 8 the 1-month post-study assessment

Dropouts: at T2 the number of participants had dropped from 664 (experimental group: 260; control group: 404) to 376 (experimental group: 231; control group: 145), and at T3 it had dropped to 304 (experimental group: 208; control group: 96).

The study involved 227 doctors however the analysis considered in the present SR were Group 4 and control (91 doctors).

Additional information by emailing official authors.

The questionnaire derived from the Physician's Belief Scale; the American Academy on Physician and Patient evaluation; common objectives of Balint-like groups across the United States; barriers to physician

recognition of psychosocial aspects of health care reports; and from surveys of previous hematology-oncology fellows to explore attitudes toward patients, colleagues, and psychosocial issues



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Author, Year	Intervention	Final Diagnosis / After intervention Results before and	Conclusions	Limitations
Bar-Sela, Lulav-Grinwald and Mitnik, 2012 ³⁰ (Israel)	Balint-type case discussion groups 1.5 h monthly (9 sessions/y)	Before intervention/Baseline MBI parameter score: - EE: Junior: 3.66 / Senior: 3.14 - RPA: Junior: 1.33 / Senior: 1.34- Dp Junior: 2.6 / Senior: 0.98 After Follow-up 1y MBI parameter score: - EE: Junior: 3.67 / Senior: 3.48 - RPA: Junior: 2.13 / Senior: 1.48	No significant difference was found in regard to Burnout. However, Balint group may improve tresidents' communication abilities and contribute to their selfaccomplishment as doctors.	- Small sample size; - Limited number of group sessions
Barzelloni et al, 2014³² (Italy)	1 day of classroom training and discussion meetings on a monthly basis for a year T0 (previous) T4 after a year	MBI - Emociona Exaustion: statistically significant reduction (p \le 0.04) T0 versus T4 in the EG No statistical differences in the other MBI test two dimensions GHQ IG T0 vs. T4, p \le 0.00), IG vs. CG at the end of surgery (T4 p \le 0.01).	There was clinical significant implications	N N
Bragard et al, 2010 ²⁵ (Belgium)	19h BT (two 8h day sessions and one 3h evening session - 2h plenary session focusing on theoretical information in the form of two lectures and 17h of small-group role-playing sessions) CW (six sessions of 3h spread over a 3-month period)	Before intervention/Baseline MBI mean (SD): - EE: BT: 21 (7) vs BT with CW: 18 (8) - RPA: BT: 39 (5) vs BT with CW: 39 (6) - Dp: BT: 7 (4) vs BT with CW: 6 (5) After intervention – after 6 monts MBI mean (SD): - EE: BT: 22 (8) vs BT with CW: 18(10) - RPA: BT: 39 (3) vs BT with CW: 39 (4) - Dp: BT: 8 (5) vs BT with CW: 7 (6)	No difference found in Burnout scores. The authors consider that the amount of clinical workload and the overuse of some facilitative communication skills were associated with cancer physicians' burnout.	- Use of role play with direct feedback focusing mainly on the acquisition of communication skills oriented towards patient; voluntary participation (and thus highly motivated physicians); Small sample size
Brown et al, 2014 ²³ (Australian/ New Zealand/ Swiss/German/ Austrian)	1 day 7h interactive face-to-face workshop (2h of written and oral materials, 30 min of video modeling ideal behavior, 4h of role-play practice, and 30 min of individualized feedback on audiotaped consultations with actual patients)	Only AFTER 1 month of intervention CG versus IG ANZ: - IG: significant increase in collaborative communication after training; - CG: decline in use of collaborative behaviors during the study period. No effect of training on other doctor communication behaviors. Trained doctors did not demonstrate increased confidence in their information provision or reduced stress and burnout	The intervention was not sufficient to reduce stress and burnout	N

Author, Year	•			
	Intervention	Final Diagnosis / After intervention Results before and after intervention	Conclusions	Limitations
I	8-months intervention involved fortnight meetings by facilitators, incorporated elements of reflection, shared experiences and managing emotions; 3-hour meetings occurring once every two weeks with an expert team in building, communication strategies and emotional management; Topics addressed were organized into 4 modules: 1 Specific training focused on personal work experience and relationship between colleagues in each department 2 Individual counselling 3 Emotion management (residential	MBI percentage of participants with each levels (2 weeks before the intervention): - EE: Low: 29.4% High: 17.6% - RPA: NR - DP: NR - DP: NR - MBI percentage of participants with each levels(6mo) After the intervention: - EE: Low: 52.9%: High: 5.9% - RPA: no difference from TO - DP: no difference from TO	The intervention was not able to change Burnout, but there was dramatic reduction in high degree of EE (Emocional Exaustion), mainly in Day Hospital	- Limited sample size - No control group
	course) 4 sent-neip Groups activation			
	CST: 1.5-day intensive	Baseline		Small sample size of
	face-to-face workshop	MBl's median:	ls of	doctors and of institutions
	incorporating presentation of	- EE IG: 18.0 versus CG: 16.0	stress and burnout	making it i impossible to
	principles, a DVD modelling	- RPA IG: 40.0 versus CG: 40.0		stratify; -skill levels were
	ideal behavior and role-play	- Dp lG: 8.0 versus CG: 2.5		high at Baseline; over-
	practice, followed by four 1.5h	MBl's median:		representation of female
	video-conferences at monthly	- EE IG: 18.0 versus CG: 13.5		doctors in the sample;
	intervals incorporating role-	- RPA IG: 39.0 versus CG: 38.5		
	play of doctor-generated	- Dp IG: 6.0 versus CG: 3.0		

Conclusions	There was no improvement of happiness nor burnout reduction. Scores which significantly changed in self-rated virtue old program and the perance, and resolution perance, and resolution bernance or improvements in happiness and reduced burnout in physicians caring for cancer patients	The intervention was highly NR valued by residents and faculty, suggesting the importance of resilience education in the Radiation Oncology residency curriculum	Social media can be an effective venue to educate physicians, augment patient care via advice, foster and improve career satisfaction among female
Final Diagnosis / After intervention Results before and after intervention	Baseline MBI score: - EE: 7 - RPA: 16 - Dp: 4 - Dp: 4 - Happiness Questionnaire: 4.2 - TW (13w of follow-up and 1mo following study completion After interventio MBI score: - EE: 13w: 7 + 1 mo: 4.5 - RPA: 13w: 16 + 1 mo: 2 - Dp: 13w: 3 + 1 mo: 2 - Happiness Questionnaire: 13w: 4.7 + 1 mo: 4.7	Resident group: - Burnout rate: NR - Burnout rate: NR - Burnout rate: NR - Burnout rate: NR - Usefulness: 7.0-8.9/10 - Usefulness: 7.0-8.9/10 - Of the resident and staff session: 8.4/10 al activities rated from 8.3-8.8/10 - Follow up 3 mo - After After - After - After - After - After - After - After	- -
Intervention	Modified Franklin's 13 virtues (the next virtue was added to the previously listed virtues and scores were requested daily) - Each day during the 13-week program, oncologists were emailed a list of virtues to focus on and scored how they felt they were complying with them (5-point Likert scale was used instead of a simple yes/no to increase potential variability in responses)	2-hour resident seminar with discussions, mentorship and teaching communication and stress management skills focused on fostering resiliency and wellness in oncology residents by a local Radiation Oncologist, covered topics such as mindfulness, healthy habits and reframing stress with an interactive focus on experiential learning and group discussion;	Online virtual Facebook (FB) community for female physicians practicing in hematology/ oncology, founded in 2015, dynamic evolved to include advice on complex deidentified cases, real-time updates from H/O conferences, designated
Author, Year (Country)	Clemons et al, 2019 ²⁶ (Canada)	Dahn et al, 2019 ²⁷ (Canada)	Graff et al, 2018³³ (United States)

Author Vear	Intervention	Final Diagnosis / After intervention Pesults hefore and	sacisulado	limitations
		after intervention	COLICIASIONS	בוווונפנוסווס
	13 weekly meetings organized by psychologist and psychologist and psychologist-art therapist: 5 used psychodrama techniques to promote communicative exchanges; 4 'play-therapy' and stimulate a sense of comfort by non-verbal communication based on play; 3 to Ericksonian relaxation techniques; and 1 to observed and discussed a video with techniques to support children during painful procedures)	Baseline MBI mean±SD: - EE: Group A: NR Group B: 15.85±6.37 - RPA: Group A: NR Group B: 60.35±11.07 - Dp: Group A: NR Group B: 3.80±4.20 After 4mo MBI mean±SD: - EE: Group A: NR Group B: 67.40±9.10 - Dp: Group A: NR Group B: 2.25±2.63	Techniques using AT such as psychodrama and relaxation were effective for the operators who were most at risk of burnout	- Intervention was composed of different techniques
Kesselheim et al, 2020 ³⁹ (United States)	A novel, 4-module, case-based curriculum entitled "Humanism and Professionalism for Pediatric Hematology-Oncology Fellows", which aims to foster pediatric hematology-oncology fellows' reflection on the feelings, challenges, and conflicts arising in the care of children and families affected by cancer or blood disorders	Baseilne PHOSAH - intervention: 8.2 (3.3); usual training: 7.4 (4.2) MBI mean (SD): - EE: intervention: 2.3 (1.0); usual training: 2.4 (1.2) - RPA: intervention 4.8 (0.7); usual training: 4.7 (0.8) - Dp: intervention: 1.1 (0.9) - usual training: 4.4 (0.4) - Empowerment at Work Scale - intervention: 4.9 (0.6) - usual training: 4.5 (0.8) - Follow up 1y After PHOSAH: - intervention: 9.0 (3.9); - usual training: 8.0 (4.0) MBI mean (SD): - EE: intervention 2.2 (1.0); usual training = 4.8 (0.8) - Dp: intervention = 1.1 (0.8); usual training = : 1.2 (0.9) PPOS: - intervention: 4.3 (0.4); - usual training: 4.4 (0.4) Empowerment at Work Scale: - intervention: 5.0 (0.7); - usual training: 4.7 (0.6)	Exposure to the curriculum did not alter fellows' self-assessed humanism and professionalism skills. However, fellows expressed significantly higher levels of satisfaction in their humanism training, indicating the curriculum potential for positive impact on their perceived learning environment	The study: did not explicitly collect information about feasibility; and, is vulnerable to selection bias, as program directors who chose to have their programs participate may already prioritize humanism training in their fellowships; - Data on the routine strategies for teaching humanism that were utilized by usual training sites were collected retrospectively and could be vulnerable to recall bias

	s		ite over ive (i.e., outcome icluded; he EG	size; rres; rres; ve bias group ysicians o learn w skills inques); ne bias time in e facing inditions ed an in the)
	Limitations	Z Z	- High attrition rate over time; - No objective (i.e., non-self-report) outcome measures were included; Interaction of the EG (training) and CG	- Small sample size; - Short follow-up; self- report measures; - Potential positive bias within the study group (participating physicians were motivated to learn and practice new skills and coping techniques); - Potential outcome bias (simply spending time in a group of people facing similar working conditions may have played an important role in the outcomes)
	Conclusions	The intervention was effective	The relatively brief, teambased intervention program, not only influenced the stress component of burnout (EE) but also was a motivational component (Dp).	A significant effect indicated that lower scores for perceived stress after intervenção and there was decrease in emotional exhaustion and and some emotion regulation skills with small to medium effect sizes
	Final Diagnosis / After intervention Results before and after intervention	Follow up 2y Reduction of burnout risk from intermediate in 60% of the oncologists to 10%,.None of the oncologists had developed burnout syndrome	MBI mean±SD: - EE: IG: 1.54±0.89 vs CG: 1.46±0.80 - RPA: NR - Dp: IG: 0.96±0.70 vs CG: 0.86±0.58 Follow pu 6 m0 After MBI mean±SD: - EE: IG: 1.53 0.92 vs CG: 1.65±1.00 - RPA: NR - Dp: IG: 0.98±0.65 vs CG: 0.93±0.62	Baseine Perceived stress (mean±SD): - IG: 3.25±0.69 - CG: 3.20±0.66 MBI-EE (mean±SD): - IG: 4.09±0.59 - CG: 4.19±0.60 Follow up 36 w after Perceived stress (mean±SD): - IG: 2.83±0.72 - CG: 3.29±0.64 MBI-EE (mean±SD): - IG: 3.71±0.68 - CG: 4.18±0.61 Preceived stress IG at T1 (F = 29.21, P < .001), T2 (F = 24.7, P < .001), and T3 (F = 8.76, P < .01) Decrease emotional exhaustion (MBI-EE) and some emotion regulation skills (P < .01) (d=0.3-0.7)
	Intervention	Anti-stress program to medical oncologists consisting of oncologists' 1d-meeting outside outside the workplace every 3mo, in which there are integrative activities, teamwork, and workshops of stress management	Team-based burnout intervention called "Take Care!" program with a training plan of 6mo sessions of 3h each, which were supervised by both team counselors	Intervention based on Lazarus transactional model of stress, including 2 strategies of coping with stressors: problem- and emotion-oriented coping; psychosocial competency training focus on current working situations and problems of junior oncologists, coping strategies, resilience, and self-efficacy training as well as developing a support system among colleagues (12w sessions of 1.5h. All training sessions involved theoretical input, watching videos, oral group discussions, experiential exercises, and home assignments) combined with cognitive behavioral and
continue	Author, Year (Country)	Landaverde et al, 2018 ²⁸ (Costa Rica)	Le Blanc et al, 2007³⁵ (the Netherlands)	Mache et al, 2017 ²⁹ (Germany)



continue				
Author Year	Intervention	Final Diagnosis / After intervention Results before and	Conclusions	Limitations
(Country)		after intervention		
Medisauskaite	Intervention of 4 modules:	Baseline	From baseline to time-2 there	- Short follow-up
and Kamau,	 Module 1 taught doctors 	Mean (SD)	were significant reductions	
201936	about stress	MBI:	in burnout (EE), burnout (Dp)	
(United	 Module 2 taught doctors 	- EE: EG: 3.26±1.41 CG: 3.2±1.4	and anxiety among doctors	
Kingdom)	about burnout	- RPA: EG: 4.42±0.83 CG: 4.41±0.82	who completed all modules	
	 Module 3 taught doctors 	- Dp: EG: 1.98±1.49 CG: 1.68±1.29	about the psychology of	
	about coping with patient	Anxiety: EG: 0.96±0.81 CG: 0.88±0.74	distress.	
	death	Psychiatric morbidity: EG: 2.14±0.57 CG: 2.17±0.61		
	 Module 4 taught doctors 	Grief: EG: 1.6±0.6 CG: 1.74±0.66		
	about methods of managing	Insomnia: EG: 1±0.84 CG: 1.18±0.84		
	distress.	Physical symptoms: EG: 1.75±0.51 CG: 1.84±0.56		
		Alcohol use habits: EG: 7.33±2.26 CG: 6.71±1.92		
		Binge-eating features: EG: 1.38±1.75 CG: 1.1±1.69		
		Drug use: EG: 0.71±0.87 CG: 0.78±0.78		
		Follow up 7 d		
		mean±SD		
		MBI:		
		- EE: EG: 2.98±1.44 CG: 3.04±1.42		
		- RPA: EG: 4.38±0.91 CG: 4.27±0.85		
		- Dp: EG: 1.68±1.41 CG: 1.72±1.35		
		Anxietv:EG: 0.73±0.72 CG: 0.81±0.74		
		Psychiatric morbidity: EG: 2.16±0.57 CG: 2.21±0.64		
		Grief: EG: 1.51±0.57 CG: 1.64±0.62		
		Insomnia: EG: 1.02±0.96 CG: 1.11±0.87		
		Physical symptoms: EG: 1.69±0.61 CG: 1.85±0.65		
		Alcohol use habits: EG: 7.39±2.38 CG: 6.99±1.95		
		Binge-eating features: EG: 1.54±1.86 CG: 1.1±1.74		
		Drug use: EG: 0.53±0.69 CG:0.69±0.81	,	
Moody	Mindfulness-based course	Baseline	Nearly 100% of participants	- Small sample size; -
et al, 2013³¹	(MBC) participants received	MBI mean±SD:	met criteria for high levels of	Overrepresentation of
(United States	8 weeks of didactic and	- EE: IG: 27.2 CG: 26.2	burnout in the categories of	women in the sample;
and Israel)	experiential mindfulness	- RPA: IG: 16.0 CG: 15.4	RPA and (Dp). in both the CG	- Lack of any intervention
	education via a structured,	- Dp: IG: 19.6 CG: 18.2		in the CG; lack of blinding
	skills-training course delivered	Follow up 8w		
	in a group setting at their		category of EE, greater than	
	hospital. The course included 1	MBI mean±SD:	95% of participants in both	
	initial 6-hour session; 6 weekly	- EE: IG: 26.9 CG: 24.2	groups, at both time points,	
	1-hour follow-up sessions; and	7	showed moderate or high	
	a final 3-hour wrap-up session	- Dp: IG: 19.3 CG: 18.7	levels of burnout	
	(15 hours total class time).			

	Limitations	N N	- Small sample size; Important effects of the intervention or of the first oncology fellowship year may have been missed; Two-group comparison analyses were used instead of paired comparisons; Fellows in this program may not have been representative of Hematology-Oncology fellows across the United States; Attitudes questionnaire we used has not been validated previously, and individual topic domains varied in their reliability
		a novel ial capital oblem of sy trainees social altruistic faculty- orically- I cohorts	
	d Conclusions	The FitFirms are a novel system using social capital to reduce the problem of burnout in oncology trainees by engaging in social connectivity and altruistic service through facultymentored, historicallynamed divisional cohorts	In conclusion, hematology- oncology fellows' attitudes change over the course of the first fellowship year. Positive attitudes and development as caring physicians can be enhanced through the institution of a physician awareness group. The impact and effectiveness of the EG can be measured, and successful groups should improve the ability of physicians to communicate with their patients, and thus patient satisfaction
	Final Diagnosis / After intervention Results before and after intervention	Before Nine pre-intervention surveys were collected with 78% of trainees describing themselves as on the burnout spectrum of feeling either ineffective, overextended, disengaged, or burned out (22% engaged) Follow up 15 mo 10 post-intervention surveys were collected in which 60% of trainees described themselves on the burnout spectrum (40% engaged)	Base line Mean (range) - Full questionnaire summary score: 3.6 (3.5-3.7) - Stress in the work environment: 3.3 (3.2-3.5) - Comfort dealing with emotional patient/clinical situations: 3.5 (3.3-3.7) - Fellow's views of him/herself as a physician: 3.8 (3.7-3.9) - Discomfort with psychosocial issues: 3.7 (3.5-3.9 Follow up 1y after Mean (range) - Full questionnaire summary score: 3.7 (3.6-3.8) - Stress in the work environment: 3.4 (3.2-3.6) - Comfort dealing with emotional patient/clinical situations: 3.7 (3.6-3.9) - Fellow's views of him/herself as a physician: 4.1 (3.9-4.2)
	Intervention	Adapted Firm System called the FitFirms, which focused on social connectivity and altruistic service as means to combat burnout in oncology trainees The faculty and fellows interacted on an at-minimum quarterly basis in casual social events and/or community service-oriented events A didactic discussion series was created to explore concepts of resiliency, worklife balance, and the role of art in medicine—mentored by faculty across the spectrum of oncologic disciplines	Balint-like physician awareness group every 2 weeks for 1.5 to 2 hours for 6 months -
continue	Author, Year (Country)	Pathak, Eapen and Zell, 2019 ⁴⁰ (United States)	Sekeres et al, 2003 ⁴¹ (United States)



continue				
Author, Year	Intervention	Final Diagnosis / After intervention Results before and	Conclusions	Limitations
(Country)		after intervention		
Tjasink and	Art Therapy: mixture of	Baseline	MBI-HS pre- and post-	- Small sample; more
Soosaipillai,	different techniques such	MBI mean±SD:	intervention demonstrated	effective techniques
2018 ³⁷	as mindfulness, relaxation,	- EE: 30.79±8.31	statistically significant	learned through the
(England)	visualization, psychodrama	- RPA: 35.38±6.51	improvements in EE	first experience may
	and skills based supervision	- Dp: 7.93±5.05		have been used in the
	alongside art therapy; Six	Follow up 1 mo		second and third groups;
	weeks of structured art	AfterMBI mean±SD:		group was self-selecting;
	therapy sessions lasting	- EE: 23.5±7.61		the authors feel that a
	90–120 minutes each,	- RPA: 38.31±5.31		reluctance for some to
	structured and divided into	- Dp: 6.79±4.68		join the course; Data was
	three broad themes: Self-	improvements in EE (p=<0.001) and RPA (p=0.011)		not collected from those
	awareness and self-care;			who chose not to respond
	Collegial connection and			to the opportunity
	the organization; Reflecting			
	on death, bereavement and			
	finding meaning			

Cl=confidence interval; d=days; CG=Control Group; EG=Experimental Group; h=hour; min=minutes; mo=month; MBI=Maslach Burnout Inventory; EE=Emotional exhaustion; RPA=Reduced personal accomplishment; Dp=Depersonalization; NR=Not reported; RCT=randomized-controlled trial; SD=Standard Deviation; U=Undear; w=weeks; y=years; (*) data calculated by the authors.

Of the 17 residents that were participating, 2 decided not to continue after 2 meetings and therefore were excluded.
13 physicians completed the baseline scores, 11 completed Maslach/Oxford scores at the end of the study, and 8 the 1-month post-study assessment

Dropouts: at T2 the number of participants had dropped from 664 (experimental group: 260; control group: 404) to 376 (experimental group: 231; control group: 145), and at T3 it had dropped to 304 (experimental group: 208; control group: 96). ⁴ The study involved 227 doctors however the analysis considered in the present SR were Group 4 and control (91 doctors).

Additional information by emailing official authors.

The questionnaire derived from the Physician's Belief Scale; the American Academy on Physician and Patient evaluation; common objectives of Balint-like groups across the United States; barriers to physician recognition of psychosocial aspects of health care reports; and from surveys of previous hematology-oncology fellows to explore attitudes toward patients, colleagues, and psychosocial issues I have an experiment on the first session, and two candidates did not complete the post-intervention MBI-HSS survey.



Two studies used art therapy, [34,37] one study used a virtue list, [26] one used an online virtual community of women oncologists on Facebook, in which, [38] and another used a novel case-based curriculum. [39]

The Maslach burnout inventory (MBI) was used to measure changes before and after the intervention in fifteen studies. [24,33,37,39,40] Other questionnaires were used in some studies to analyze the interventions' effect on stress and burnout levels, [23,29,31,32,35,36,38] but only three of them also used MBI. [29,32,36] In two studies non-validated questionnaires were used to examine the participants' behavior and satisfaction after the intervention. [38,41]

Results of individual studies

The follow-up periods ranged from 7 days^[36] to 2 years,^[28] with eight studies with follow-ups longer than 1 year^[24,28-30,33,39,40] and ten shorter than 6 months.^[23,25-27,31,34-37,41] One cross-sectional study did not have follow-up, and evaluated participants' outcomes only after the intervention.^[38]

Art-therapy was evaluated in two studies, both with before and after design. [34,37] Italia et al. (2008) [34] organized weekly meetings that included psychodrama, games, relaxation techniques and videos. Tjasink and Soosaipillai (2018) [37] applied relaxation techniques, visualization, mindfulness meditation, psychodrama and skill-based supervision with art therapy. Through these techniques, authors addressed three broad themes: self-awareness and self-care; collegial connection and the organization; reflecting on death, bereavement and finding meaning alongside art therapy. [37]

A cross-sectional study published by Graff et al. (2018)^[38] impact of a closed community on Facebook in reducing professional burnout in women oncologists. In this group, they had the opportunity to discuss complex clinical cases, promote updates to clinical practice and magazine clubs, relocate transferred patients and disseminate research protocols. In addition, they were able to share ideas on the balance between life and work, and on the art of oncology.

Kesselheim et al. (2020)^[39] developed a curriculum with the objective to promote reflections on the feelings, challenges, and conflicts that arise in the care of children and families affected by cancer or blood disorders by pediatric hematology-oncology fellows.

Other approaches that worked to reduce physician stress and burnout were based on small groups, and their curriculum included the following strategies: a training program supervised by counselors, [27,33,35] coping with problems based on the Lazarus, [29] and educational approach on stress, burnout and methods to combat them. [36] Landaverde et al. (2018)[28] developed an anti-stress program that consisted of taking oncologists for a day out of the workplace for integration activities, teamwork and workshops to deal with stress, and Pathak et al. (2019)[40] used an adapted system to combat burnout in oncology trainees by focusing on social connectivity and altruistic service with a later didactic discussion.

Three studies included in this systematic review attempted to combat the burnout of the oncologist by promoting communication skills between doctors and patients. [23-25] These studies, however, have failed to demonstrate effectiveness.

Two studies evaluated Balint groups' impact on reducing burnout in residents and oncology fellows. [30,41] Intervention promoted the improvement of communication skills and contributed to the doctors' sense of self-realization, but it was not effective in preventing or reducing burnout.

In a randomized clinical trial conducted by Moody et al. (2013),^[31] despite mindfulness caused positive changes in the participants, both at work and at home, it did not reduce their burnout.

Interventions which had a significant effect in the reduction of stress and burnout were experience sharing between women doctors in virtual groups, [38] art therapy, [34,37] team monthly meetings outside the work environment, [28] training sessions supervised by counselors, [35] sessions with strategies for coping with stressors focused on both the problem and the solution, [29] and teaching doctors about stressors, burnout, how to deal with death and with stress. [36]

Interventions that did not reduce were simulated communication skills training, [23-25] Balint groups, [30,41] Franklin's new virtue model, [26] mindfulness, [31] adapted systems, [40] and the created humanism and professionalism curriculum. [39]

Risk of bias within and across studies

Nine of the 19 studies included fulfilled all the applicable questions regarding the methodological quality criteria, being classified with low-risk of bias: three RCT,^[35,36,39] the cross-sectional study,^[38] and five quasi-randomized studies,^[26,30,34,37,40] Six articles had a moderate RoB^[24,25,29,31,33,41] and four were classified as a high RoB^[23,27,28,32] score according to the specific checklist.

The main topics that introduced potential bias into the studies were the selection of the reported results and the existence of other interventions on the compared groups. Figures 2A-2D and Appendix 3 present the RoB graphs and detailed information regarding the RoB assessment, respectively.

Synthesis of results

All studies were based on interventions focused on the participants individually, or in group, and any intervention made a structural change within the work environment. [28] There has been no study that made a structural change within the work environment.

Confidence in cumulative evidence

Certainty in cumulative evidence was considered low and very low for randomized and observational studies, respectively. Further explanations regarding evidence appraisal are presented in Table 3.



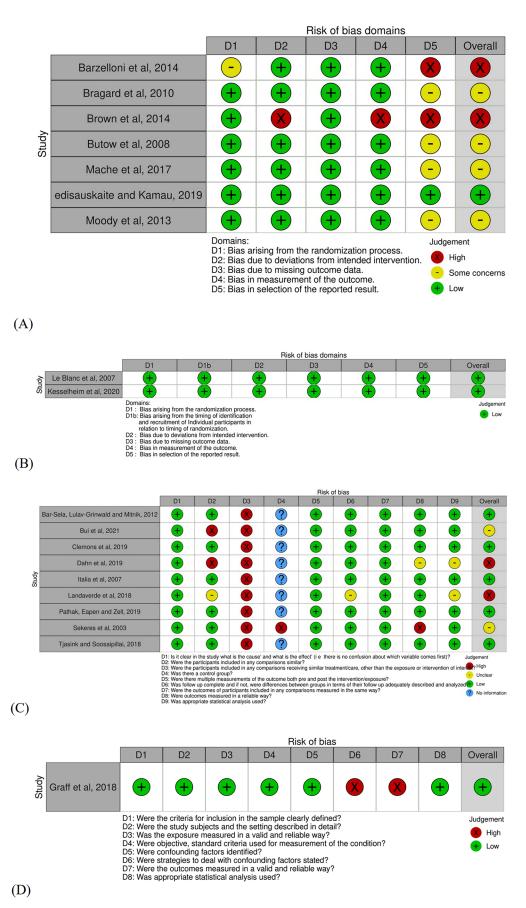


Figure 2. RoB summary author's judgments for each included study, assessed by the Cochrane risk of bias tool RoB 2.0 for randomized trials¹⁹ (A), the Cochrane risk of bias tool RoB 2.0 for cluster-randomized trials¹⁹ (B), the JBI Critical Appraisal Checklist for Quasi-experimental for non-randomized experimental studies¹⁷ (C), and the JBI Critical Appraisal Checklist for Analytical Cross Sectional Studies¹⁸ for descriptive studies (D). Author's judgements were graphically represented by "Traffic-light" plot (generated using the online tool *robvis* (Risk-Of-Bias VISualization) (National Institute for Health Research).²⁰



Table 3. The Grading of Recommendation Assessment, Development and Evaluation (GRADE) summary of findings table.

Tubic 5. 11	ic drading of the	Commendation	7.556551116116, DCV	ciopinent and i		ADL) Sulfillially Of II	rianigs table.
	Certainty assessment						
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Certainty
Impact o	mpact on Burnout symptoms (follow up: range 7 days to 1 years)						
6	randomised trials	serious ^a	serious ^b	not serious	not serious	none	⊕⊕⊜⊝ LOW
Impact o	n Burnout sym	ptoms (follow	up: range 1 mor	ths to 2 years)			
9	observational studies	very serious ^c	very serious ^b	not serious	not serious	none	⊕○○○ VERY LOW
Impact o	n Burnout sym	ptoms					
1	observational studies	not serious	not serious	not serious	serious ^d	none	⊕○○○ VERY LOW
Prevention	on on Burnout						
1	randomised trials	very serious ^e	not serious	not serious	not serious	none	⊕⊕⊜⊝ LOW
Stress lev	vel (follow up: r	ange 1 months	to 9 months)				
3	randomised trials	very serious ^f	very serious ^b	serious ^g	not serious	none	⊕○○○ VERY LOW

CI: Confidence interval

CI: Confidence interval Explanations:

a. Some included studies presented problems on domains 1 (randomization process), 4 (measurement of the outcome) and 5 (selection of the reported results). Overall risk of bias was considered Moderate.

b. Studies have methodological differences - Measurement method to burnout detection, coping strategy and / or follow-up period.

c. Included studies showed concerns in domain related to control group and it comparision treatment/care.

d. Intervention group with small size.

e. The study present concerns in the randomization process.

f. The studies showed some concerns in the domain related to selection of the reported results.

G. Stress is one of the predictor factors to burnout, however isolated is not conclusive.

DISCUSSION

The objective of this systematic review was to identify interventions which are effective to prevent or reduce the signs and symptoms of stress and burnout on oncology physicians compared to oncology physicians who did not participate in the interventions.

Our findings showed that eight studies had interventions which were effective.

Two of these studies[34,37] were their interventions based on art therapy. This type of intervention has been used in oncology and palliative care environments with the aim of helping to overcome grief, boost morale and reduce the burnout of the multidisciplinary team.[42] As both studies used several techniques,[34,37] despite the good results, there are doubts about which strategy or strategies in the interventions were the real agents of change.

Another intervention which reduced burnout was a virtual community of practices to sharing experiences among women oncologists' doctors.[38] This was an innovative work, without any previous reference in literature. However, the study was cross-sectional and evaluated the community members (female physician practicing hematology and oncology) through a voluntary anonymous 12-question online survey using a visual analog scale, that is not the gold standard way of measuring burnout before and after any intervention. Thus, the results found in this study may be led to a more subjective effect of stress than burnout itself.

Other interventions that reduced stress and burnout were based on small groups and were conducted at a time protected (paid) by the employer. It's important because both doctors and employers share the responsibility to promote the doctor's well-being.[12]

On the other hand, eleven studies had interventions which had no effect on stress and burnout.

Three of them used interventions to promote communication skills between doctors and patients.[23-25] Ineffective communication, in addition to negatively influencing the patient's well-being, interferes with the multidisciplinary team, which can cause increased stress and burnout, as well as less satisfaction with work.[43] These studies, however, have failed to demonstrate effectiveness.[23-25] Data are in line with literature. A systematic review published in the Cochrane Library and updated for the third time in 2018 shows that training communication skills are not effective in reducing the burnout of professionals working with cancer.[43]

Two of them had Balint groups[30,41] and showed no effect on stress and burnout. Balint groups were created in the 1950s by the psychoanalyst Michael Balint^[44] and aims at improving physician-patient relationship, by reflecting on the patterns of action of the patient and reaction of the physician (transference and counter-transference). Results must be interpreted with caution, due to the small number of participants in the studies and to the limited number of sessions. However, they are in agreement with the findings of other studies.[30,41]



A systematic review published in 2017^[45] with the aim of reviewing and evaluating evidence on psychosocial interventions to reduce occupational stress and burnout among physicians in general showed that none of the studies that used Balint groups reported significant effects.

Studies containing attempts to promote reflections on the feelings^[39] and focusing on social connectivity and altruistic service^[40] also failed in reducing burnout among oncologists.

Furthermore, one randomized clinical trial with the effects of mindfulness could not identify its effect on stress and burnout.[31] However, it caused positive changes in the participants, both at work and at home. Mindfulness has appeared in recent studies as a potential treatment for work-related burnout.[46,47] Defined as an awareness obtained when paying close attention to the moment, on purpose and without judgement, it originates from Buddhist values and can be learned.[46] Most programs currently offered are based on a program developed in the 1970s called mindfulness-based stress reduction (MBSR).[46] Some studies explore the relationship between mindfulness and burnout in healthcare professionals and suggest benefits, but with no significant impact on reducing burnout.[47]

Some of the interventions of the studies included in our systematic review, such as mindfulness and Balint groups, require a period of participations, learning or practice, [48] and perhaps the studies that evaluated these interventions did not have time enough to demonstrate effectiveness.

In regard to the interventions' predominant focus on individuals, similar findings have been found in other reviews.^[12]

West et al. (2016)^[12] evaluated the effectiveness of interventions in doctors with burnout in a systematic review and meta-analysis involving 15 randomized studies and 37 observational studies. Only three of these were institutional structural studies.

Petrie et al. (2019),^[48] in a meta-analysis of interventions that could reduce the symptoms of mental disorders and suicidal ideation among doctors was not able to include any controlled study at the organizational level. This was considered quite worrying for the authors, since an institutional study has the advantage of identifying risk factors present in the work environment, the potential to be more acceptable to the participants and to have greater preventive power.^[48]

Despite some important findings in the studies included in this systematic review,^[23-41] many limitations are present. Among them the small sample size, the limited number of sessions, the voluntary recruitment, the interaction between control and intervention groups and only one institution included in the study.

In addition, most studies had evidenced the low quality and the follow-up periods were quite different.

Also, the studies had methodological differences, both in terms of design and in the population, in which not only oncologists were included. In some studies, measures were also differently obtained, through distinctive self-assessment questionnaires.^[23-26,28]

Even with failures, all studies that have evaluated possible ways to reduce stress and burnout among oncologist physician are still valid, since the prevalence of burnout varies from 23% to 48% around the world. [36] Any way of trying to mitigate such public health problem must be recognized and encouraged.

Individual approaches are useful, but in order to improve physicians' conditions to get involved and really care for people, the healthcare organization must work to control or eliminate known causes of wear and tear and enhance their defense and support systems. [49] Medical societies, hospitals, governments have a great responsibility to improve working conditions, but they commonly have difficulties in allocating resources to bring about effective changes. [1]

It cannot be said with certainty that the interventions which did not presented impact on burnout in the included studies do not actually work. Likewise, data are also insufficient to show exactly the most effective interventions. Therefore, additional studies with interventions to prevent or decrease stress and burnout are still needed, including not only individual but also organizational and work environment changes.

CONCLUSION

In this systematic review, interventions which have effect in the reduction of stress and burnout were virtual oncologists' women community of practices to share experiences, art therapy, team monthly meetings outside the work environment, training sessions supervised by counselors, the teaching of stress coping strategies, and about stressors, burnout, and ways to identify and handle them. Interventions that did not reduce were communication skills training included Balint groups, Franklin's new virtue model, mindfulness, adapted systems, and the created humanism and professionalism curriculum for pediatric hematology-oncology.

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APPENDIX

Appendix 1. Database search strategy

Database	Search (on July 22 nd , 2021)
CINAHL	("Burnout" OR "Burn out" OR "burned out" OR ((stress* OR distress*) AND (work* OR
	"job"))) AND (Oncologist* OR "medical oncology" OR "oncology physician" OR "oncology
	physicians") AND ("psychological adaptation" OR "psychological adaptations" OR "Coping"
	OR "Behavior" OR resilienc* OR "happiness" OR "well being" OR "wellness" OR therapeutic*
	OR treatment* OR therap* OR "quality of life" OR "Life satisfaction" OR intervention*)
COCHRANE	("Burnout" OR "Burn out" OR "burned out" OR ((stress* OR distress*) AND (work* OR
	"job"))) AND (Oncologist* OR "medical oncology" OR "oncology physician" OR "oncology
	physicians") AND ("psychological adaptation" OR "psychological adaptations" OR "Coping"
	OR "Behavior" OR resilienc* OR "happiness" OR "well being" OR "wellness" OR therapeutic*
	OR treatment* OR therap* OR "quality of life" OR "Life satisfaction" OR intervention*)
EMBASE	EMBASE ("Burnout" OR "Burn out" OR "burned out" OR ((stress* OR distress*) AND (work*
	OR "job"))) AND (Oncologist* OR "medical oncology" OR "oncology physician" OR "oncology
	physicians") AND ("psychological adaptation" OR "psychological adaptations" OR "Coping"
	OR "Behavior" OR resilienc* OR "happiness" OR "well being" OR "wellness" OR therapeutic*
	OR treatment* OR therap* OR "quality of life" OR "Life satisfaction" OR intervention*)
LILACS	(tw:("Burnout" OR "Burn out" OR "burned out" OR ((stress* OR distress* OR estres*) AND (work*
	OR "job" OR trabalho OR trabajo)))) AND (tw:(oncologist* OR "medical oncology" OR "oncology
	physician" OR "oncology physicians" OR oncolog* OR cancerolog* OR radioterapeuta*)) AND
	(tw:("psychological adaptation" OR "Psychological Adaptations" OR "psychological adaptations"
	OR "Coping" OR "Behavior" OR resilienc* OR "happiness" OR "well being" OR "wellness" OR therapeutic* OR treatment* OR therap* OR "quality of life" OR "Life satisfaction" OR
	intervention* OR "adaptação" OR "comportamento adaptativo" OR "felicidade" OR "alegria" OR
	"equilibrio" OR "bem-estar" OR "terapia" OR "Terapeutica" OR tratamento* OR "qualidade de
	vida" OR prevenc* OR intervenc* OR "adaptacion" OR "felicita" OR "bienestar" OR tratamiento*
	OR "calidad de vida")) AND (instance: "regional") AND (db:("LILACS") AND type:("article"))
PSYCINFO	("Burnout" OR "Burn out" OR "burned out" OR ((stress* OR distress*) AND (work* OR "job")))
	AND (Oncologist* OR "medical oncology" OR "oncology physician" OR "oncology physicians")
	AND ("psychological adaptation" OR "psychological adaptations" OR "Coping" OR "Behavior"
	OR resilienc* OR "happiness" OR "well being" OR "wellness" OR therapeutic* OR treatment*
	OR therap* OR "quality of life" OR "Life satisfaction" OR intervention*)
PUBMED	("Burnout, Professional"[Mesh] OR "Burnout, Psychological"[Mesh] OR "Burn out" OR
	"Burnout"[Title/Abstract] OR "burned out" OR (("Stress, Physiological"[Mesh:NoExp]
	OR Stress* OR distress*) AND (work* OR "job"))) AND ("Oncologists" [Mesh] OR
	"Oncologists"[Title/Abstract] OR "Oncologist"[Title/Abstract] OR "Oncologists"[Title/Abstract]
	OR "medical oncology"[Title/Abstract] OR "oncology physician"[Title/Abstract] OR "oncology
	physicians"[Title/Abstract])AND ("adaptation, psychological"[MeSH Terms] OR "psychological
	adaptation" OR "psychological adaptations" OR "Coping" OR "Behavior" OR "resilience" OR
	"Resiliences" OR "happiness" [MeSH Terms] OR "happiness" OR "well being" OR "wellness" OR "therapeutics" [MeSH Terms] OR "therapeutics" OR "treatment" OR
	"treatments" OR "therapy" OR "therapies" OR "quality of life"[MeSH Terms] OR "quality of
	life" OR "Life satisfaction" OR "intervention" OR "interventions")
SCOPUS	TITLE-ABS-KEY(("Burnout" OR "Burn out" OR "burned out" OR ((stress* OR distress*) AND
3001 03	(work* OR "job"))) AND (Oncologist* OR "medical oncology" OR "oncology physician" OR
	"oncology physicians") AND ("psychological adaptation" OR "Psychological Adaptations"
	OR "Coping" OR "Behavior" OR resilienc* OR "happiness" OR "well being" OR "wellness"
	OR therapeutic* OR treatment* OR therap* OR "quality of life" OR "Life satisfaction" OR
	intervention*)) AND (LIMIT-TO (DOCTYPE,"ar"))
WEB OF	("Burnout" OR "Burn out" OR "burned out" OR ((stress* OR distress*) AND (work* OR "job")))
SCIENCE	AND (Oncologist* OR "medical oncology" OR "oncology physician" OR "oncology physicians")
	AND ("psychological adaptation" OR "psychological adaptations" OR "Coping" OR "Behavior"
	OR resilienc* OR "happiness" OR "well being" OR "wellness" OR therapeutic* OR treatment*
	OR therap* OR "quality of life" OR "Life satisfaction" OR intervention*)
	("Burnout" OR "Burn out" OR "burned out") AND (Oncologist) AND ("psychological
GOOGLE	
GOOGLE SCHOLAR	adaptation" OR "Coping" OR "Behavior" OR resilienc* OR "happiness" OR "well being" OR "wellness" OR "quality of life" OR "Life satisfaction")



...continue

Database	Search (on July 22 nd , 2021)
OPENGREY	("Burnout" OR "Burn out" OR "burned out" OR ((stress* OR distress*) AND (work* OR "job"))) AND (Oncologist* OR "medical oncology" OR "oncology physician" OR "oncology physicians") AND ("psychological adaptation" OR "psychological adaptations" OR "Coping" OR "Behavior" OR resilienc* OR "happiness" OR "well being" OR "wellness" OR therapeutic* OR treatment* OR therap* OR "quality of life" OR "Life satisfaction" OR intervention*)
PROQUEST	noft(("Burnout" OR "Burn out" OR "burned out" OR ((stress* OR distress*) AND (work* OR "job"))) AND (Oncologist* OR "medical oncology" OR "oncology physician" OR "oncology physicians") AND ("psychological adaptation" OR "psychological adaptations" OR "Coping" OR "Behavior" OR resilienc* OR "happiness" OR "well being" OR "wellness" OR therapeutic* OR treatment* OR therap* OR "quality of life" OR "Life satisfaction" OR intervention*))

Appendix 2. Articles excluded and the reasons for exclusion (n=178)

Reference	Author	Reasons for Exclusion [*]
1.	Allegra et al, 2005	2
2.	Alorabi et al, 2015	2
3.	Ansmann et al, 2013	2
4.	Armstrong et al, 2004	5
5.	Asai et al, 2007	2
6.	Back et al, 2017	5
7.	Balbay, 2011	2
8.	Balch, 2010	2
9.	Balch, 2007	5
10.	Balch, 2009	5
11.	Balch et al, 2011	2
12.	Banerjee et al, 2017	2
13.	Barberio et al, 2015	2
14.	Beas et al, 2016	5
15.	Berman et al, 2007	2
16.	Bittner et al, 2011	1
17.	Blanchard et al, 2009	5
18.	Blanchard et al, 2010	2
19.	Borate, 2017	5
20.	Bragard et al, 2012	2
21.	Bragard et al, 2010	2
22.	Bressi et al, 2008	2
23.	Burki, 2018	5
24.	Camps et al, 2009	2
25.	Cano, 2016	2
26.	Caruso et al, 2012	2
27.	Carvalho et al, 2014	2
28.	Catt et al, 2005	2
29.	Chatwal et al, 2018	6
30.	Ciammella et al, 2011	2
31.	Cotton, 2018	5
32.	Creagan, 1993	5
33.	Crowe, 2016	5



Reference	Author	Reasons for Exclusion
34.	Cubero et al, 2013	2
35.	Cumbe et al, 2017	2
36.	Dahn et al, 2019	2
37.	Daruvala et al, 2019	2
38.	De Rezende et al, 2011	5
39.	Dix et al, 2012	5
40.	Dougherty et al, 2009	2
41.	Dubois et al, 2020	6
42.	Eelen et al, 2014	2
43.	Essaadi et al, 2013	2
44.	Fang et al, 2010	2
45.	Fiore et al, 2016	6
46.	Fitzgerald et al, 2012	2
47.	Flores et al, 2014	2
48.	Ftanou, 2017	5
49.	Fujimori et al, 2015	1
50.	Fusco et al, 2019	2
51.	Giansante et al, 2012	2
52.	Girgis et al, 2009	2
53.	Glasberg et al, 2007	2
54.	Granek et al, 2016 (A)	2
55.	Granek et al, 2017	2
56.	Granek et al, 2016 (B)	2
57.	Granek et al, 2015	2
58.	Granek et al, 2017	2
59.	Granek et al, 2016 (C)	2
60.	Granek et al, 2013	2
61.	Grant, 2018	6
62.	Grootenhuis et al, 1996	2
63.	Gross et al, 2014	2
64.	Grunfeld et al, 2000	2
65.	Guadagna et al, 2012	6
66.	Guest et al, 2011 (A)	2
67.	Guest et al, 2011 (B)	2
68.	Guveli et al, 2015	2
69.	Haley et al, 2004	2
70.	Hamilton, 2019	2
71.	Hegedus, 2012	1
72.	Henrik et al, 2021	2
73.	Hipp et al, 2015	2
74.	Hlubocky et al,	2
75.	Hlubocky et al, 2017	5
76.	Hlubocky et al,	2
77.	Holliday et al, 2017	2



Reference	Author	Reasons for Exclusion*
78.	Hudson et al, 2018	2
79.	Hunter, 2019	6
80.	Huynh et al, 2019	2
81.	lsc-En et al, 2011	2
82.	Jackson et al, 2008	2
83.	Jasperse et al, 2012	2
84.	Jasperse et al, 2014	2
85.	Joaquim et al, 2018	2
86.	Jørgensen et al, 2009	2
87.	Joubert et al, 2013	1
88.	Jutagir et al, 2017	1
89.	Karnyski et al, 2017	5
90.	Kash et al, 2000	2
91.	Kattlove et al, 1992	5
92.	Kavalieratos et al, 2017	1
93.	Kaymak et al, 2010	2
94.	Kearney et al, 2009	5
95.	Kiguchi et al, 2018	5
96.	Kinderman and Gross, 2014	3
97.	Kleiner and Wallace, 2017	2
98.	Knight et al, 2014	2
99.	Koh et al, 2015	1
100.	Koo et al, 2013	2
101.	Koocher, 1980	5
102.	Korones, 2010	5
103.	Kracen, 2011	2
104.	Kuerer et al, 2007	2
105.	Laurent et al, 2015	2
106.	Leones et al, 2020	6
107.	Liakopoulou et al, 2008	2
108.	Lievrouw et al, 2016	2
109.	López-Castillo et al, 1999	2
110.	Lyckholm, 2001	5
111.	Lyckholm, 2007	5
112.	Mahendran et al, 2014	1
113.	Mampuya et al, 2016	2
114.	Mampuya et al, 2017	2
115.	Manochakian, 2014	5
116.	Martins et al, 2016	2
117.	McFarland et al, 2017	2
118.	McLean et al, 2011	2
119.	Mehlis et al, 2018	2
120.	Mougalian et al, 2013	2
		<u>~</u>



Reference	Author	Reasons for Exclusion
122.	Mukherjee et al, 2014	2
123.	Murali et al, 2019	5
124.	Muriel et al, 2009	1
125.	Na, 2019	5
126.	Nabhan, 2009	5
127.	Nissim et al, 2019	2
128.	Nowakowski et al, 2016	2
129.	O'Byrne et al, 1997	2
130.	Paiva et al, 2018	2
131.	Paula Vega et al, 2017	2
132.	Penson et al, 2000	5
133.	Penson et al, 2005	5
134.	Poulsen et al, 2018	2
135.	Poulsen et al, 2011	2
136.	Poulsen et al, 2012	2
137.	Pye et al, 2013	2
138.	Ramey et al, 2017	2
139.	Ramirez et al, 1995	2
140.	Raphael et al, 2019	2
141.	Rath et al, 2014	2
142.	Rath et al, 2015	2
143.	Ratti et al, 2019	2
144.	Rohan et al, 2009	2
145.	Romeo et al, 2016 (A)	2
146.	Romeo et al, 2016 (B)	2
147.	Rosenstein, 2019	2
148.	Roth et al, 2011	2
149.	Royce et al, 2019	5
150.	Russo et al, 2014	2
151.	Sargsyan et al, 2017	2
152.	Sarra and Feuz, 2019	2
153.	Schirmers, 2016	5
154.	Schraub et al, 2004	2
155.	Shanafelt et al, 2006	5
156.	Shanafelt, 2005	5
157.	Shanafelt et al, 2014	2
158.	Shanafelt et al, 2014 (A)	2
159.	Shanafelt et al, 2005	2
160.	Shanafelt et al, 2014 (B)	2
161.	Shanafelt et al, 2014 (C)	2
162.	Shayne et al, 2012	
163.	Shereck et al, 2014	3
164.	Sheth et al, 2019	2
	5car cay 2013	_



continue		
166.	Shinan-Altman et al, 2018	1
167.	Sundquist, 2009	6
168.	Swetz et al, 2009	1
169.	Tamura et al, 2020	2
170.	Thompson et al, 2018	5
171.	Torres et al, 2013	2
172.	Tucunduva et al, 2006	2
173.	Turner et al,	2
174.	Vachon et al, 2016	1
175.	Vasylyeva et al, 2011	2
176.	Vetter et al, 2018	2
177.	Vici et al, 2021	2
178.	Wan, 2008	5

^{*}Legend: 1) involved only non-medical cancer professionals or medical students; 2) did not involve interventions to prevent or handle with stress and burnout; 3) had duplicated data from another included study or insufficient data; 4) were conducted in animals; 5) were reviews, letters, books, case report, case series with less than 10 participants, opinion article, technique articles and guidelines; 6) did not have their complete text available online/published and if the texts were not accessible after three contact attempts in a 15-day period by electronic mail to corresponding authors.

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Appendix 3. Risk of bias assessed by Cochrane risk of bias tool and Joanna Briggs Institute (JBI) critical appraisal tools. Risk of bias was categorized as High when the study reaches up to 49% score "yes", Moderate when the study reached 50% to 69% score "yes", and Low when the study reached more than 70% score "yes".

Cochrane Risk of Bias tool for Randomized Trials (RoB 2.0)

Study	Bias	Signalling question	Coments	Authors' judgement
	Bias arising from the randomization 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions?	1.1. Was the allocation sequence random?		PY
		Study: Abstract - Few information	NI	
	process	1.3. Were there baseline imbalances that suggest a problem with the randomization process?		N
		Domain-level judgement	Some co	ncerns
		2.1. Were participants aware of their assigned intervention during the trial?	Intervention: 1 day of classroom	Y
		2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial?	training Control: No intervention	Y
Barzelloni et al, 2014	Bias due to deviations from intended	2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context?		N
ni et a	interventions (effect of	tions 2.4. If Y/PY to 2.3 : Were these deviations likely to have		NA
rzelloı	assignment to intervention)	2.5. If Y/PY/NI to 2.4: Were these deviations from intended intervention balanced between groups?		NA
Bal		2.6. Was an appropriate analysis used to estimate the effect of assignment to intervention?		PY
		2.7. If N/PN/NI to 2.6 : Was there potential for a substantial impact (on the result) of the failure to analyse participants in the group to which they were randomized?		NA
		Domain-level judgement	Low	risk
		3.1. Were data for this outcome available for all, or nearly all, participants randomized?		NI
	Bias due	3.2. If N/PN/NI to 3.1 : Is there evidence that the result was not biased by missing outcome data?		N
	to missing outcome data	3.3. If N/PN to 3.2 : Could missingness in the outcome depend on its true value?		PN
		3.4. If Y/PY/NI to 3.3 : Is it likely that missingness in the outcome depended on its true value?		PN
		Domain-level judgement	Low	risk



	Bias	Signalling question	Coments	Authors' judgement
		4.1. Was the method of measuring the outcome inappropriate?		N
		4.2. Could measurement or ascertainment of the		N
		outcome have differed between intervention groups?		
		4.3. If N/PN/NI to 4.1 and 4.2: Were outcome assessors		NA
	measurement	aware of the intervention received by study participants?		
	of the outcome	4.4. If Y/PY/NI to 4.3: Could assessment of the outcome have		NA
	outcome	been influenced by knowledge of intervention received? 4.5. If Y/PY/NI to 4.4: Is it likely that assessment of the outcome		
		was influenced by knowledge of intervention received?		NA
		Domain-level judgement	Low	risk
		, ,	Study:	T S R
		5.1. Were the data that produced this result analysed in	Abstract	
4	accordance with a pre-specified analysis plan that was finalized	- Few	NI	
20.		before unblinded outcome data were available for analysis?	information	
<u>a</u> ,			- Study:	
et			Abstract	
Ē			- Few	
Barzelloni et al, 2014			information	
rze			- Evaluations	
Ва	Bias in	5.2. Is the numerical result being assessed likely to have	were	
	selection of the reported	been selected, on the basis of the results, from multiple eligible outcome measurements (e.g. scales, definitions,	performed at T0 and	PY
	result	time points) within the outcome domain?	quarterly	
	result	time points) within the outcome domain.	(T1, T2, T3, T4)	
		butonlyresults		
		comparing T0		
			and T4 were	
			showed.	
		5.3. Is the numerical result being assessed likely to have		
		been selected, on the basis of the results, from multiple		PY
		eligible analyses of the data?		
			111.1	
		Domain-level judgement	High	
		Domain-level judgement Overall bias	High High	risk
		Domain-level judgement Overall bias 1.1. Was the allocation sequence random?		
	Bias arising	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until		risk
	from the	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions?		risk PY
	from the randomization	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem		risk PY
	from the	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process?	High	risk PY Y N
	from the randomization	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement	High	risk PY Y N
	from the randomization	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention	Low Intervention:	risk PY Y N
10	from the randomization	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial?	Low Intervention: consolidation	risk PY Y N
2010	from the randomization	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants'	Low Intervention:	risk PY Y N
al, 2010	from the randomization	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial?	Low Intervention: consolidation workshops	risk PY Y N risk Y
et al, 2010	from the randomization process	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial?	Low Intervention: consolidation workshops Control: No	risk PY Y N risk Y
ard et al, 2010	from the randomization	Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial? 2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context?	Low Intervention: consolidation workshops Control: No	risk PY Y N risk Y
agard et al, 2010	from the randomization process	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial? 2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context? 2.4. If Y/PY to 2.3: Were these deviations likely to have	Low Intervention: consolidation workshops Control: No	risk PY N N risk Y N
Bragard et al, 2010	from the randomization process Bias due to deviations from intended interventions	Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial? 2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context? 2.4. If Y/PY to 2.3: Were these deviations likely to have affected the outcome?	Low Intervention: consolidation workshops Control: No	risk PY Y N risk Y
Bragard et al, 2010	from the randomization process Bias due to deviations from intended interventions (effect of	Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial? 2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context? 2.4. If Y/PY to 2.3: Were these deviations likely to have affected the outcome? 2.5. If Y/PY/NI to 2.4: Were these deviations from	Low Intervention: consolidation workshops Control: No	risk PY N N risk Y N
Bragard et al, 2010	from the randomization process Bias due to deviations from intended interventions (effect of assignment to	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial? 2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context? 2.4. If Y/PY to 2.3: Were these deviations likely to have affected the outcome? 2.5. If Y/PY/NI to 2.4: Were these deviations from intended intervention balanced between groups?	Low Intervention: consolidation workshops Control: No	risk PY N risk Y N N NA
Bragard et al, 2010	from the randomization process Bias due to deviations from intended interventions (effect of	Domain-level judgement Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial? 2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context? 2.4. If Y/PY to 2.3: Were these deviations likely to have affected the outcome? 2.5. If Y/PY/NI to 2.4: Were these deviations from intended intervention balanced between groups? 2.6. Was an appropriate analysis used to estimate the	Low Intervention: consolidation workshops Control: No	risk PY N risk Y N N NA
Bragard et al, 2010	from the randomization process Bias due to deviations from intended interventions (effect of assignment to	Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial? 2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context? 2.4. If Y/PY to 2.3: Were these deviations likely to have affected the outcome? 2.5. If Y/PY/NI to 2.4: Were these deviations from intended intervention balanced between groups? 2.6. Was an appropriate analysis used to estimate the effect of assignment to intervention?	Low Intervention: consolidation workshops Control: No	risk PY N N risk Y N N NA NA
Bragard et al, 2010	from the randomization process Bias due to deviations from intended interventions (effect of assignment to	Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial? 2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context? 2.4. If Y/PY to 2.3: Were these deviations likely to have affected the outcome? 2.5. If Y/PY/NI to 2.4: Were these deviations from intended intervention balanced between groups? 2.6. Was an appropriate analysis used to estimate the effect of assignment to intervential for a substantial	Low Intervention: consolidation workshops Control: No	risk PY Y N risk Y N NA NA NA
Bragard et al, 2010	from the randomization process Bias due to deviations from intended interventions (effect of assignment to	Overall bias 1.1. Was the allocation sequence random? 1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions? 1.3. Were there baseline imbalances that suggest a problem with the randomization process? Domain-level judgement 2.1. Were participants aware of their assigned intervention during the trial? 2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial? 2.3. If Y/PY/NI to 2.1 or 2.2: Were there deviations from the intended intervention that arose because of the trial context? 2.4. If Y/PY to 2.3: Were these deviations likely to have affected the outcome? 2.5. If Y/PY/NI to 2.4: Were these deviations from intended intervention balanced between groups? 2.6. Was an appropriate analysis used to estimate the effect of assignment to intervention?	Low Intervention: consolidation workshops Control: No	risk PY N N risk Y N N NA NA



...continue

Study	Bias	Signalling question	Coments	Authors' judgemen
	participants randomized? 3.2. If N/PN/NI to 3.1 : Is there evidence the was not biased by missing outcome	3.1. Were data for this outcome available for all, or nearly all, participants randomized?		PY
		3.2. If N/PN/NI to 3.1 : Is there evidence that the result was not biased by missing outcome data?		NA
		<u>'</u>		NA
		3.4. If Y/PY/NI to 3.3 : Is it likely that missingness in the outcome depended on its true value?		NA
		Domain-level judgement	Low	risk
		4.1. Was the method of measuring the outcome inappropriate?		N
0		4.2. Could measurement or ascertainment of the outcome have differed between intervention groups?		N
, 201(Bias in measurement	4.3. If N/PN/NI to 4.1 and 4.2 : Were outcome assessors aware of the intervention received by study participants?		NI
et al	of the outcome	4.4. If Y/PY/NI to 4.3 : Could assessment of the outcome have been influenced by knowledge of intervention received?		N
Bragard et al, 2010		4.5. If Y/PY/NI to 4.4: Is it likely that assessment of the outcome was influenced by knowledge of intervention received?		N
Bra		Domain-level judgement	Low	risk
_	Bias in selection of the reported result	5.1. Were the data that produced this result analysed in accordance with a pre-specified analysis plan that was finalized before unblinded outcome data were available for analysis?		NI
		5.2. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible outcome measurements (e.g. scales, definitions, time points) within the outcome domain?		N
		5.3. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible analyses of the data?		N
		Domain-level judgement	Some concerns	
		Overall bias	Some co	ncerns
		1.1. Was the allocation sequence random?	Study:	PY
	Bias arising from the randomization	1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions?	Abstract – Few information	NI
	process	1.3. Were there baseline imbalances that suggest a problem with the randomization process?		PN
		Domain-level judgement	Low	risk
, 2014		2.1. Were participants aware of their assigned intervention during the trial?	Intervention: face-to-face workshop Control: No intervention	Y
et al	Bias due to	2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial?		Y
Brown et al, 2014	deviations from intended	2.3. If Y/PY/NI to 2.1 or 2.2 : Were there deviations from the intended intervention that arose because of the trial context?		N
	interventions (effect of	2.4. If Y/PY to 2.3 : Were these deviations likely to have affected the outcome?		NA
	assignment to intervention)	2.5. If Y/PY/NI to 2.4 : Were these deviations from intended intervention balanced between groups?		NA
		2.6. Was an appropriate analysis used to estimate the effect of assignment to intervention?		NI
		2.7. If N/PN/NI to 2.6 : Was there potential for a substantial impact (on the result) of the failure to analyse participants in the group to which they were randomized?		PY
	_	Domain-level judgement	High	risk



.continue	Bias	Signalling question	Coments	Authors' judgement
	3.1. Were data for this outcome available for all, or nearly all, participants randomized? 3.2. If N/PN/NI to 3.1: Is there evidence that the result was not biased by missing outcome data? 3.3. If N/PN to 3.2: Could missingness in the outcome depend on its true value?	participants randomized?		NI
			N	
		depend on its true value?		PN
		3.4. If Y/PY/NI to 3.3 : Is it likely that missingness in the outcome depended on its true value?		PN
		Domain-level judgement	Low	risk
		4.1. Was the method of measuring the outcome inappropriate?	LOW	Y
		4.2. Could measurement or ascertainment of the		
		outcome have differed between intervention groups?		Y
4	Bias in	4.3. If N/PN/NI to 4.1 and 4.2 : Were outcome assessors		NA
20,	measurement	aware of the intervention received by study participants?		
Brown et al, 2014	of the outcome	4.4. If Y/PY/NI to 4.3 : Could assessment of the outcome have been influenced by knowledge of intervention received?		NA
ē		4.5. If Y/PY/NI to 4.4 : Is it likely that assessment of the outcome		
Š		was influenced by knowledge of intervention received?		NA
Bro		Domain-level judgement	High	risk
		5.1. Were the data that produced this result analysed in accordance with a pre-specified analysis plan that was finalized before unblinded outcome data were available for analysis?		NI
	Bias in selection of the reported result	Bias in selection of the reported 5.2. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible outcome measurements (e.g. scales, definitions)		Υ
		5.3. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible analyses of the data?		Y
		Domain-level judgement	High	
		Overall bias	High	
		1.1. Was the allocation sequence random?		Υ
	Bias arising from the	1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions?		PY
	randomization	1.3. Were there baseline imbalances that suggest a		N
	process	problem with the randomization process? Domain-level judgement	Low	viels
		2.1. Were participants aware of their assigned	Intervention:	IISK
m		intervention during the trial?	Communication	Υ
Butow et al, 2008		2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial?	skills training Control: No intervention	Y
w et	Bias due to deviations	2.3. If Y/PY/NI to 2.1 or 2.2 : Were there deviations from the intended intervention that arose because of the trial context?		N
Buto	from intended interventions	2.4. If Y/PY to 2.3 : Were these deviations likely to have affected the outcome?		NA
	(effect of assignment to	2.5. If Y/PY/NI to 2.4 : Were these deviations from intended intervention balanced between groups?		NA
	intervention)	2.6. Was an appropriate analysis used to estimate the effect of assignment to intervention?		Υ
		2.7. If N/PN/NI to 2.6 : Was there potential for a substantial impact (on the result) of the failure to analyse participants in the group to which they were randomized?		NA
		Domain-level judgement	Low	risk



Study	Bias	Signalling question	Coments	Authors' judgemen	
	Bias due to missing outcome data	3.1. Were data for this outcome available for all, or nearly all, participants randomized?		Y	
		3.2. If N/PN/NI to 3.1 : Is there evidence that the result was not biased by missing outcome data?		NA	
		3.3. If N/PN to 3.2 : Could missingness in the outcome depend on its true value?		NA	
		3.4. If Y/PY/NI to 3.3 : Is it likely that missingness in the outcome depended on its true value?		NA	
		Domain-level judgement	Low	risk	
		4.1. Was the method of measuring the outcome inappropriate?		N	
		4.2. Could measurement or ascertainment of the outcome have differed between intervention groups?		N	
Butow et al, 2008	Bias in measurement of the outcome	4.3. If N/PN/NI to 4.1 and 4.2 : Were outcome assessors aware of the intervention received by study participants?		Y	
		4.4. If Y/PY/NI to 4.3 : Could assessment of the outcome have been influenced by knowledge of intervention received?		N	
utow		4.5. If Y/PY/NI to 4.4 : Is it likely that assessment of the outcome was influenced by knowledge of intervention received?		NA	
8		Domain-level judgement	Low	risk	
	Bias in selection of the reported result	5.1. Were the data that produced this result analysed in accordance with a pre-specified analysis plan that was finalized before unblinded outcome data were available for analysis?		NI	
		5.2. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible outcome measurements (e.g. scales, definitions, time points) within the outcome domain?		N	
		5.3. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible analyses of the data?		N	
		Domain-level judgement Overall bias	Some concerns		
		Some co	ncerns		
	Bias arising from the randomization process	1.1. Was the allocation sequence random?		Y	
		1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions?		Y	
		1.3. Were there baseline imbalances that suggest a problem with the randomization process?		N	
		Domain-level judgement	Low	risk	
		2.1. Were participants aware of their assigned intervention during the trial?	Intervention: problem- and emotion-	Y	
Mache et al, 2017	Bias due to deviations from intended interventions (effect of	2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial?	oriented coping Control: No intervention	N	
		2.3. If Y/PY/NI to 2.1 or 2.2 : Were there deviations from the intended intervention that arose because of the trial context?		N	
		2.4. If Y/PY to 2.3 : Were these deviations likely to have affected the outcome?		NA	
	assignment to intervention)	2.5. If Y/PY/NI to 2.4 : Were these deviations from intended intervention balanced between groups?		NA	
		2.6. Was an appropriate analysis used to estimate the effect of assignment to intervention?		Y	
		2.7. If N/PN/NI to 2.6 : Was there potential for a substantial impact (on the result) of the failure to analyse participants in the group to which they were randomized?		NA	
		Domain-level judgement	Low	risk	



Study	Bias	Signalling question	Coments	Authors' judgement	
	Bias due to missing outcome data	3.1. Were data for this outcome available for all, or nearly all, participants randomized?		Υ	
		3.2. If N/PN/NI to 3.1 : Is there evidence that the result was not biased by missing outcome data?		NA	
		3.3. If N/PN to 3.2 : Could missingness in the outcome depend on its true value?		NA	
		3.4. If Y/PY/NI to 3.3 : Is it likely that missingness in the outcome depended on its true value?		NA	
		Domain-level judgement	Low		
		4.1. Was the method of measuring the outcome inappropriate?		N	
7		4.2. Could measurement or ascertainment of the outcome have differed between intervention groups?		N	
Mache et al, 2017	Bias in measurement	4.3. If N/PN/NI to 4.1 and 4.2 : Were outcome assessors aware of the intervention received by study participants?		Y	
e et a	of the outcome	4.4. If Y/PY/NI to 4.3 : Could assessment of the outcome have been influenced by knowledge of intervention received?		N	
lache		4.5. If Y/PY/NI to 4.4 : Is it likely that assessment of the outcome was influenced by knowledge of intervention received?		N	
2		Domain-level judgement	Low	risk	
	Bias in selection of the reported result	5.1. Were the data that produced this result analysed in accordance with a pre-specified analysis plan that was finalized before unblinded outcome data were available for analysis?		NI	
		5.2. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible outcome measurements (e.g. scales, definitions, time points) within the outcome domain?		PN	
		5.3. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible analyses of the data?		PN	
		Domain-level judgement	Some concerns		
		Overall bias	Some concerns		
	Bias arising from the randomization process	1.1. Was the allocation sequence random?		Y	
		1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions?		Υ	
		1.3. Were there baseline imbalances that suggest a problem with the randomization process?		N	
		Domain-level judgement	Low risk		
2019	Bias due to deviations from intended interventions (effect of assignment to intervention)	2.1. Were participants aware of their assigned intervention during the trial?	Intervention: learning	Y	
Medisauskaite and Kamau, 2019		2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial?	modules that presented doctors with information about stress Control: No intervention	N	
		2.3. If Y/PY/NI to 2.1 or 2.2 : Were there deviations from the intended intervention that arose because of the trial context?		N	
		2.4. If Y/PY to 2.3 : Were these deviations likely to have affected the outcome?		NA	
		2.5. If Y/PY/NI to 2.4 : Were these deviations from intended intervention balanced between groups?		NA	
		2.6. Was an appropriate analysis used to estimate the effect of assignment to intervention?		Υ	
		2.7. If N/PN/NI to 2.6 : Was there potential for a substantial impact (on the result) of the failure to analyse participants in the group to which they were randomized?		NA	
		Domain-level judgement	Low	risk	



Study	Bias	Signalling question	Coments	Authors' judgement	
	Bias due to missing outcome data	3.1. Were data for this outcome available for all, or nearly all, participants randomized?		Υ	
		3.2. If N/PN/NI to 3.1 : Is there evidence that the result was not biased by missing outcome data?		NA	
		3.3. If N/PN to 3.2 : Could missingness in the outcome depend on its true value?		NA	
		3.4. If Y/PY/NI to 3.3 : Is it likely that missingness in the outcome depended on its true value?		NA	
		Domain-level judgement	Low risk		
919		4.1. Was the method of measuring the outcome inappropriate?		N	
au, 2		4.2. Could measurement or ascertainment of the outcome have differed between intervention groups?		N	
Medisauskaite and Kamau, 2019	Bias in measurement	4.3. If N/PN/NI to 4.1 and 4.2 : Were outcome assessors aware of the intervention received by study participants?		N	
	of the outcome	4.4. If Y/PY/NI to 4.3 : Could assessment of the outcome have been influenced by knowledge of intervention received?		NA	
		4.5. If Y/PY/NI to 4.4 : Is it likely that assessment of the outcome was influenced by knowledge of intervention received?	NA		
an		Domain-level judgement	Low	risk	
Medis	Bias in selection of the reported result	5.1. Were the data that produced this result analysed in accordance with a pre-specified analysis plan that was finalized before unblinded outcome data were available for analysis?		Y	
		5.2. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible outcome measurements (e.g. scales, definitions, time points) within the outcome domain?		N	
		5.3. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible analyses of the data?		N	
		Domain-level judgement Overall bias	Low risk		
		Low	risk		
	Bias arising from the randomization process	1.1. Was the allocation sequence random?		Y	
		1.2. Was the allocation sequence concealed until participants were recruited and assigned to interventions?		PY	
		1.3. Were there baseline imbalances that suggest a problem with the randomization process?		N	
		Domain-level judgement	Low	risk	
Moody et al, 2013	Bias due to deviations from intended interventions (effect of assignment to intervention)	2.1. Were participants aware of their assigned intervention during the trial?	Intervention: Mindfulness-	Y	
		2.2. Were carers and trial personnel aware of participants' assigned intervention during the trial?	based course Control: No intervention	Y	
dy et		2.3. If Y/PY/NI to 2.1 or 2.2 : Were there deviations from the intended intervention that arose because of the trial context?		N	
Моос		2.4. If Y/PY to 2.3 : Were these deviations likely to have affected the outcome?		NA	
		2.5. If Y/PY/NI to 2.4 : Were these deviations from intended intervention balanced between groups?		NA	
		2.6. Was an appropriate analysis used to estimate the effect of assignment to intervention?		Y	
		2.7. If N/PN/NI to 2.6 : Was there potential for a substantial impact (on the result) of the failure to analyse participants in the group to which they were randomized?		NA	
		Domain-level judgement	Low	risk	



...continue

Study	Bias	Signalling question	Coments	Authors' judgement	
	Bias due to missing outcome data	3.1. Were data for this outcome available for all, or nearly all, participants randomized?		Y	
		3.2. If N/PN/NI to 3.1 : Is there evidence that the result was not biased by missing outcome data?		NA	
		3.3. If N/PN to 3.2 : Could missingness in the outcome depend on its true value?		NA	
		3.4. If Y/PY/NI to 3.3 : Is it likely that missingness in the outcome depended on its true value?		NA	
		Domain-level judgement	Low risk		
	Bias in measurement of the outcome	4.1. Was the method of measuring the outcome inappropriate?		N	
2013		4.2. Could measurement or ascertainment of the outcome have differed between intervention groups?		N	
		4.3. If N/PN/NI to 4.1 and 4.2 : Were outcome assessors aware of the intervention received by study participants?		N	
Moody et al, 2013		4.4. If Y/PY/NI to 4.3 : Could assessment of the outcome have been influenced by knowledge of intervention received?		N	
Mo		4.5. If Y/PY/NI to 4.4 : Is it likely that assessment of the outcome was influenced by knowledge of intervention received?		N	
		Domain-level judgement	Low risk		
	Bias in selection of the reported result	5.1. Were the data that produced this result analysed in accordance with a pre-specified analysis plan that was finalized before unblinded outcome data were available for analysis?		NI	
		5.2. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible outcome measurements (e.g. scales, definitions, time points) within the outcome domain?		PN	
		5.3. Is the numerical result being assessed likely to have been selected, on the basis of the results, from multiple eligible analyses of the data?		PN	
		Domain-level judgement	Some concerns		
		Some concerns			

Legend – Y=Yes, PY=Probably yes, PN= Probably no, N=No, NA=Not applicable, NI=No information.



JBI Critical Appraisal Checklist for Quasi-Experimental Studies (non-randomized experimental studies).

Question	1. Is it clear in the study what is the cause' and what is the effect' (i.e. there is no confusion about which variable comes first)?	2. Were the participants included in any comparisons similar?	3. Were the participants included in any comparisons receiving similar treatment/care, other than the exposure or intervention of interest?	4. Was there a control group?	5. Were there multiple measurements of the outcome both pre and post the intervention/exposure?	6. Was follow up complete and if not, were differences between groups in terms of their follow up adequately described and analyzed?	7. Were the outcomes of participants included in any comparisons measured in the same way?	8. Were outcomes measured in a reliable way?	9. Was appropriate statistical analysis used?	%yes/risk
Bar-Sela, LulavGrinwald and Mitnik, 2012	Υ	Υ	N	NA	Υ	Υ	Υ	Υ	Υ	77.78
Bui et al, 2021	Υ	N	N	NA	Υ	Υ	Υ	Υ	Υ	66.67
Clemons et al, 2019	Υ	Υ	N	NA	Υ	Υ	Υ	Υ	Υ	77.78
Dahn et al, 2019	Υ	N	N	NA	Υ	Υ	Υ	U	U	44.44
Italia et al, 2007	Υ	Υ	N	NA	Υ	Υ	Υ	Υ	Υ	77.78
Landaverde et al, 2018	Υ	U	N	NA	Υ	U	Υ	Υ	U	44.44
Pathak, Eapen and Zell, 2019	Υ	Υ	N	NA	Υ	Υ	Υ	Υ	Υ	77.78
Sekeres et al, 2003	Υ	Υ	N	N	Υ	Υ	Υ	N	Υ	66.67
Tjasink and Soosaipillai, 2018	Υ	Υ	N	NA	Υ	Υ	Y	Υ	Υ	77.78

Legend - Y=Yes, N=No, U=Unclear, NA=Not applicable.

JBI Critical Appraisal Checklist for Analytical Cross Sectional Studies

Legend - Y=Yes, N=No, U=Unclear, NA=Not applicable.