scientific reports



OPEN The impact of preoperative oral nutrition supplementation on outcomes in patients undergoing gastrointestinal surgery for cancer in lowand middle-income countries: a systematic review and meta-analysis

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Malnutrition is an independent predictor for postoperative complications in low- and middle-income countries (LMICs). We systematically reviewed evidence on the impact of preoperative oral nutrition supplementation (ONS) on patients undergoing gastrointestinal cancer surgery in LMICs. We searched EMBASE, Cochrane Library, Web of Science, Scopus, WHO Global Index Medicus, SciELO, Latin American and Caribbean Health Sciences Literature (LILACS) databases from inception to March 21, 2022 for randomised controlled trials evaluating preoperative ONS in gastrointestinal cancer within LMICs. We evaluated the impact of ONS on all postoperative outcomes using random-effects metaanalysis. Seven studies reported on 891 patients (446 ONS group, 445 control group) undergoing surgery for gastrointestinal cancer. Preoperative ONS reduced all cause postoperative surgical complications (risk ratio (RR) 0.53, 95% CI 0.46–0.60, P < 0.001, $l^2 = 0\%$, n = 891), infection (0.52, 0.40– 0.67, P = 0.008, $l^2 = 0\%$, n = 570) and all-cause mortality (0.35, 0.26–0.47, P = 0.014, $l^2 = 0\%$, n = 588). Despite heterogeneous populations and baseline rates, absolute risk ratio (ARR) was reduced for all

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cause (pooled effect -0.14, -0.22 to -0.06, P = 0.006; number needed to treat (NNT) 7) and infectious complications (-0.13, -0.22 to -0.06, P < 0.001; NNT 8). Preoperative nutrition in patients undergoing gastrointestinal cancer surgery in LMICs demonstrated consistently strong and robust treatment effects across measured outcomes. However additional higher quality research, with particular focus within African populations, are urgently required.

Malnutrition is a major public health issue in low- and middle-income countries (LMICs) and forms part of the United Nations 2030 Agenda for Sustainable Development¹. The predominant focus in LMICs has been child nutrition, yet as many as two-thirds of hospitalised adult patients are malnourished in this setting². Malnutrition is associated with higher postoperative mortality and morbidity, including longer length of in-patient stay and increased healthcare-associated costs²⁻⁴. Furthermore, preoperative nutrition has been identified as an area of high research priority in LMICs⁵.

Provision of safe and equitable surgical care is becoming increasingly recognised as an essential part of cancer care and population health⁶. In the majority of solid tumours, surgery provides the best chance of cure, particularly where chemotherapy and radiotherapy are unavailable⁷. Oral nutritional supplementation (ONS) provided at the time of surgery in LMICs could provide a low-cost and sustainable intervention, requiring minimal specialist training and equipment to administer. Several reviews and meta-analyses have demonstrated the beneficial effects of preoperative nutrition on surgical site infection, peri-operative complication rate and length of stay⁸⁻¹⁰. However, data is lacking from a systematic review of the evidence exploring the impact of standard oral nutritional supplementation an LMIC setting.

This systematic review and meta-analysis investigated the effect of preoperative oral nutrition on postoperative outcomes for patients undergoing gastrointestinal surgery for cancer in LMICs.

Materials and methods

Search strategy and selection criteria. The systematic review protocol was registered prospectively with the PROSPERO database (CRD42019125161)¹¹. A systematic search of the EMBASE, Cochrane Library, Web of Science, Scopus, WHO Global Index Medicus, SciELO, Latin American and Caribbean Health Sciences Literature (LILACS) databases, together with a grey literature search using Google Scholar, was performed in accordance with the PRISMA guidelines¹².

Search terms relating to preoperative oral nutritional intervention in patients undergoing surgery for solid tumours were combined with LMIC filters as specified by the Cochrane Collaboration¹³. The following exploded Medical Subject Headings (MeSH) were used: "surgery", "cancer", "malignancy", "nutrition", "diet" combined with "postoperative outcomes" (Supplementary Material). Databases were searched from inception, with no limits on publication year or language placed. The reference list of all studies that met the inclusion criteria and review articles were searched manually for additional studies. The trial registry clinicaltrials.gov was searched to identify any unpublished studies. The final literature search was performed on 21st March 2022. Non-randomised, retrospective, review articles, letters to the editor, case reports and conference abstracts with no access to the entire study were excluded.

All studies identified were screened independently by two reviewers from a pool of seven (SB, UQ, TMD, CML, MM, EY, SS) using the online systematic review tool Covidence¹⁴. All disagreements were adjudicated by a third reviewer (SK).

Randomised controlled trials reporting at least one clinical outcome in an LMIC population based on the World Bank classification at the time of study publication¹⁵ were included, as described previously¹⁶. Studies were required to be in patients aged 18 years or above undergoing surgery for gastrointestinal cancer, defined as any procedure requiring a skin incision under regional or general anaesthesia. The intervention required the use of an oral nutritional supplement (ONS) containing macronutrients (fat, carbohydrate and protein) with or without micronutrients (vitamins and minerals). The control arm was patients receiving routine care with no additional dietary supplementation. Therefore, the only difference between the intervention and control groups was the additional preoperative intake of ONS.

Studies with a nutritional intervention using single nutrient substrates, complementary food substances, probiotic formulas or as part of a multimodal preoperative intervention (such as an enhanced recovery programme) and those delivered by enteral tubes or parenteral routes were excluded. Those studies that only reported on postoperative administration of a nutritional intervention were also excluded. Additionally, studies that met the inclusion criteria but did not report data separately for malignant and non-malignant surgery, or between patients within a high-income and LMIC setting were also excluded if attempts to obtain the relevant data failed. A summary of inclusion and exclusion criteria is provided in Supplementary Table S1.

Data extraction and statistical analysis. Non-English articles were translated by medically qualified individuals where appropriate. Data were retrieved from published articles using a standardised data extraction form for all included studies, including publication details, study design, country, participant number, proportion of malnourished participants, cancer type, surgical procedures performed, participant age, oral nutritional intervention used, follow-up period, 30-day complication rate, all-cause mortality and length of stay. Attempts were made to contact study authors if any data were unclear within the published manuscript or study protocol. Assessment of methodological quality was performed for all included studies using the Cochrane Risk-of-Bias tool¹⁷. Publication bias was assessed through funnel plot symmetry and statistical analysis using Egger's test for each outcome.

All binary outcome measures were summarised as risk ratios (RRs) with 95% confidence intervals (CIs), with individual study weights calculated for pooled analysis. Risk ratios (RR) were reported in accordance with the Cochrane Collaboration to avoid overestimation of any potential treatment effect¹⁸. For individual trials with zero event data in one or more groups, a continuity correction of 0.5 was performed to provide a more conservative estimate of effect size¹⁹. The presence of statistical heterogeneity was expected, due to the in-between study variability in cancer type, geographical setting, malnutrition rate and oral nutrition provided. Therefore, pooled data analyses were performed using the Mantel–Haenszel random-effects model using the R meta package (v3.6.3).

Absolute Risk Reduction (ARR) was used to estimate population and baseline rate heterogeneity for each outcome, excluding those including zero event data¹⁹, with pooled estimates calculated as previously stated. The number needed to treat to benefit (NNT), estimating the number of patients that need to be treated in order to have an impact on one person, was defined as the reciprocal of the absolute risk reduction.

Influence analyses, using the leave-one-out method, were performed to determine robustness of pooled effect estimates^{19,20} Heterogeneity was assessed using the I^2 statistic and defined as low, moderate or high with the corresponding upper limits of 25%, 50% and 75%, respectively¹⁷.

Role of funding source. The study sponsors had no role in study design, data collection, data analysis, data interpretation, or writing of the report. All authors had full access to all the data in the study and had final responsibility for the decision to submit for publication.

Patient and public involvement statement. Patient representatives for the NIHR Global Health Research Unit on Global Surgery, from both the UK and Rwanda, guided development of the research question.

Results

Literature search. The systematic search yielded 6615 studies. After the removal of 259 duplicates, 6146 articles were excluded by publication type, or on the basis of title or abstract. Of the remaining 210 articles, 203 did not meet inclusion criteria, with a final seven articles included in the meta-analysis (Fig. 1)²¹⁻²⁷. The reasons for article exclusion are stated within Fig. 1. A summary of study and patient characteristics are provided in Table 1 and Supplementary Table S2. Six studies originated from China²¹⁻²⁶ and one from India²⁷. All studies included patients undergoing surgery for gastrointestinal malignancy, with four²²⁻²⁵ investigating preoperative oral nutrition in gastric cancer surgery only. A total of 891 patients (446 oral nutrition group, 445 control group) were included within the meta-analysis. Six studies reported the recruitment of a full cohort with malnutrition as measured by validated tools (Table 1), with one study not reporting nutritional status²⁶.

Included study design. A number of oral nutritional formulations were used, each compared to a standard diet control. Nutrison liquid (Nutricia^{*}), commonly given by feeding enteral tube in high-income countries, was used as an oral supplement in three studies²²⁻²⁴. Treatment regimen and duration varied between studies, with nutrition commenced for at least five days preoperatively in four studies^{21,24,25,27}. Nutritional supplementation characteristics are summarised in Supplementary Table S1.

Patient follow-up to 30 days occurred in all studies, with the majority evaluating postoperative complications as their primary outcome. Some studies additionally reported nutrition status and serum biomarkers as outcomes. Length of stay was not sufficiently reported in three studies for meta-analysis inclusion^{21,27}. A summary pooled estimates across all measured outcomes can be found in Supplementary Table S3.

Outcomes. All studies provided incident rates for postoperative complications at 30 days. The pooled event rate was 17.9% (80/446) in the oral nutrition group compared with 33.9% (151/445) in the control arm. The pooled RR for complications after preoperative treatment with oral nutrition was 0.53 (95% CI 0.46 to 0.60, P < 0.001, $I^2 = 0\%$; Fig. 2a). This effect persisted when including only studies which recruited malnourished patients (RR 0.53; 95% CI 0.41 to to 0.68, P = 0.009, $I^2 = 0\%$; Fig. 2b). The type of nutrition intervention did not modify the overall effect, with a pooled RR of 0.54 (95% CI 0.40 to 0.73, P = 0.007, $I^2 = 0\%$) following exclusion of studies using Nutrison^{*} liquid (Supplementary Fig. S1). Influence and sensitivity analysis demonstrated the persistence in overall effect for the nutritional supplement group (Supplementary Figs. S2 and S3). The intervention had a consistently strong positive effect despite baseline rate heterogeneity across included studies (ARR range -0.22 to -0.06, pooled effect -0.14, 95% CI -0.22 to -0.06; Fig. 2c), with a corresponding number needed to treat (NNT) of 7 (95% CI 5 to 17).

Three studies reported incidence of infectious complications (Fig. 3a)^{21,22,27}. The pooled event rates for infectious complications were 14.3% (41/286) in the nutrition group and 27.8% (79/284) in the control arm. The pooled RR for infectious complications was 0.52 (95% CI 0.40 to 0.67, P = 0.008, $I^2 = 0\%$; Fig. 3a). The intervention had a consistently strong positive effect despite baseline rate heterogeneity across included studies (ARR range -0.20 to -0.07, pooled effect -0.13, 95% CI -0.22 to -0.06, P < 0.001; Fig. 3b), with a corresponding NNT of 8 (95% CI 5 to 14). Meanwhile the incidence of surgical site infection (SSI) was reported in three studies^{21,22,27}, however none reported the criteria used to diagnose SSI. There were no SSI events up to 30 days postoperatively in one study²². SSI rates were 5.9% (17/286) in the nutrition group and 10.2% (29/284) in the control arm, with the pooled RR for SSI 0.59 (95% CI 0.33 to 1.04, P = 0.058, $I^2 = 0\%$; Fig. 4a).

Three studies reported the incidence of non-infectious complications^{21,22,27}, with one study reporting no events at 30 days²². Pooled event rates were 3.1% (9/286) in the nutrition group and 5.3% (15/284) in the control arm. The pooled RR for non-infectious complications were 0.61 (95% CI 0.42 to 0.88, P = 0.029, I^2 = 0%; Fig. 4b). Three studies reported on 30-day mortality^{21,26,27}, with the pooled event rate 2.0% (6/295) in the nutrition group





	Year	Country	Cancer type(s)	Patient number Intervention/Control	Patients malnourished (%)	Screening tool used
Wu et al ²¹	2006	China	Gastric, colon and rectal cancer	235/233	100	SGA
Ding et al ²²	2009	China	Gastric cancer	21/21	100	NRS-2002
Zheng et al ²³	2010	China	Gastric cancer	18/18	100	NRS-2002
Kharbuja et al ²⁴	2013	China	Gastric cancer	92/93	100	NRS-2002
Chen et al ²⁶	2013	China	Rectal cancer	30/30	ns	ns
Zhou et al ²⁵	2016	China	Gastric cancer	20/20	100	NRS-2002
Sagar et al ²⁷	2019	India	Oesophageal and gastric cancer	30/30	100	SGA

Table 1. Summary of included randomised control trials. SGA Subjective Global Assessment, NRS NutritionalRisk Screening, ns not stated.

and 5.8% (17/293) in the control arm. The pooled RR for mortality was 0.37 (95% CI 0.18 to 0.76, P = 0.027, I^2 = 0%; Fig. 4c).

Assessment of bias. Publication bias was demonstrated to be low across all measured outcomes. The distribution of RR was evenly distributed across the funnel plot, with no significant outliers (Supplementary

All complications

a

	Nutritio	n	Control			Risk		
Author	Events	Total	Events	Total	i	Ratio	(95% CI)	Weight
Wu et al Ding et al Zheng et al Kharbuja et al Chen et al Zhou et al Sagar et al	43 1 21 5 3 6	235 21 18 92 30 20 30	78 2 43 6 6 14	233 21 - 18 - 93 30 20 30		0.55 0.50 0.50 0.49 0.83 0.50 0.43	$\begin{matrix} [0.39; 0.76] \\ [0.05; 5.10] \\ [0.05; 5.04] \\ [0.32; 0.76] \\ [0.28; 2.44] \\ [0.14; 1.73] \\ [0.19; 0.96] \end{matrix}$	49.2% 1.2% 1.2% 29.8% 5.4% 4.0% 9.2%
Overall Heterogeneity: $I^2 = 0$ Test for overall effect:	80 9% : p < 0.00	446	151	445	0.1 0.5 1 2 10	0.53	[0.46; 0.60]	100.0%

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b

	Nutrition		Control			Bisk			
Author	Events	Total	Events	Total		Ratio	(95% CI)	Weight	
Wu et al Ding et al Sagar et al	43 1 6	235 21 30	78 2 14	233 21 30		0.55 0.50 0.43	[0.39; 0.76] [0.05; 5.10] [0.19; 0.96]	84.2% 1.7% 14.1%	
Overall Heterogeneity: <i>I</i> ² Test for overall eff	50 = 0% ect: p = 0.00	286 9	94	284	0.1 0.5 1 2 10	0.53	[0.41; 0.68]	100.0%	

Favours nutrition Favours control

Favours control

6	Nutritio	n	Control			Absolute		
Author	Events	Total	Events	Total		Risk Ratio	(95% CI)	Weight
Wu et al Ding et al Zheng et al Kharbuja et al Chen et al Zhou et al Saqar et al	43 1 21 5 3 6	235 21 18 92 30 20 30	78 2 43 6 6 14	233 21 18 93 30 20 30 -		-0.15 -0.05 -0.06 -0.23 -0.03 -0.15 -0.27	[-0.23; -0.07] [-0.20; 0.11] [-0.24; 0.12] [-0.37; -0.10] [-0.23; 0.16] [-0.40; 0.10] [-0.50; -0.04]	27.5% 15.0% 12.4% 17.9% 11.0% 7.4% 8.7%
Overall Heterogeneity: $I^2 = 8$ Test for overall effect:	80 3% : p = 0.00	446	151	445	-0.4 -0.2 0 0.2 0.4	- 0.14	[-0.22; -0.06]	100.0%

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Figure 2. Random-effects meta-analysis of the effects of preoperative oral nutrition on postoperative complications (**a**), when nutritional support was provided for at least 5 days pre-operatively (**b**), and risk difference (**c**) in patients undergoing surgery for gastrointestinal cancer.

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Fig. S4). The risk of bias for all included studies is summarised in Supplementary Fig. S5. Incomplete outcome data had a low risk of bias in all studies, with adequate sequence generation in five studies^{21,22,24,26,27}. However the majority of other domains contained unclear or high risk, particularly for allocation concealment and outcome assessment.

Discussion

This meta-analysis has demonstrated that the risks of infection, complications and all-cause mortality after surgery for gastrointestinal cancer in a LMIC setting were reduced in patients receiving preoperative nutrition. The intervention had a consistently strong effect on complication rates across each study population despite heterogeneity in baseline rates. However, the intervention did not impact upon SSI, with insufficient data available to assess length of stay, hospital costs, and return to work or household activity.

The analysis included patient populations from two countries, undergoing operations for gastrointestinal cancer. Statistical heterogeneity was found to be low for all outcomes measured and the risk of publication bias was also low. Overall treatment effects were robust during sensitivity analysis and due to methodology are likely to be conservative estimates. However, interventions were predominantly performed in China for patients undergoing surgery for gastric cancer and evidence of methodological bias was demonstrated.

Malnutrition is a major public health issue in LMICs and forms part of the United Nations 2030 Agenda for Sustainable Development¹, reducing a patient's ability to compensate for stressful events, such as major

а

b

	Nutritio	n	Control						Rick		
Author	Events	Total	Events	Total					Ratio	(95% CI)	Weight
Wu et al Ding et al Sagar et al	35 1 5	235 21 30	65 2 12	233 21 — 30		-			0.53 0.50 0.42	[0.37; 0.77] [0.05; 5.10] [0.17; 1.04]	84.1% 2.1% 13.7%
Overall Heterogeneity: <i>I</i> ² = 0' Test for overall effect:	41 $\% p = 0.008$	286	79	284	0.1	0.5 1	2	10	0.52	[0.40; 0.67]	100.0%
				Favour	rs nutri	tion	Favou	ırs co	ntrol		

Nut	rition	Control		Risk		
Author Eve	ents Total	Events	Total	Ratio	(95% CI)	Weight
Wu et al 35 Ding et al 1 Sagar et al 5	235 21 30	65 2 12	233 21 30	-0.13 -0.05 -0.23	[-0.20; -0.06] [-0.20; 0.11] [-0.45; -0.01]	82.1% 7.4% 10.5%
Overall 41 Heterogeneity: <i>I</i> Test for overall e	286 ${}^{2} = 0\%$ effect: $p < 0$.	79 001	284 -0.2 0 0.2 Favours nutrition Favours of	- 0.13	[-0.20; -0.07]	100.0%

Figure 3. Random-effects meta-analysis of the effects of preoperative oral nutrition on infectious complications (a), and risk difference (b) in patients undergoing surgery for gastrointestinal cancer.

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surgery²⁸. While associated with poorer outcomes, malnutrition is potentially reversible²⁹. The European Society for Parenteral and Enteral Nutrition (ESPEN) guidelines suggest nutritional support should be initiated without delay in patients undergoing surgery if oral intake reduction is expected 7 days perioperatively³⁰, however the effectiveness of nutritional intervention in LMICs is uncertain.

Previously the ability of nutrition interventions to reduce infectious complications and length of hospital stay in a global population has been demonstrated¹⁰, however inclusion of high-income populations and parenteral routes limits generalizability to LMIC settings. A recent meta-analysis including only studies conducted in East Asia demonstrated no benefit of preoperative oral nutrition for postoperative complications⁹, however the sole inclusion of gastric cancer patients undergoing surgery, high weighting towards high-income country settings and significant study heterogeneity (58%) may explain differences with our findings.

Baseline rates for measured outcomes differed across studies, similar to variation demonstrated globally in large population cohorts across LMICs^{31,32}. However, a consistently strong positive treatment effect was shown across measured outcomes, demonstrated by similar risk ratios and small confidence intervals. This suggests preoperative oral nutrition confers a positive effect independent of baseline complication rate and our findings are applicable across LMIC settings.

The method of administration was not always obvious and tube feeding may have been used in three studies²²⁻²⁴, yet sub-group analysis found treatment effects persisted following their exclusion. Some studies failed to report the formal criteria used to classify postoperative complications and SSI. Furthermore, disease stage, nutrition dose variation and potential unmeasured confounders will have introduced elements of clinical heterogeneity. However, the use of random-effects models, consistent treatment effects and low statistical heterogeneity overall supports our conclusions. Only four studies provided at least five days preoperative nutrition, in keeping with current guideline recommendations^{33,34}. Therefore, our results may underestimate the overall effect of preoperative nutrition in LMICs.

Surgical site infection rates were low within included studies (overall rate 8.1%; range 0 – 12.6%), which suggests SSI may be under-reported³¹. The absence of definitive diagnostic criteria, such as those stated by the Centre for Disease Control and Prevention³⁵, may explain this variation and the null effect of nutrition on SSI rates. In contrast, infectious complications reduced following preoperative nutrition, similar to another recent meta-analysis of immune modulating nutrition in high-income settings⁸.

Some limitations within our analysis exist. Cancer-focused studies commonly report longer-term survival, particularly at one and five years^{36,37}, and the impact of preoperative oral nutrition on these outcomes remains unknown, with included studies only reporting data on short-term outcomes. However, the demonstrated absolute risk reduction in mortality may also influence longer-term survival in patients undergoing surgery for localised, potentially curative disease. Furthermore, the reduction in postoperative complications is likely to reduce delays to adjuvant treatment, which have been associated with worse survival^{38–40} and unfavourable oncological outcomes⁴¹ in a wide range of cancers.

Secondly, particular patient groups are under-represented within the meta-analysis. The effectiveness of interventions remains uncertain in some globally common malignancies, for example gynaecological and oral cancer⁴², and in a broader range of settings across Africa and the Asian subcontinent. More conclusive statements on the effectiveness of preoperative nutrition across LMICs is limited by the majority of studies conducted

a

Surgical site infection

	Nutritio	n	Control			Risk		
Author	Events	Total	Events	Total		Ratio	(95% CI)	Weight
Wu et al Ding et al Sagar et al	15.0 0.5 2.0	235 21 30	27.0 0.5 2.0	233 21 — 30		0.55 —1.00 1.00	[0.30; 1.01] [0.02; 48.09] [0.15; 6.64]	88.8% 2.2% 9.1%
Overall Heterogeneity: $I^2 = 0$ Test for overall effect	17.5)% : <i>p</i> = 0.05	286 8	29.5	284	0.1 0.51 2 10	0.59	[0.33; 1.04]	100.0%
				Favours	s nutrition Favours co	ontrol		

b

Non-infectious complications

	Nutritio	n	Control			Risk		
Author	Events	Total	Events	Total		Ratio	(95% CI)	Weight
Wu et al Ding et al Sagar et al	8.0 0.5 1.0	235 21 30	13.0 0.5 2.0	233 21 — 30		0.61 —1.00 0.50	[0.26; 1.44] [0.02; 48.09] [0.05; 5.22]	84.4% 4.2% 11.4%
Overall Heterogeneity: $I^2 = 0$ Test for overall effect:	9.5 % p = 0.029	286 9	15.5	284	0.1 0.51 2 10	0.61	[0.42; 0.88]	100.0%

Favours nutrition Favours control

С

Mortality

	Nutrition		Control			Risk			
Author	Events	Total	Events	Total		Ratio	(95% CI)	Weight	
Wu et al Chen et al Sagar et al	5.0 0.5 1.0	235 30 30	14.0 0.5 3.0	233 30 — 30		0.35 —1.00 0.33	[0.13; 0.97] [0.02; 48.77] [0.04; 3.03]	78.5% 5.2% 16.3%	
Overall Heterogeneity: $I^2 = 0$ Test for overall effect	6.5 % : p = 0.02	295	17.5	293	0.1 0.51 2 10	0.37	[0.18; 0.76]	100.0%	

Favours nutrition Favours control

Figure 4. Random-effects meta-analysis of the effects of preoperative oral nutrition on surgical site infection (**a**), non-infectious complications (**b**), and mortality (**C**) in patients undergoing surgery for gastrointestinal cancer.

in China. Furthermore, despite being LMICs, advanced medical care is available in many parts of China and India. It is unclear if the circumstances of these included studies are truly representative of the challenges other countries might face with implementation, and the populations they will be predominantly treating. Lastly, the majority of interventions were commercially sourced, with no cost-effectiveness data to support this strategy within LMIC settings. It remains to be demonstrated whether nutritionally balanced, locally sourced low-cost supplements would be as effective.

If these research gaps are addressed, preoperative oral nutrition is likely to form part of future global surgical guidelines as a simple measure that can improve outcomes after surgery for cancer. Planned trials should particularly focus on determining the impact of oral nutrition in Africa and the Asian subcontinent, with at least one expected in the near future⁴³.

Conclusion

This meta-analysis provides substantial evidence that preoperative oral nutrition in patients undergoing surgery for gastrointestinal cancer has a significant impact on postoperative complications and all-cause mortality. Treatment effects remained consistent despite variation in baseline complication rates and suggest generalisability across income strata. However, high quality randomised control trials across a wider LMIC surgical population are required to validate our findings based on current low to moderate quality of evidence.

Data availability

All data included within the meta-analysis is freely available within the public domain as all studies are published. The search strategy is available in the Supplementary material, and any additional data are available on reasonable request to the corresponding author.

Received: 23 March 2022; Accepted: 11 July 2022 Published online: 21 July 2022

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Acknowledgements

Funding for this study were provided by the UK Medical Research Council (Grant reference MR/S014527/1). We thank Dr. Kai Nie and Dr. James Ng for providing English translations for included articles originally published in Chinese.

Author contributions

S.R.K., S.B., S.S. and E.M.H. conceived the idea for the study and contributed to study design. S.R.K., A.U.Q., T.M.D., C.M.L., M.M. and E.Y. performed data extraction and quality assessments. S.R.K. performed the metaanalysis. S.R.K. wrote the first manuscript draft and all authors made substantial contributions, including interpretations of findings and critical revision of the manuscript. All authors approved the final version of the manuscript prior to submission.

Funding

This work was supported by the UK Medical Research Council MRC (Grant reference MR/S014527/1); ET is supported by a Cancer Research UK fellowship (C31250/A22804).

Competing interests

The authors declare no competing interests.

Additional information

Supplementary Information The online version contains supplementary material available at https://doi.org/ 10.1038/s41598-022-16460-4.

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