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European Society of Organ Transplantation (ESOT) Consensus Statement on Prehabilitation for Solid Organ Transplantation Candidates

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There is increasingly growing evidence and awareness that prehabilitation in waitlisted solid organ transplant candidates may benefit clinical transplant outcomes and improve the patient's overall health and quality of life. Lifestyle changes, consisting of physical training, dietary management, and psychosocial interventions, aim to optimize the patient's physical and mental health before undergoing surgery, so as to enhance their ability to overcome procedure-associated stress, reduce complications, and accelerate

Abbreviations: AGREE, Appraisal of Guidelines for Research & Evaluation; CET, Centre for Evidence in Transplantation; ESOT, European Society of Organ Transplantation; GRADE, Grading of Recommendations, Assessment, Development, and Evaluations; PICO, Population, Intervention, Comparator and Outcome; PRISMA, Preferred Reporting Items for Systematic reviews and Meta-Analyses; RCT, Randomized Controlled Trial.

post-operative recovery. Clinical data are promising but few, and evidence-based recommendations are scarce. To address the need for clinical guidelines, The European Society of Organ Transplantation (ESOT) convened a dedicated Working Group “Prehabilitation in Solid Organ Transplant Candidates,” comprising experts in physical exercise, nutrition and psychosocial interventions, to review the literature on prehabilitation in this population, and develop recommendations. These were discussed and voted upon during the Consensus Conference in Prague, 13–15 November 2022. A high degree of consensus existed amongst all stakeholders including transplant recipients and their representatives. Ten recommendations were formulated that are a balanced representation of current published evidence and real-world practice. The findings and recommendations of the Working Group on Prehabilitation for solid organ transplant candidates are presented in this article.

Keywords: prehabilitation, solid organ transplant candidates, exercise, nutrition, psychosocial interventions

INTRODUCTION

Patients who need a solid organ transplant often have a compromised overall condition due to end-stage organ failure, comorbidities, deconditioning, and treatment-related adverse effects such as dialysis in end-stage kidney disease (ESKD), left ventricular assist device (LVAD) in heart failure, and oxygen therapy in end-stage pulmonary disease (ESPD) [1–3]. Although considered a frail patient population with malnutrition, low physical fitness, fatigue, and often secondary psychological challenges, it is imperative for such patients to attain, and maintain, their optimal physical and mental wellbeing, as this will help them tolerate the waiting time and the stress of transplant surgery and expedite recovery after the transplant. The time spent on the transplant waitlist provides a window of opportunity to work towards enhancing the overall condition of such patients.

Prehabilitation refers to the optimization of patient’s overall physical and psychological condition before undergoing surgery, in order to enhance his/hers ability to overcome the stress associated with the procedure, to reduce the risk of complications and to accelerate post-operative recovery, with the ultimate goal to improve survival and quality of life [4]. The approach focuses on achieving lifestyle changes and should consist of physical training, dietary management, and psychological interventions [4]. By providing a multimodal program, the complex interaction between the physical and psychological health of a patient is addressed, which is important to maximize the outcomes of the interventions [5].

Prehabilitation has shown promising results in non-transplant patients undergoing major abdominal or orthopedic procedures [6–10], with reduced overall post-operative complications and morbidity, improved aerobic capacity, and improved functional recovery and shorter length of stay. The conclusions from two systematic reviews supported the feasibility and safety of such interventions in waitlisted solid organ transplant candidates [11, 12]. In addition, observed beneficial effects included improvements in cardiorespiratory function, exercise capacity, muscular strength, mental/physical composite scores and health-

related quality of life [11, 12]. There is a growing awareness and evidence that prehabilitation may not only benefit clinical transplant outcomes, but may also improve the transplant candidate’s overall health and quality of life, through adoption of a sustainable, healthy lifestyle. Despite this growing awareness and promising data, evidence-based recommendations for physical exercise, nutritional, or psychological prehabilitation interventions in candidates for solid organ transplants are not available. With regard to exercise interventions, recommendations on the role of exercise in solid organ transplantation were made by Janaudis-Ferreira et al in a position statement paper in 2019 [13].

The limited clinical guidance on how to implement prehabilitation for solid organ transplant candidates was presented as one of the priority themes at the first European Society of Organ Transplantation (ESOT) consensus conference in November 2022. Under the oversight of the ESOT guideline taskforce, and in keeping with the procedures recently established by the ESOT Consensus Platform for Organ Transplantation, leading experts presented in-depth literature evidence and proposed recommendations, which were publicly discussed and assessed by an independent jury, and consensus was formed [14]. Participants in the consensus process included not only transplant, prehabilitation and medical specialists, but also allied health professionals, patients and patient representatives.

This document presents the 2022 ESOT consensus findings and recommendations on implementing prehabilitation in the care for solid organ transplant candidates. These guidelines and recommendations undergo continual review and will be updated to reflect new evidence as it becomes available.

METHODS

The consensus development process was governed by the dedicated ESOT Guidelines Taskforce and co-organized by the ESOT sections European Liver and Intestine Transplant Association, European Kidney Transplant Association,

European Pancreas and Islet Transplant Association, European Cardio Thoracic Transplant Association, European Transplant Allied Healthcare Professionals, the ESOT Education Committee and Young Professionals in Transplantation.

The consensus development process followed the methodology stipulated by the ESOT Consensus Platform as recently published in detail [14]. In brief, the subsequent steps were as follows:

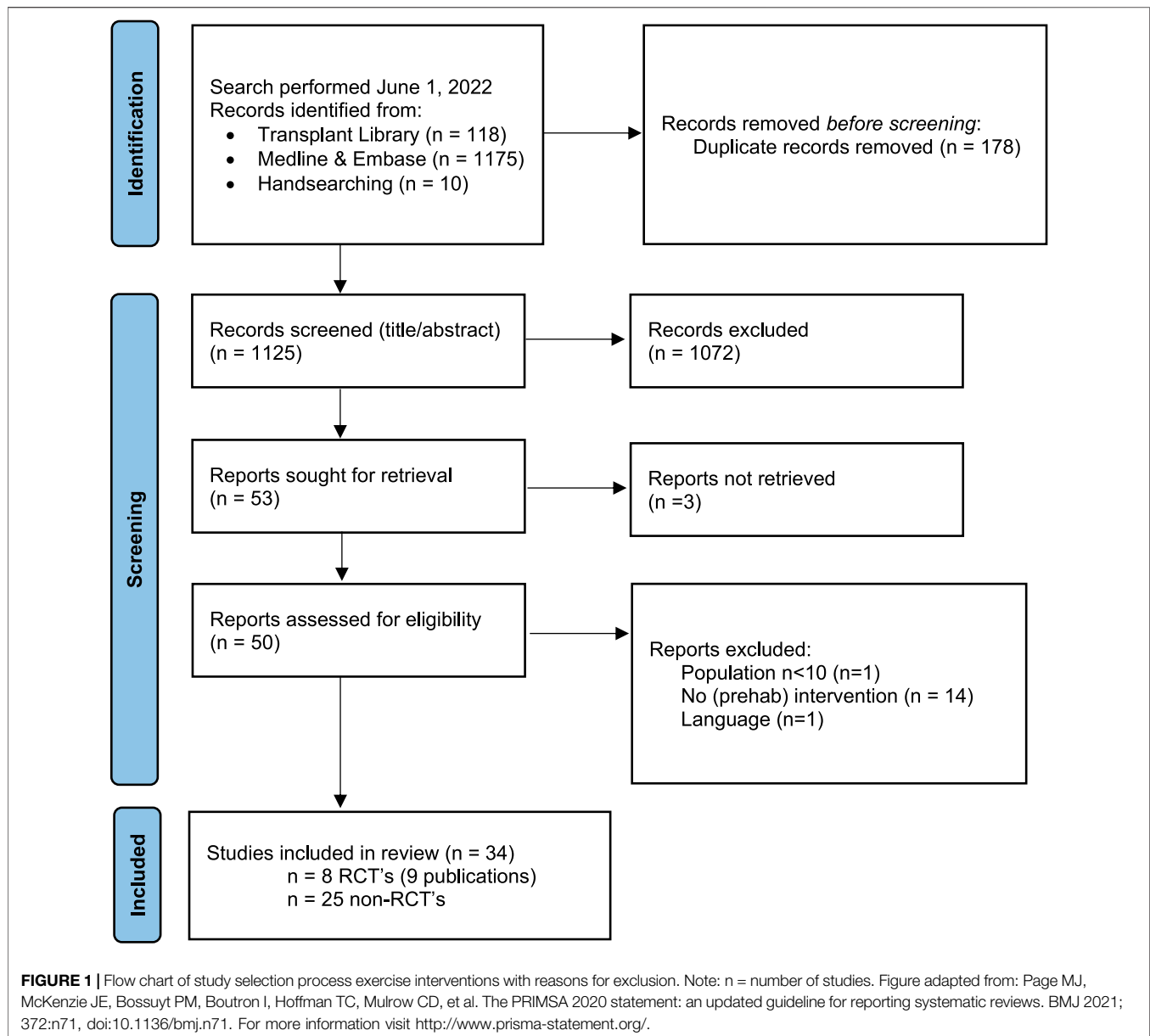
- i) Prehabilitation for solid organ transplant candidates was selected as a priority topic for the first ESOT Consensus Conference, as published [14].
- ii) A specific steering committee was selected, consisting of experts in the topic field, members from the Centre for Evidence in Transplantation, a Young Professional in Transplantation representative, and a guideline taskforce member to liaise with ESOT.
- iii) The steering committee identified key relevant questions related to prehabilitation of solid organ transplant candidates (heart, lung, liver, kidney) using to the Population, Intervention, Comparator and Outcome (PICO) methodology [15] (Table 1).
- iv) The staff of the Centre for Evidence in Transplantation performed systematic literature reviews that were informed by the PICO questions and thus related to exercise, nutritional and psychological interventions in solid organ transplant candidates. The search strategy is presented in Supplementary Table S1. The PRISMA diagrams from the evidence review are shown in Figures 1–3. As the number of publications was expected to be limited, selection was not limited to randomized clinical trials but also included studies that used a pre/post or case-control design, prospective and retrospective studies (cohorts or registry), feasibility studies and pilot studies. Reviews and meta-analyses were included for hand searching of bibliographies for additional literature. Studies were included only if a minimum of 80% of study participants were formally waitlisted for a solid organ transplant. Case reports on fewer than 10 patients, conference abstracts, and letters to the editor were excluded, as was non-English literature. The literature evidence relating to the PICO questions was summarized, as shown in Tables 2–4 and Supplementary File S1.
- v) The steering committee integrated the literature evidence and formulated recommendations (Supplementary File S1). When proposing recommendations for each question, the quality of evidence was considered as evaluated by the GRADE approach [16]. This included risk of bias (Figures 4–6), which was assessed by two independent reviewers, and an additional third one if disagreement occurred. The strength of the individual recommendations was rated as strong or weak.
- vi) Jury members, who were not part of the steering committee were selected and vetted by the guideline taskforce and were comprised of allied health professionals, patients (representatives), transplant physicians, and transplant surgeons.
- vii) Consensus was generated using discussion within the entire working group and modified Delphi methodology including consensus polling, followed by jury voting of the recommendations during a session at the ESOT Consensus conference in Prague [17].
- viii) A committee of validating experts validated the recommendations using the AGREE II guidelines [18].

RESULTS

A total of 4 PICO questions were identified, along with key criteria for analysis, as presented in Table 1. The systematic

TABLE 1 | PICO questions and criteria for analysis.

1	In adult candidates for lung, liver, kidney and heart transplantation: What is the evidence for the effectiveness of pre-transplant exercise training, nutritional support and psychosocial interventions, as measured by the criteria prehabilitation efficacy outcomes, clinical outcomes and patient-reported outcomes Criteria for analysis: Effectiveness of the prehabilitation program: Maximal exercise capacity, Functional exercise capacity, Muscle strength, Nutritional status, Body composition, BMI, Cardio metabolic risk profile, Distress (anxiety/depression), Fatigue, Frailty Clinical outcomes: Mortality (pre-/post-transplant); Hospital (re-)admissions (pre-/post-transplant); Length of hospital stay (pre-/post-transplant); Complications after transplant surgery; Graft survival; Rejection episodes Patient reported outcomes: Health related Quality of Life (HRQoL), Activities of daily living
2	In adult candidates for lung, liver, kidney, and heart transplantation: What is the evidence for the type of pre-transplant exercise, nutritional support and psychosocial interventions?
3	In adult candidates for lung, liver, kidney, and heart transplant candidates: Which relevant outcomes need to be measured to evaluate the effect of the pre-transplant exercise and physical therapy, nutritional support and psychosocial interventions?
4	In adult candidates for lung, liver, kidney and heart transplantation: What is the evidence for the feasibility of prehabilitation, as measured by the criteria enrolment, retention, acceptability, fidelity, safety? Criteria for analysis: Enrolment: the number of screened patients who met the eligibility criteria (n%), the number of eligible patients who were recruited for the study (n%) Retention: the number of participants that were retained in the intervention study, drop-out rate (n%), reasons for drop-out Acceptability: the perception among professionals and participants that the intervention is agreeable, appropriate, or satisfactory Fidelity: the degree to which the intervention was implemented as it was intended, as measured by adherence to the program protocol by the interventionist and participants, Safety: occurrence of adverse events



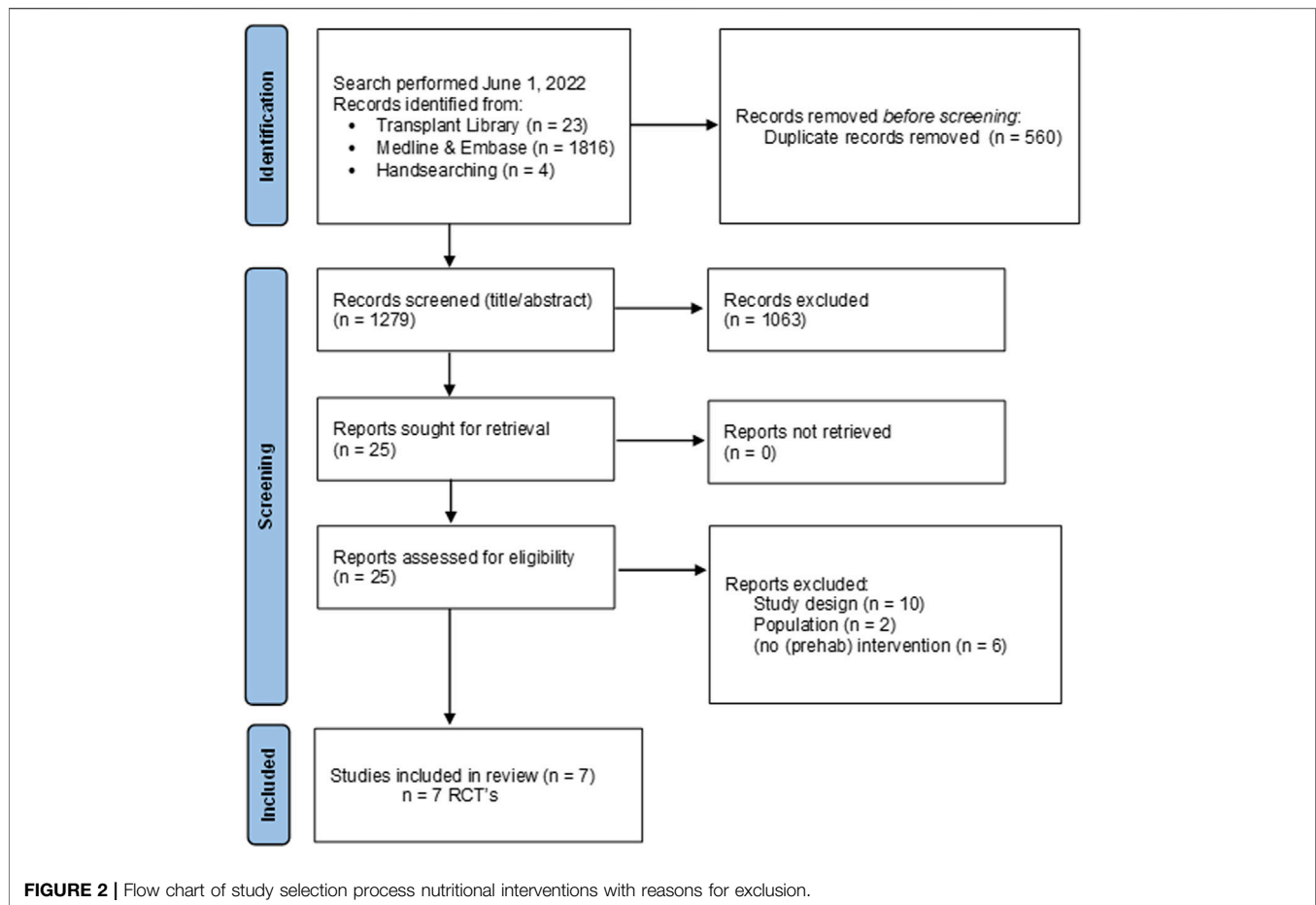
review of literature yielded 34 studies on exercise, 7 on nutritional, and 10 on psychological interventions (Figures 1–3). Summaries of the literature evidence were generated and are presented in Tables 2–4 and Supplementary File S1. A total of 26 recommendations were formulated (Supplementary File S2). At the Consensus Conference, the literature summaries and recommendations were presented, discussed, and amended according to the ESOT consensus-finding process. In response to the considerations voiced during the discussion, and in an attempt to avoid overlap, the number and nature of the recommendations was revised to 10 well-defined recommendations, i.e., 4 general and 6 specific ones. In a first voting round, 100% agreement was achieved on 7 out of 10 recommendations, whereas 3 recommendations reached 86% agreement (1.2, 2.4, 2.5) due to being considered as too

exclusive of certain patient groups. Consensus was reached to amend these recommendations to be more inclusive, and in a second voting round, 100% agreement was achieved on all 10 recommendations.

RECOMMENDATIONS

PICO Question 1

In adult candidates for lung, liver, kidney, & heart transplantation: what is the evidence for the effectiveness of pre-transplant exercise training, nutritional support and psychosocial interventions as measured by prehabilitation efficacy outcomes, clinical outcomes and patient-reported outcomes.



To date, multimodal prehabilitation programs that offer a combination of exercise, nutritional, and psychosocial interventions, have not been studied in solid organ transplant candidates. Rather, literature is limited to studies investigating a single type of intervention. Based on the committee's literature review and analysis of the predefined criteria prehabilitation effectiveness, clinical and patient-reported outcomes, one general recommendation and two specific recommendations were made.

Recommendation 1.1

Studies are needed that evaluate multi-modal prehabilitation interventions in candidates for all types of solid organ transplantation and that focus on core outcomes and implementation. Such studies should be of high quality, and preferably—but not exclusively—adequately powered RCTs.

Quality of Evidence: not applicable.

Strength of Recommendation: Strong.

Rationale: Although supportive, the current evidence (Table 2–4) based on the effectiveness of pre-transplant exercise, nutritional, and psychosocial interventions is weak because of the limited number of randomized studies; 8 for exercise interventions [19–25], 7 for nutritional interventions [26–32], and 6 for psychosocial interventions [33–38]. In

addition, 25 non-randomized studies regarding exercise [39–64], and 4 non-randomized studies on psychosocial interventions [65–68] were retrieved by literature review (Supplementary File S1). The small sample size per study and the limited size and heterogeneity of the total populations studied, the variability in interventions and outcomes measures, the generally low-to-moderate quality of the methodology, and—as a result—the inconsistency of findings across studies (Tables 2–4), warrants high-quality studies on multimodal prehabilitation before solid organ transplantation.

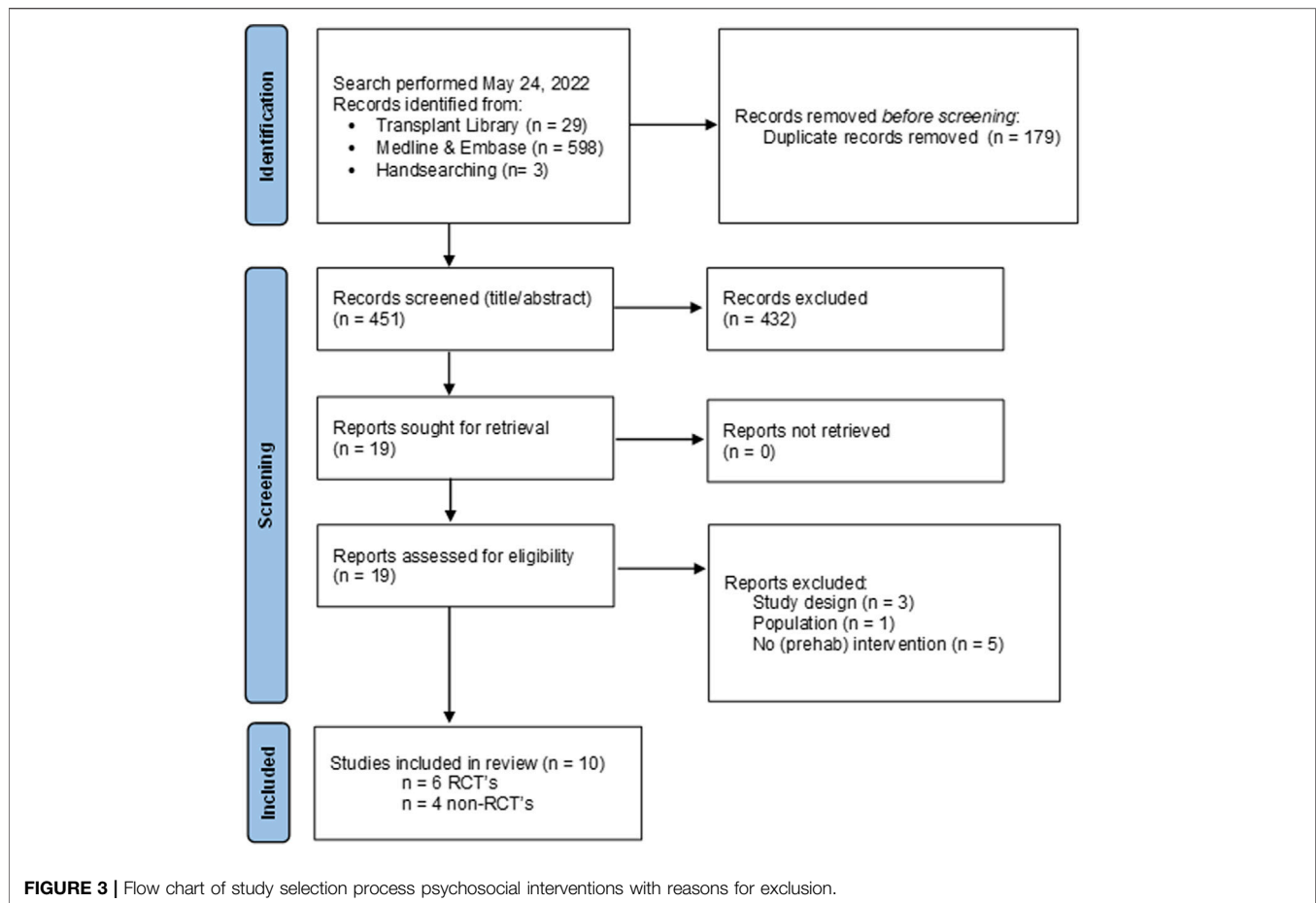
Recommendation 1.2

It is suggested that exercise-based interventions are included in the prehabilitation care of solid organ transplant candidates, with the objective to improve cardiorespiratory fitness and/or inspiratory muscle strength.

Quality of Evidence: Low.

Strength of Recommendation: Weak.

Rationale: Although the number and size of RCTs is limited, studies have shown that exercise training was associated with clinically meaningful improvement in cardiorespiratory fitness in heart transplant candidates [19, 21, 22, 24] and a clinically meaningful gain in inspiratory muscle strength in heart and in liver transplant candidates [19, 23, 24].



Recommendation 1.3

It is suggested that probiotic therapy be used in candidates for liver transplantation to reduce their susceptibility to post-transplant infections.

Quality of Evidence: Very low.

Strength of Recommendation: Weak.

Rationale: Two studies were identified in which pre-transplant probiotic and symbiotic therapy were associated with reduced post-transplant infection rates in recipients of a liver transplant [26, 28]. However, both studies had small sample sizes ($n = 44/n = 50$) and used different products.

PICO Question 2

In adult candidates for lung, liver, kidney, & heart transplantation: which type(s) of exercise, nutritional support and psychosocial interventions are recommended in the pre-transplant phase?

As there are no established prehabilitation programs for solid organ transplant candidates, evidence review was focused on studies that addressed interventions that could be of value in a multimodal prehabilitation program. One general recommendation and four specific recommendations were established.

Recommendation 2.1

Studies are needed to identify the optimal component(s) and the mode of delivery of pre-transplant multimodal prehabilitation

programs in solid organ transplant candidates. Such studies should be of high quality and be preferably -but not exclusively-adequately powered RCTs.

Quality of Evidence: not applicable.

Strength of Recommendation: Strong.

Rationale: Because of the heterogeneity in the study populations and in the nature and delivery mode of the interventions described in the current literature (**Tables 2–4; Supplementary File S1**), it remains unclear which organ transplant candidates would benefit most from which intervention program. Most exercise intervention studies used aerobic training [19–22, 24], peripheral muscle training [41], inspiratory muscle strength training [25, 51], or a combination of these training modalities. Nutrition intervention studies mostly used nutritional support to optimize energy intake and/or obtain weight loss [29–32, 69, 70]. Whilst psychosocial interventions predominantly included cognitive behavioral therapy [33, 34, 36, 37, 65, 67], psycho-educational interventions [35, 68] and stress management and relaxation techniques [38, 67] or a combination of these interventions. Studies are needed that will help determine the modalities of the intervention, and for each modality (exercise, nutrition or psychosocial), the intervention characteristics (frequency, intensity and timing), the delivery mode (type of interventionist, level of supervision, home-based versus in- or outpatient) for each type of donor organ recipient.

TABLE 2 | Summary of RCTs- Exercise interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – Feasibility outcomes
1 Laoutaris, 2011 (Greece)	HTx candidates with LVAT or BIVAT n = 21 I n = 14 100% male Age 37 (±18) C n = 7 80% male Age 42 (±15)	All participants were advised to walk every day for 30–45 min. I: 10 week, homebased aerobic exercise (45 min, 3–5x/week, intensity y12–14/20 RPE) and hospital-based IMT (until exhaustion, 2–3x/week, intensity 60% MIP) C: Usual care Assessments: - Pre-intervention - Post-intervention	VO ₂ peak 6MWT QOL (MLwHFQ) PIMax and sustained PImax Lung volumes (inspiratory capacity) Dyspnea after 6MWT	↑ within I; I ≈ C ↑ within I; I ≈ C ↑ within I; I ≈ C ↑ within I; I ≈ C ↑ within I; I ≈ C I ≈ C	Enrolment: 21/23 (91%) of eligible patients Attrition: 15/21 (71%) completed (I 10/14 (71%); C 5/7 (71%)), all drop out due to Tx Fidelity (participants): NR Fidelity (interventionist): NR Acceptability: patients enjoyed training and seemed more enthusiastic compared with patients in the control group Safety: no adverse events occurred during the training period
2 Gloeckl, 2012 (Germany)	LuTx candidates following COPD stage IV diagnosis n = 71 I n = 36 49% male Age 52 (±6) C n = 35 44% male Age 55 (±7)	All participants received strength training (four to six exercises, 3 sets of 30 repetitions, at maximal tolerated load), breathing therapy, education, and psychological support. I: 3-weeks, hospital-based high-intensity interval training, 10–36 min per session, 5–6x/week, 1–2 sessions/day, intensity repeated bouts of 30 s at 100% Wmax alternated by 30 s rest C: 3-weeks, hospital-based, moderate-intensity aerobic exercise, 10–30 min/session, 5–6x/week, 1-2 sessions/day, intensity 60% Wmax Assessments: - Pre-intervention - Post-intervention	6MWT QOL (SF-36 PCS and MCS) W _{max} During exercise - SpO ₂ - TCPCO ₂ - Dyspnea - leg fatigue Unintended breaks (number and time) during exercise PaO ₂ and PaCO ₂ lung function (DLCO, FEV1, FEV1/IVC)	≈ increase in I and C PCS: ≈ within I; ↑ within C; I ≈ C MCS: ↑ within I; ≈ within C; I ≈ C ≈ increase in I and C I ≈ C I ≈ C ↓ in I I ≈ C ↓ number and ↓ duration in I I ≈ C I ≈ C	Enrolment: 71/97 of eligible patients Attrition: 60/71 completed (I 30/36 (83%); C 30/35 (86%)). Dropout due to: I acute exacerbation (n = 4), non-compliance with study protocol (n = 1), other (n = 1); C acute exacerbation (n = 3), Tx (n = 1), other (n = 1) Fidelity (participants): no difference in number of exercise sessions or total work performed per group. I: 14.9 (±1.9); C: 14.7 (±1.5) Fidelity (interventionist) NR Acceptability: NR Safety: no serious adverse events occurred

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TABLE 2 | (Continued) Summary of RCTs- Exercise interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – Feasibility outcomes
3 Hayes, 2012 (Australia)	HTx candidates with LVAT n = 14	All participants followed a progressive walking program: They were advised to walk a minimum of 5 days per week at 13 RPE and increased their walk progressively up to 60 min.	VO ₂ peak	↑ in; I ≈ C	Enrolment: 14/18 (78%) of eligible patients
	I n = 7 86% male Age 48.7 (±14.5)	I: 8-week, gym-based aerobic and strength training (60 min/session, 3x/week, intensity cycling at 50% VO ₂ reserve; treadmill at 60% of the speed averaged during the 6MWT; strength: three upper limb and three lower limb exercises using weight machines and free weights, 2 sets of 10 reps).	6MWT	↑ in I and C; I ≈ C	Attrition: 14/14 (100%) completed
	C n = 7 86% male Age 45.9 (±14.6)	C: Usual care, which included a walking program.	QOL (SF36)	↑ within I; I ≈ C	Fidelity (participants): I: participation in 21.3 ± 1.5 of possible 24 sessions. Reasons for missed sessions: conflicting medical appointment (79%) and conflicting family demands (21%). C: 100% compliance to the walking program Fidelity (interventionist): NR
	Assessments: - Pre-intervention - Post-intervention	W _{max}	↑ in I and C	Acceptability: NR Safety: no adverse events occurred	
4 Adamopoulos, 2013 (Greece)	HTx candidates with LVAT or BIVAT n = 22	All participants were advised to walk every day for 30–45 min.	Thyroid hormone signalling (TRα1, p/t-AKT and p/t-JNK)	↑ within I; I ↑ C	Enrolment: 22/26 (85%) of eligible patients
	I n = 11 91% male Age 39.7 (±4.3)	I: 12 weeks, aerobic training (home-based, 45 min/session, 4x/week, intensity 12-14/20 RPE) and IMT (hospital-based, until exhaustion, 3x/week, intensity 60% P _{lmax})	VO ₂ peak	↑ within I; I ↑ C	Attrition: 22/22(100%) completed
	C n = 11 82% male Age 40.9 (±4.9)	C: Usual care	NT-proBNP (marker of heart failure)	↓ within I; I ↓ C	Fidelity (participants): NR
	Assessments: - Pre-intervention - Post-intervention			Fidelity (interventionist): NR Acceptability: NR Safety: NR	

(Continued on following page)

TABLE 2 | (Continued) Summary of RCTs- Exercise interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – Feasibility outcomes
5 Limongi, 2014 and 2016 (Brazil)	LiTx candidates n = 49 I n = 22 79% male Age 55.8 (±5.4) C n = 27 78% male Age 55.4 (±9.9)	I: 3-months, home-based, daily exercises illustrated in a manual (3 × 15 repetitions of diaphragmatic breathing exercises, diaphragmatic isometric exercise, Threshold IMT®, lifting upper limbs with a bat and strengthening the abdomen). Duration training sessions varied by patient. Intensity reported only for diaphragmatic breathing (1 kg on the belly). Supervision once a month at distance. C: Usual care Assessments: - Pre-intervention - Post-intervention	MIP MEP Spirometry (FVC, FEV1) QOL (SF-36) Surface EMG of diaphragm Surface EMG of rectus abdominis Ascites presence	↑ in I and C ↑ in I and C no changes ↑ in I and C on general health and mental health subscale; ↑ within I on functional capacity, not in C, but without between group differences I ↓ C no changes I ≈ C	Enrolment: 49/49 (100%) of eligible patients Attrition: 37/49 (76% completed; I 14/22 (64%); C 23/27 (85%)). Dropouts due to I: LiTx (n = 2), death (n = 3), declined to perform exercise (n = 3), C LiTx (n = 1), death (n = 3) Fidelity (participants): NR Fidelity (interventionist): NR Acceptability: NR Safety: NR
6 Forestieri, 2016 (Brazil)	HTx candidates n = 24 I n = 12 71% male Age 48.3 (±10.2) C n = 12 82% male Age 48.0 (±11.2)	I: ~22 days, hospital-based, intermittent aerobic (stationary cycle ergometer exercise: 5 periods of 3 min cycling and 1 min res, 20 min/session, 2x/day, intensity 3–4/10 RPE C: ~19 days, hospital-based, breathing exercises and global active exercises of the upper and lower limbs in the upright seated position (2x/day, intensity: 3–4/10 RPE) Assessments: - Pre-intervention - Post-intervention	6MWT MIP FVC FEV1 NT-proBNP	↑ within I; I ↑ C ↑ within I; I ↑ C NR NR NR	Enrolment: 24/27 (89%) of eligible patients Attrition: 18/24 (75%) completed (I 7/12 (58%); C 11/12 (92%)). Dropouts due to: I: incapacity to complete the stationary cycle ergometer exercise (n = 5); C: acute severe arrhythmias (n = 1) Fidelity (participants): 42% lost for follow-up Fidelity (interventionist): NR Acceptability: 42% were incapable to complete the intervention program Safety: NR

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TABLE 2 | (Continued) Summary of RCTs- Exercise interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – Feasibility outcomes
7 Pehlivan, 2018 (Turkey)	LuTx candidates n = 34	All participants participated in a home-based pulmonary rehabilitation program: breathing exercises (local expansion exercises, diaphragmatic breathing, and pursed lip breathing), free walking, and upper and lower body strengthening with resistance bands. Participants completed a weekly chart that was reviewed by the physiotherapist. I: 3-months, 5x/week (supervised 2x/week; home-based 3x/week) standard pulmonary rehabilitation (aerobic exercises: treadmill, cycle and arm ergometer, 15 min per exercise modality/ session, intensity: 50%–70% of HRmax and resistance exercises: dumbbell and free weight bags, 8–12 reps, one to two sets/ session, intensity 20%–40% 1-RM) + IMT with Powerbreathe device (15 min/ session, 2x/day, 5 days/week, intensity initial 30% of MIP, progressed to 60% MIP) C: Usual care, including standard pulmonary rehabilitation (see above) Assessments: - Pre-intervention - Post-intervention	MIP	↑ within I; I ↑ C	Enrolment: 34/38 (89%) of eligible patients
	I n = 17 59% male Age 36.1 (±15.9)		MEP	↑ in I and C	Attrition: 34/34 (100%) completed
	C n = 17 65% male Age 39.0 (±12.4)		6MWT	↑ in I and C, but greater in I than C	Fidelity (participants): NR
			mMRC dyspnea scale	↓ in I and C	Fidelity (interventionist): NR
			FVC FEV1 DLCO DLCO/VA	no changes no changes no changes I ↑ C	Acceptability: NR Safety: NR
8 Manzetti, 1994 (United States)	LuTx candidates n = 21 22% male Age 40 (±10)	I: 6-week, health education program + supervised aerobic training (treadmill, bicycle ergometer, 30 min/session, 2x/ week, around aerobic threshold or 80% maximal ventilation) + strength training of upper extremity (low intensity)	W _{max}	no changes	Enrolment: 36/91 (40%) eligible for participation, 15/36 (42%) of eligible patients declined participation due to financial reasons (n = 10) or transport issues or inability to perform activities of daily living independently (n = 5) Attrition: 9/21 (43%) completed; drop-outs due to Tx (n = 9) or hospitalization (n = 3). Number of drop-outs per group NR
	I: n = 5 % Male NR Age NR		6MWT	↑ in I and C, I ≈ C ↑ or ≈ I and C, I ≈ C	Fidelity (participants): NR (Continued on following page)

TABLE 2 | (Continued) Summary of RCTs- Exercise interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – Feasibility outcomes
	C: n = 4 % male NR Age NR	C: 6-week, health education program Assessments: - Pre-intervention - Post-intervention	QoL (QWB, QLI, SFSD)		Fidelity (interventionist): NR Acceptability: NR Safety: NR

I, Intervention group; *C*, comparator group; *Tx*, transplantation; *NR*, not reported; *1RM*, one-repetition maximum; *6MWD*, six-minute walking distance; *6MWT*, six-minute walking test; *BIVAT*, biventricular assist device; *COPD*, chronic obstructive pulmonary disease; *DLCO/VA*, alveolar volume ratio of carbon-monoxide diffusion capacity; *DLCO*, diffusion capacity of the lung for carbon monoxide; *EMG*, electromyography; *FEV1*, forced expiratory volume in 1 second; *HRmax*, maximal heart rate; *HTx*, heart transplantation; *IMT*, inspiratory muscle training; *IVC*, inspiratory vital capacity; *LiTx*, liver transplantation; *LuTx*, lung transplantation; *LVAT*, left ventricular assist device; *MEP*, maximal expiratory pressure; *MIP*, maximal inspiratory pressure; *MLwHFQ*, Minnesota Living with Heart Failure Questionnaire; *mMRC* dyspnea scale, modified Medical Research Council dyspnea scale; *NA*, Not applicable; *NR*, not reported; *NT-proBNP*, N-terminal prohormone of brain natriuretic peptide; *P_aCO₂*, partial pressure arterial carbon dioxide; *P_aO₂*, partial pressure arterial oxygen; *P_lmax*, maximal inspiratory pressure; *QOL*, quality of life; *RPE*, rate of perceived exertion; *SpO₂*, saturation of peripheral oxygen; *T_cPCO₂*, transcutaneously measured pressure of arterial carbon dioxide; *VO₂peak*, peak or maximal oxygen consumption; *W_{max}*, peak work rate at the end of a cardiopulmonary exercise test; *QWB*, Quality of Well-being scale; *QLI*, Quality of Life Index; *SFSD*, Symptom Frequency/Symptom distress scale.

Recommendation 2.2

Solid organ transplant candidates who are underweight may be offered nutritional interventions with the aim to achieve optimal target weight before the transplant.

Quality of Evidence: Very low.

Strength of Recommendation: Weak.

Rationale: Evidence from two intervention studies in lung transplant candidates [29, 30] have indicated that increased caloric intake before transplantation may allow solid organ transplant candidates, especially those who are underweight, to reach a pre-transplant target weight. However, these studies had a small sample size and were conducted in different settings (hospital vs. outpatient clinic).

Recommendation 2.3

Solid organ transplant candidates who are overweight may be offered nutritional interventions with the aim to achieve optimal target weight before the transplant.

Quality of Evidence: Very low.

Strength of Recommendation: Weak.

Rationale: One study ($n = 43$) [32] showed that a weight-loss program, consisting of bibliotherapy and voice call counselling by a dietician, was successful in reducing body weight in adult candidates for heart transplantation.

Recommendation 2.4

It is suggested that cognitive behavioral therapy and psychoeducational interventions are considered for solid organ transplant candidates who have symptoms of anxiety and/or depression.

Quality of Evidence: Very low.

Strength of Recommendation: Weak.

Rationale: Six studies utilized elements of cognitive behavioral therapy (CBT) and psychoeducational interventions [33, 34, 36, 37, 65, 67] of which five reported a significant decrease in symptoms of anxiety and depression or mood [33, 34, 36, 37, 65] in lung, liver, and kidney transplant candidates. However, studies differed regarding duration (8–12 weeks), modality (group vs. individual; remote vs. in person), and most studies had small sample sizes ($n = 29$ to $n = 71$) (Table 4). Only the study of Blumenthal et al (2006) [36] had an adequate sample size ($n = 328$).

Recommendation 2.5

It is suggested to consider stress-reducing interventions such as mindfulness-based stress reduction or relaxation techniques in candidates for solid organ transplantation to reduce anxiety or stress levels.

Quality of Evidence: Very low.

Strength of Recommendation: Weak.

Rationale: In two studies among kidney and kidney-pancreas transplant candidates, stress-reducing interventions were associated with alleviated symptoms of anxiety [65] or depression [38, 65] directly after the intervention. However, this effect was not maintained long-term. In addition, sample sizes were small ($n = 41/n = 63$) and the intervention differed regarding content and interventionist.

PICO Question 3

In adult candidates for lung, liver, kidney, & heart transplantation: what are the outcomes relevant to exercise and physical activity, nutritional support and psychosocial interventions that should be measured pre-transplant?

In order to reliably assess the effects of prehabilitation interventions, it is imperative to standardize outcome

TABLE 3 | Summary of RCTs- Nutritional interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness Outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – feasibility outcomes
1 Grat, 2017 (Poland)	Liver Tx candidates N = 55 I n = 26 81% male Age 52 (47–58) C n = 29 74% male Age 50 (35–61)	I: once daily intake of a 4-strain probiotic preparation before breakfast (ProBacti 4 Enteric®: 3 × 10 ⁹ colony-forming units of <i>Lactococcus lactis</i> PB411 (50.0%), <i>Lactobacillus casei</i> PB121 (25.0%), <i>Lactobacillus acidophilus</i> PB111 (12.5%), and <i>Bifidobacterium bifidum</i> PB211 (12.5%) from enrolment until transplantation. Duration of intervention was <2–>10 weeks depending upon timing Tx C: placebo Assessments: - Baseline - Pre-Tx: follow-up with intervals of 10 weeks - Post-Tx: 90 days follow-up	90-day mortality rate 30-day and 90-day infection rate 5-days post-Tx: - AST - ALT - Bilirubin concentration - INR Pre-transplant: - Waitlist mortality - Hospitalizations - Infections - Complications Post-transplant - Primary non-function - Early allograft dysfunction - Complications MELD-score changes CTP changes	I ≈ C I ↓ C I ↑ C I ↑ C I ↓ C I ≈ C None I ≈ C I ≈ C I ≈ C I ≈ C I ≈ C I ≈ C I ≈ C I ≈ C	Enrolment: 209/491 (43%) eligible for participation; 55/209 (26%) of eligible patients participated. Refusal to participate probably due to administrative factors Attrition: 50/55 (91%) completed (I 24/26 (92%); C 26/29 (90%)). Dropouts (n = 5) all discontinued treatment Post-Tx outcomes available of I: 21/26 (81%) and C: 23/29 (79%) Fidelity (participants): I 2/26 (8%) and C: 3/29 (10%) discontinued treatment Fidelity (interventionist): NR Acceptability: NR Safety: NR
2 Plank, 2015 (New Zealand)	Liver Tx candidates N = 101 I n = 52 Male 63% Age 53 (25–68) C n = 49 Male 73% Age 50 (22–59)	I: daily intake of immuno-nutrition, two 74 g sachets per day until the day of transplant, consisting of 7.5 g arginine, 2 g omega-3 fatty acids + 0.8g Ribonucleic acid. for 56–65 days (median) C: daily intake with a similar amount of an isocaloric, but not isonitrogenous, control product Assessments: - Baseline - Prior to Tx - 10, 30, 90, 180, 360 days after Tx	Body composition - Body weight (kg) - Total body protein - Total body fat Muscle function - Hand grip strength - Respiratory muscle strength Plasma phosphatidylcholine fatty acids Fatigue (NR) Graft rejection Length of stay at ICU Length of stay at hospital	I ≈ C I ≈ C I ≈ C I ≈ C I ≈ C I ↑ C at pre-Tx and day 10 measurements I ≈ C I ≈ C I ≈ C I ≈ C	Enrolment: NR Attrition: 101/120 (84%) completed (I 52/60 (87%); C 49/60 (82%)). Dropouts: I: delisting (n = 8), C: death (n = 4), delisting (n = 7) Fidelity (participants): NR Fidelity (interventionist): NR Acceptability: NR Safety: intolerance to immune-nutrition in four participants

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TABLE 3 | (Continued) Summary of RCTs- Nutritional interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness Outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – feasibility outcomes
3 Eguchi, 2011 (Japan)	Living donor Liver TX candidates N = 50 I n = 25 52% male Age 56 (33–66) C n = 25 64% male Age 57 (25–68)	I group 1: 2 days preoperative and group 2: 2 weeks post-operative synbiotic therapy (<i>Bifidobacteriu breve</i> , <i>Lactobacillus casei</i> and <i>Galactooligosa charides</i>) C: placebo Assessments: Not specified	Infectious complications Mortality Length of stay at ICU Length of stay at hospital	I ↓ C I ≈ C I ≈ C I ≈ C	Enrolment: NR Attrition: NR Fidelity (participants): NR Fidelity (interventionist): NR Acceptability: NR Safety: all participants tolerated synbiotic therapy
4 Park, 2003 (United States)	Heart Tx candidates with BMI > 25 kg/m ² N = 43 I n = 21 81% male Age 47.8 (±8.5) C n = 22 68% male Age 48.1 (±9.4)	All participants had one consultation session by a graduate student in clinical psychology under the supervision of the study's registered dietitian, who provided the recommendations such as energy balance I : 3-months weight-loss program comprised of bibliotherapy (written, 20-page manual containing brief lessons about cognitive and behavioral weight loss strategies), and telephone-based counseling (1x/week, 15–20 min) delivered by a therapist who has a bachelor's or master's degree in psychology. C: 3-months weight-loss program comprised of bibliotherapy without counseling Assessments: - Pre-intervention - Post-intervention	Body weight change	I ↑ C	Enrolment: 43/54 (80%) of referred patients Attrition: 36/43 (84%) completed (I 17/21 (81%); C 19/22 (86%)) Fidelity (participants): I returned more food diaries than C, but not significant; I returned more postcards than C, but not significant Fidelity (interventionist): NR Acceptability: NR Safety: NR
5 Forli, 2001(a) (Norway)	Lung Tx candidates N = 65 I n = 18 44% male Age 49 (44–53) C1 n = 19 53% male Age 48 (44–52) C2 n = 28 43% male Age 51 (48–55)	I: intensified nutritional support comprised of energy-rich diet and supplements, provided by a dietician during hospital stay C1: normal hospital diet C2: normal weight lung Tx candidates Assessments: - During hospitalization for lung Tx screening, exact moments NR	Change in body weight. BMI (kg/m ²) Total energy intake/kg Total energy intake/REE predicted	I ↑ C1 and C2 C1 ↓ I and C2 C2 ↓ I and C1 I ↑ C1 and C2, C1 ↑ C2	Enrolment: 6/71(8%) of eligible patients excluded for various reasons: refused intervention (n = 1), dietary wishes (n = 1), absent during night/weekend (n = 1), short hospital stay (n = 1), death (n = 1). Attrition: 49/65 (75%) completed. Drop-outs due to: not willing to record data (n = 1), missing data (n = 3), oedema (n = 2), death (n = 1) Fidelity (participants): NR Fidelity (interventionist): NR Acceptability: NR Safety: NR

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TABLE 3 | (Continued) Summary of RCTs- Nutritional interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness Outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ = no difference	Results – feasibility outcomes
6 Forli, 2001(b) (Norway)	Lung Tx candidates with underweight N = 71 I n = 21 48% male Age 47 (28–59) C1 n = 21 48% male Age 46 (25–60) C2 n = 29 41% male Age 52 (26–60)	I: intensified sessions dietary counselling with suggestions for individual meal plans facilitating weight gain, booklet with dietary information and recipes, supplements, and support by telephone by the dietitian each month after hospital discharge. Mean intervention time was 22 weeks. C1: one session of individual dietary counselling with the dietitian. No follow-ups. The mean intervention time was 20 weeks. C2: normal weight Lung Tx candidates Assessments: - Pre-intervention - 4–5 months after discharge	Body composition - Change in body weight - Change in Fat mass - Change in Fat free mass Blood samples - Albumin concentration - Phosphate concentration Lung function test: - PaO ₂ - PaCO ₂ - FVC - FEV1 - TLCO Exercise testing: - handgrip strength - 6MWT	↑ in I (+2.9 kg) and C1 (+2.3 kg) group, not in C2 group I ↑ C1 ≈ C2 C1 ↑ ≈ C2 I ≈ C1 ≈ C2 I ≈ C1 ≈ C2 I ↓ C1 ≈ C2 I ≈ C1 ≈ C2 I ≈ C1 ≈ C2 I ≈ C1 ≈ C2 I ≈ C1 ≈ C2 I ≈ C1 ≈ C2	Enrolment: NR Attrition: 54/71 (76%) completed (I 18/21 (86%); C1 13/21 (62%); C2 23/29 (79%)). Dropouts: I and C1 death (n = 8), Tx (n = 3), infection (n = 14); C2 death (n = 2), Tx (n = 4), infection (n = 7) Fidelity (participants): NR Fidelity (interventionist): NR Acceptability: NR Safety: NR
7 Le Cornu, 2000 (United Kingdom)	Liver Tx candidates N = 82 I n = 42 69% male Age 52 (27–67) C n = 40 79% male Age 50 (24–68)	No information who provided the advice in both groups nor the number of sessions I: Standard dietary advice to increase energy intake on top of the dietary recommendations they already had to follow for underlying medical conditions and daily enteral supplementation (750 calories out of 20 g protein and 33.5 g fat). C: Standard dietary advice to increase energy intake on top of the dietary recommendations they already had to follow for underlying medical conditions. Assessments: - Screening - Monthly follow-up until Tx or death	Biochemical parameters - Bilirubin - Creatinine - Urea - Alkaline phosphatase - Aspartate transaminase - INR Anthropometric measurements: - Mid-arm circumference - Triceps skinfold thickness Handgrip strength Energy Intake Survival (pre-transplant) Days on ventilatory support Length of ICU stay Length of Hospital stay	≈ within ≈ within I and C ↑ within I; ≈ within C ≈ within I; ↓ within C ≈ within I and C ≈ within I; ↑ within C I ≈ C I ≈ C I ≈ C I ≈ C I ≈ C I ≈ C I ≈ C	Enrolment: 116/328 (35%) patients were eligible, 82/116 (71%) of eligible patients consented Attrition: 80-28 (98%) completed (I 41/42 (98%); C 39/40 (98%)). Dropouts due to I: lost to follow-up n = 1; C: delisted (n = 1) Fidelity (participants): NR Fidelity (interventionist): NR Acceptability: NR Safety: NR

I, Intervention group; C, comparator group; Tx, transplantation; NR, not reported; AST, aspartate; ALT, alanine aminotransferase; INR, Internationalized Normalized Ratio; MELD, Model for End-stage Liver Disease; CTP, Chil-Turcotte-Pugh; ICU, Intensive Care Unit; BMI, Body Mass Index; PaO₂, Arterial O₂; PaCO₂, Arterial CO₂; FVC, Forced Vital Capacity; FEV1, Forced Expiratory Volume/1s; TLCO, Lung transfer factor carbon monoxide; 6MWT, six Minutes Walking Test.

TABLE 4 | Summary of RCTs - Psychosocial interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness Outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – feasibility outcomes
1 Napolitano, 2002 (United States)	Lung Tx candidates N = 71		Health-related quality of life (SF36, PQLS)	SF36: I ↑ C on overall quality of life, mental health, role limitations due to emotional functioning, and vitality score PQLS: I ↑ C on overall score and subscales psychological functioning and physical functioning	Enrolment: 81/91 (89%) of eligible patients
	I n = 36 31% male Age 44.2 (±12.7)	I: 8-weeks, weekly, telephone-based psychological treatment comprised of supportive counselling and CBT, delivered by clinical psychology graduate	Anxiety (GHQ) Depression (GHQ)	I ↓ C on total score and subscales scores (anxiety, depression, social dysfunction, and somatic symptoms)	Attrition: 71/81 (88%) completed baseline (n = 2 delisted, n = 1 Tx, n = 6 withdrew consent, n = 1 died). 66/71 (93%) completed follow-up (I 34/36 (94%), missing data due to Tx (n = 2); C 32/35 (91%), missing data due to loss to follow-up (n = 3)) Fidelity (participants): all participants received all sessions
	C n = 35 31% male Age 46.6 (±12.4)	C: care as usual Assessments: - Pre-intervention - Post-intervention	Social support (PSSTx) Distress (PSTx)	I ↑ C I ≈ C	Fidelity (interventionist): NR Acceptability: NR Safety: NR
2 Rodrigue, 2005 (United States)	Lung Tx candidates N = 35		Quality of Life (QOLI)	I ↑ C at 1 and 3-month follow-up	Enrolment: 35/58 (60%) of eligible patients
	I n = 17 35% male Age 48.8 (±10.0)	I: 8–12-weeks, weekly, telephone-based CBT delivered by clinical psychology graduate students and interns	Mood (POMS)	I ↓ C at 3 month follow-up	Attrition: 35/35 (100%) at baseline, 31/35 (89%) completed all assessments
	C n = 18 33% male Age 49.0 (±11.3)	C: supportive treatment, delivery NR Assessments: - Pre-intervention - 1 month post-intervention - 3 month post-intervention	Social intimacy (MSIS) FEV1 Physical functioning (6MWT)	I ↑ C at 1 month follow-up I ≈ C I ≈ C	Fidelity (participants): I: 88% full treatment, C: 89% full treatment Fidelity (interventionist): NR Acceptability: I: high levels of comfort, rapport, helpfulness and convenience, low levels of distraction. 87% would participate again, 32% preference telephone counselling Safety: NR

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TABLE 4 | (Continued) Summary of RCTs - Psychosocial interventions.

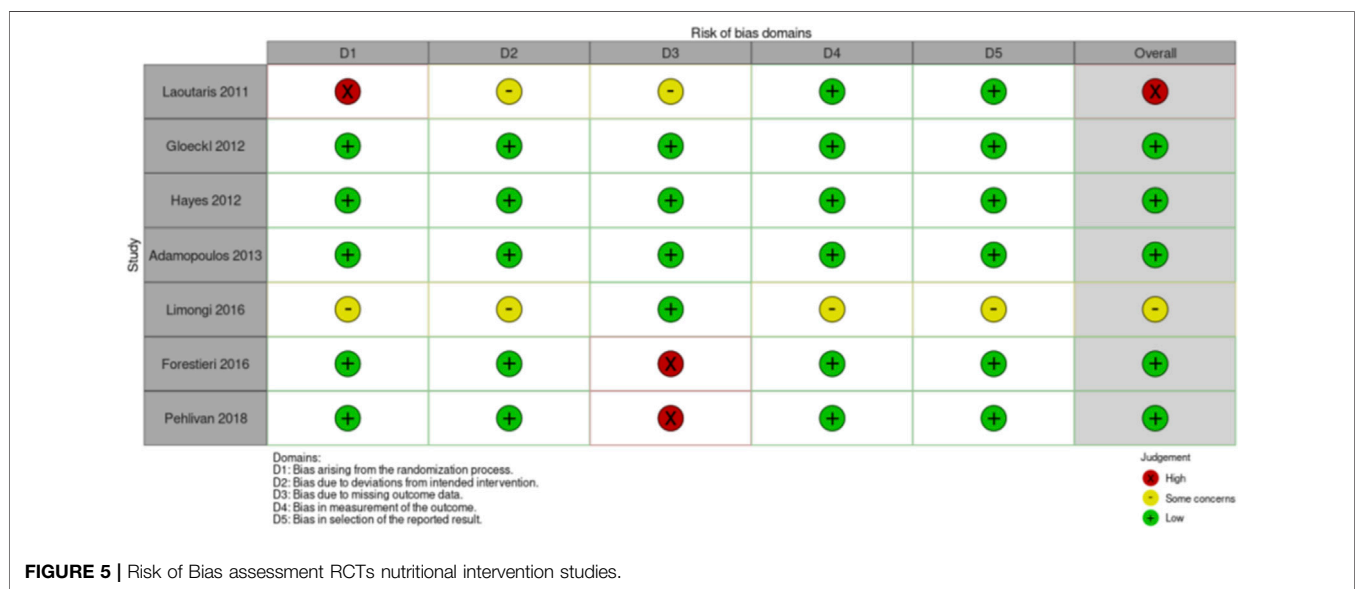
First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness Outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – feasibility outcomes
3 Sharif, 2005 (Iran)	Liver Tx candidates N = 110 I n = 55 76% male Age NR C n = 55 75% male Age NR	I: 4-weeks, three individual sessions and one group session, 90 min/week, psycho-educational treatment, mode of delivery NR C: educational booklet Assessments: - Pre-intervention - Post-intervention	Health-related Quality of life (CLDQoL)	I ↑ scores on domains fatigue, emotional function, and total QoL score at 1-month follow-up. I ↑ on all domains of QoL at 3-month follow-up Comparison with control group not reported	Enrolment: NR Attrition: NR Fidelity (participants): NR Fidelity (interventionist): NR Acceptability: NR Safety: NR
4 Blumenthal, 2006 (United States)	Lung Tx candidates N = 328 I n = 166 45% male Age 50 (±11) C n = 162 43% male Age 50 (±12)	I: 12-weeks, telephone-based, 30 min/week, supportive counseling and training in cognitive-behavioral coping skills coping skills delivered by trained social worker or psychologists C: care as usual Assessments: - Pre-intervention - Post-intervention	Health-related Quality of life (PQLS, SF36, GHQ) Anxiety (STAI) Depression (BDI) Perceived stress (PSS) Life-orientation (LOT-R) Social Support (PSSC) Shortness of breath (SDS-BQ) Survival pre-transplant	PQLS: I ≈ C SF36: I ↑ C on PSC and subscales mental health and vitality; I ≈ C on subscales general health, physical functioning, pain, physical role GHQ: I ↑ C I ↓ C I ↓ C I ≈ C I ↑ C I ≈ C I ≈ C I ≈ C	Enrolment: 389/625 (62%) of eligible patients. Drop out after randomization I: 34/200 (17%); C 27/189 (14%) due to death, Tx or delisting Attrition: I: 126/166 (76%) and C: 147/162 (91%) completed all assessments Fidelity (participants): 10.6 out of 12 sessions, 77% completed all sessions Fidelity (interventionist): 97.6% adhered to the protocol Acceptability: NR Safety: no adverse events

(Continued on following page)

TABLE 4 | (Continued) Summary of RCTs - Psychosocial interventions.

First author, year (country of origin)	Sample characteristics Tx-type, total N, n per group, % male, age (y) (mean (sd) or median (range))	Intervention(s) and measurement points	Effectiveness Outcomes	Results – Effectiveness outcomes ↑ = significant increase ↓ = significant decrease ≈ no difference	Results – feasibility outcomes
5 Rodrigue, 2011 (United States)	Kidney Tx candidates N = 62 I1 n = 22 64% male Age 53.2 (±11.1) I2 n = 20 40% male Age 48.6 (±11.9) C n = 20 45% male Age 52.7 (±12.7)	I1: 8-weeks, in person, 50 min/ week, QoL-therapy, delivered by trained social workers or psychologists I2: 8-weeks, in person, 50 min/ week, supportive care, delivered by trained social workers or psychologists C: care as usual Assessments: - Pre-intervention - 1-week post-intervention - 12-week post-intervention	Health-related Quality of life (QoL, SF-36) Mood (POMS) Distress (HSCL) Social Intimacy (MSIS) No mental unhealthy days	QoL: I1 ↑ (clinical relevant) C, I2 ≈ C SF36: I1 ↑ (clinical relevant) C, I2 ≈ C I1 ↑ (clinical relevant) C, I2 ≈ C I1 ↑ (clinical relevant) C, I2 ≈ C I1 ≈ I2 ≈ C	Enrolment: 65/110 (59%) of eligible patients (n = 18 excluded, n = 27 refused) Attrition: 62/65 (95%) completed baseline, 51/62 (82%) completed all assessments Fidelity (participants): I1 17/22 (77%) and I2 17/20 (85%) received full treatment Fidelity (interventionist): NR Acceptability: high level of comfort, rapport, supportiveness and overall helpfulness Safety: NR
6 Gross, 2017 (United States)	Kidney and kidney pancreas Tx candidates N = 63 I n = 32 43% male Age 50 (±12) C n = 31 43% male Age 50 (±12)	I: 8-weeks, group-based, combined in-person and telephone-based, mindfulness stress reduction training, delivered by a certified mindfulness-based stress reduction teacher C: 8-week, group based, combined in-person and telephone-based, weekly, structured support group, delivered by a group facilitator Assessments: - Baseline - 2 months after baseline - 6 months after baseline	Health-related quality of life (SF36) Anxiety (STAI) Depression (CES-D) Sleep quality (PSQI) Pain (SF12 pain item) Fatigue (PROMIS-Fatigue SF)	I ↑ C on MCS at 6-month follow-up I ↑ within group PCS score at 2-month follow-up I ≈ C I ↑ C at 2-month follow-up I ≈ C at 6-month follow-up I ≈ C I ≈ C I ≈ C	Enrolment: 63/388 (16%) of eligible patients Attrition: 51/63 (81%) completed assessment at 2 months after baseline (I 27/32 (84%); C 24/31 (77%)). 42/63 (67%) completed all assessments (I 22/32 (69%); C 20/31 (65%)) Fidelity (participants): attendance seven out of eight sessions; in both groups; n = 4 never attended Fidelity (interventionist): no treatment contamination found Acceptability: 90% reported continuing meditation practices, 67%–80% indicated that MBSR was helpful Safety: No intervention-related adverse events occurred

Tx, transplant; I, intervention group; C, comparator group; NR, Not reported; CBT, cognitive behavioural therapy; SF-36, short form 36 questionnaire; GHQ, general health questionnaire; PQLS, pulmonary-specific quality-of-life-scale; PSSTx, perceived social support related to transplantation; PSTx, perceived stress related to transplantation QOLI, quality of life inventory; POMS, Profile of Mood States Short-Form; MSIS, 17-item Miller Social Intimacy Scale; FEV1, Forced expiratory volume; 6MWT, six minute walk test; CLDQoL, Chronic Liver Disease Quality of Life; PQLS, pulmonary specific Quality of Life Scale; STAI, State-Trait Anxiety Inventory- State form; BDI, Beck Depression Inventory; PSS, perceived stress scale; LOT-R, life orientation test- revised; PSSC, Perceived social support scale; SDS-BQ, University of California San Diego Shortness of breath Questionnaire; COPE Inventory; POMS, Profile of Mood States-Short Form; HSCL, Hopkins Symptom Checklist-25; No of unhealthy mental health days in the past month, subjective reporting of number of days experiencing stress; depression or anxiety in the past month; CES-D, The Centre for Epidemiologic Studies Depression Scale; PSQI, The Pittsburg Sleep Quality Index; SF-12, Short-Form 12 questionnaire; MCS, Mental Composite Score of the SF-12; PCS, Physical Composite Score of the SF-12; PROMIS-Fatigue, PROMIS-Fatigue Short Form v1.



measures, their definitions, and the tools to measure them. Literature was reviewed with respect to the outcomes evaluated as well as the tools to measure them. One general recommendation was formulated.

Recommendation 3.1

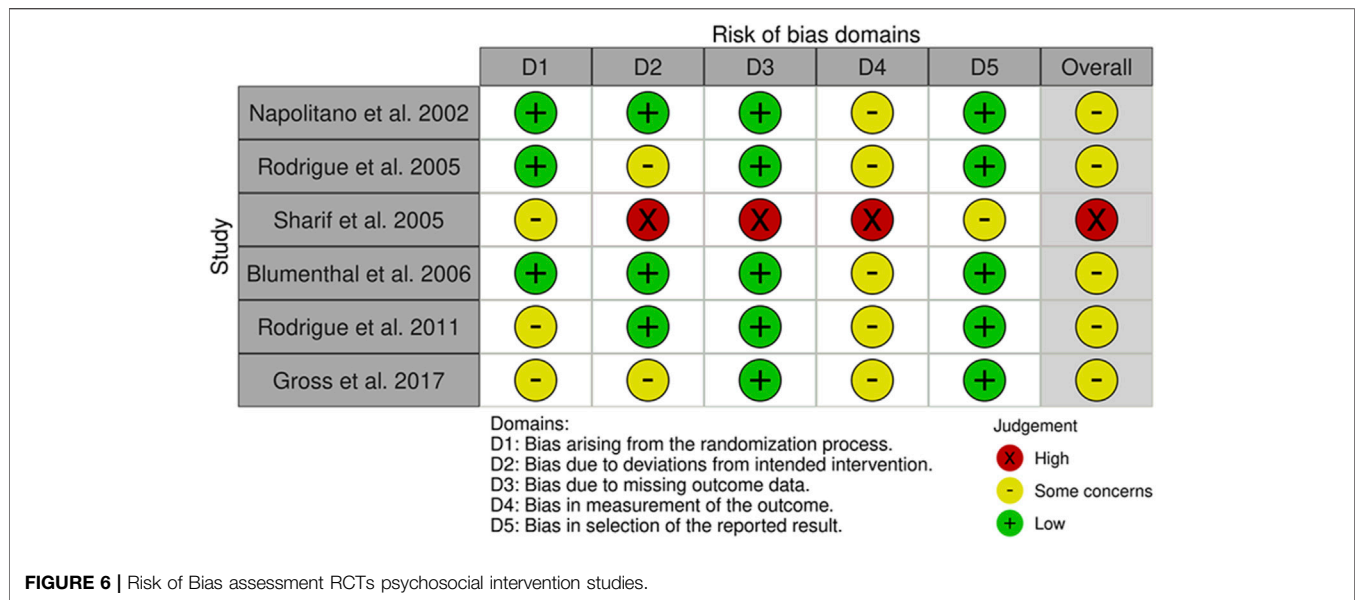
It is strongly recommended that a *core outcome measurement set* is defined for future multimodal prehabilitation studies in solid organ transplant candidates.

Quality of Evidence: not applicable.

Strength of Recommendation: Strong.

Rationale: The studies retrieved during this review varied widely with respect to the clinical and patient-reported

outcomes that were utilized, and the methods to assess them (Tables 2–4). Most exercise intervention studies included cardiorespiratory fitness including peak or maximal oxygen consumption (VO_{2peak}) and/or six-minute walking distance (6MWD)], Health-related Quality of Life (HRQoL), dyspnea, or maximal inspiratory pressures outcome measures. Nutritional intervention studies mostly monitored weight changes, infection rates, body composition and survival as either primary or secondary outcomes. The outcomes in studies that used psychosocial interventions included mostly HRQoL as well as parameters of mood, social intimacy and coping, while the use of clinical outcomes was rare. All stakeholders including solid organ transplant candidates and recipients, transplant



professionals, and researchers in the field of transplantation strongly supported that a *core outcome set* be defined to facilitate comparative studies and give impetus to the field. A *core outcomes set* refers to a minimum set of outcome measures that are critical to patients, caregivers, and health professionals for decision making [71]. Selected outcomes that have so far not been considered but do carry clinical relevance during the pre-transplant waiting time are health-related physical fitness parameters such as muscular fitness, motor fitness, body composition and (cardio)metabolic health, as well as patient-reported outcomes such as fatigue, medication adherence and lifestyle, and clinical outcomes such as duration of intensive care stay, hospitalization, (re-)admissions, complications, graft function and survival, and waitlist and post-transplant mortality.

PICO Question 4

In adult candidates for lung, liver, kidney, & heart transplantation: what is the evidence for the feasibility (enrolment, retention, acceptability, fidelity, safety) of prehabilitation?

Implementation of prehabilitation in clinical practice of solid organ transplantation should be supported by evidence of feasibility. Two systematic reviews have previously concluded that exercise prehabilitation is feasible and safe for solid organ transplant candidates [11, 12]. The review by Wallen et al was performed with focus on the feasibility outcomes enrolment, retention, acceptability, fidelity and safety [11]. One general recommendation was made.

Recommendation 4.1

It is strongly recommended that future studies on multimodal prehabilitation in solid organ transplant candidates include the specific assessment of feasibility.

Quality of Evidence: Moderate.

Strength of Recommendation: Strong.

Rationale: One study was identified that was specifically designed to assess the feasibility of delivering a psychosocial prehabilitation in solid organ transplant candidates. This study showed that a stress management and relaxation training program in liver transplant candidates was efficiently deliverable and considered acceptable and tolerable by the patients [67]. However, the enrolment rate was low, (29%) and the attrition rate was moderate (68%). Amongst the remainder of the literature, most studies reported on some aspect(s) of feasibility as a secondary outcome, mainly regarding enrolment and attrition (Tables 2–4). The feasibility measures fidelity of participants and/or interventionist and safety were less reported (Tables 2–4). For the exercise intervention studies, the enrolment rate was approximately 86%, while the average attrition rate ranged between 71% and 100% [19–25]. However, drop-outs were often due to transplant surgery. In the studies on nutritional interventions, feasibility measures were poorly reported. If reported, the enrolment rate was found to be low to moderate [26, 31, 32]. Attrition rates ranged between 62% and 98% [26, 27, 29–32]. In the psychosocial intervention studies, enrolment rates ranged between 24% and 59%, attrition rates between 69% and 88%, and acceptability of the intervention was high [33, 34, 36–38]. Only two studies reported the occurrence of adverse events [27, 65], but no serious adverse events occurred.

Overall, the consensus was that these studies do support the notion that it is feasible, acceptable and safe for adults to participate in exercise, nutritional, and psychosocial interventions during the waiting-list period (Tables 2–4). Although enrolment in studies differed significantly across studies, the overall willingness to participate in studies was found to be good and the attrition rates are adequate, and few adverse events are reported. Fidelity of participants as well as the

interventionist and acceptability of the intervention are less reported. Nonetheless, implementation of prehabilitation in a clinical practice has not been established so far. Future dedicated studies should focus on the feasibility of implementation in clinical practice by assessing factors related to potential implementation strategy effects (e.g., adoption, fidelity, reach, sustainability) and factors to inform the design or development of the implementation strategy (e.g., acceptability, adaptability, feasibility, compatibility, complexity, self-efficacy, context, costs) [72].

DISCUSSION AND FUTURE PERSPECTIVES

The newly established ESOT consensus platform has proven successful in supporting the development of evidence-based consensus recommendations for prehabilitation in candidates for solid organ transplantation. Ten recommendations were formulated for which full consensus was reached within two voting rounds. This indicated that a high degree of consensus existed amongst all stakeholders from the prehabilitation, rehabilitation and transplantation fields, including transplant recipients and their representatives, and that the recommendations are a balanced representation of current published evidence and expert opinion.

Published evidence on prehabilitation before solid organ transplantation was found to be limited and consisted of studies addressing unimodal prehabilitation interventions with heterogeneous design, methodology and relatively small sample sizes. Nevertheless, by consensus and expert opinion, the available evidence on effects of prehabilitation on physical functioning, nutritional status, and psychosocial wellbeing and the evidence on safety of prehabilitation interventions was felt sufficiently strong to recommend that multimodal, patient-tailored prehabilitation should be offered as standard of care to patients awaiting solid organ transplantation. Specific recommendations included exercise-based intervention as well as psychological and stress management support for all solid organ transplant candidates, nutritional intervention for those who are over- or underweight, and probiotic supplementation for candidates for liver transplantation.

Because of the shortage in clinical evidence, however, particularly strong recommendations were formulated regarding the urgent need for high quality, but not exclusively, randomized controlled trials and implementation research studies that address the feasibility and effectiveness of pre-transplant multimodal prehabilitation. Two RCTs on multimodal prehabilitation interventions in kidney transplant candidates are currently underway: the FRAIL-MAR-study (NCT04701398) [73] and the PreCareTx-study (NCT05489432).

In addition, it was strongly recommended that priority should be given to the definition and consistent use of a Core Outcome Set to be measured by all future trials modalities, timing, duration and delivery modes of an optimal prehabilitation program.

From the in-person public discussions during the Consensus ESOT Conference, additional constructive perspectives emerged. It was advocated that clinical guidelines should be broadly applicable to

transplant candidates irrespective of organ type, while leaving room for organ-specific recommendations, such as probiotics for liver transplant candidates. It was noted that intoxication-related interventions are not included in the recommendations, as intoxication (i.e., tobacco smoking or alcohol abuse) is typically addressed prior to patients joining the waitlist. The suggestion was made to formulate recommendations regarding pre-transplant peer support, however, such was considered premature as no evidence base could be found in the literature review. Lastly, the consideration was made that the designing of future studies or the future revisiting of the new guidelines may benefit of being informed by the prehabilitation literature in the broader field of surgery. However, unlike elective surgery, the waiting time is often unpredictably long while physical and mental condition may deteriorate due to the underlying disease. Therefore, prehabilitation should be offered throughout the waiting period from the moment of listing until transplantation.

These new evidence-based recommendations on prehabilitation serve to support best clinical practice in solid organ transplantation and help identify priorities for future research, thus optimizing patient health and post-transplant clinical outcome. The final recommendations will be included in the ESOT guidelines for transplant management, and under the auspices of the ESOT consensus development platform, will undergo continuous review and updating as new evidence becomes available.

DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/**Supplementary Material**, further inquiries can be directed to the corresponding author.

AUTHOR CONTRIBUTIONS

Involved in the conception or design of the work: CA, SS, EC, YO, JK, TJ-F, SM, EK, MS, PF, DM, and SG. Literature screen and review: CA, SS, EC, YO, JK, TJ-F, SM, EK, MS, CM, FD, DM, and SG. Drafted the article: CA, SS, EC, YO, CM, DM, and SG. Critically revised the article: JK, TJ-F, SM, EK, MS, FD, PF, JG, AM, PG, MP, VL-L, CW, DK, DM, and SG. Finally approved the version to be published: CA, SS, EC, YO, JK, TJ-F, SM, EK, MS, CM, FD, PF, JG, AM, PG, MP, VL-L, CW, DK, DM, and SG. All authors contributed to the article and approved the submitted version.

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CONFLICT OF INTEREST

DM is a senior researcher of The Research Foundation- Flanders.

The remaining authors declare that the research was conducted in the absence of any commercial or financial

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontierspartnerships.org/articles/10.3389/ti.2023.11564/full#supplementary-material>

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