

Assessment of nutritional status and nutrition impact symptoms in patients undergoing resection for upper gastrointestinal cancer: Results from the multi-centre nourish point prevalence study

The Nourish Point Prevalence Study Group; Deftereos, Irene; Yeung, Justin M.C.; Arslan, Janan; Carter, Vanessa M.; Isenring, Elizabeth; Kiss, Nicole

Published in:
Nutrients

DOI:
[10.3390/nu13103349](https://doi.org/10.3390/nu13103349)

Licence:
CC BY

[Link to output in Bond University research repository.](#)

Recommended citation(APA):

The Nourish Point Prevalence Study Group, Deftereos, I., Yeung, J. M. C., Arslan, J., Carter, V. M., Isenring, E., & Kiss, N. (2021). Assessment of nutritional status and nutrition impact symptoms in patients undergoing resection for upper gastrointestinal cancer: Results from the multi-centre nourish point prevalence study. *Nutrients*, 13(10), [3349]. <https://doi.org/10.3390/nu13103349>

General rights

Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

For more information, or if you believe that this document breaches copyright, please contact the Bond University research repository coordinator.

Article

Assessment of Nutritional Status and Nutrition Impact Symptoms in Patients Undergoing Resection for Upper Gastrointestinal Cancer: Results from the Multi-Centre NOURISH Point Prevalence Study

Irene Deftereos ^{1,2,*} , Justin M. C. Yeung ^{1,3,4} , Janan Arslan ¹ , Vanessa M. Carter ², Elizabeth Isenring ^{5,6}, Nicole Kiss ^{7,8}  and on behalf of The NOURISH Point Prevalence Study Group [†]

- ¹ Department of Surgery, Western Precinct, Melbourne Medical School, The University of Melbourne, St Albans, VIC 3021, Australia; justin.yeung@unimelb.edu.au (J.M.C.Y.); janan.arslan@unimelb.edu.au (J.A.)
- ² Department of Nutrition and Dietetics, Western Health, Footscray, VIC 3011, Australia; Vanessa.carter@wh.org.au
- ³ Department of Colorectal Surgery, Western Health, Footscray, VIC 3011, Australia
- ⁴ Western Health Chronic Disease Alliance, Western Health, Footscray, VIC 3011, Australia
- ⁵ Faculty of Health Sciences and Medicine, Bond University, Robina, QLD 4226, Australia; Lisenrin@bond.edu.au
- ⁶ Department of Nutrition and Dietetics, Princess Alexandra Hospital, Brisbane, QLD 4102, Australia
- ⁷ Institute for Physical Activity and Nutrition, Deakin University, Geelong, VIC 3220, Australia; nicole.kiss@deakin.edu.au
- ⁸ Allied Health Research, Peter MacCallum Cancer Centre, Melbourne, VIC 3000, Australia
- * Correspondence: irene.deftereos@unimelb.edu.au; Tel.: +61-3-8395-8116
- † Membership of the NOURISH Point Prevalence Study Group is provided in the Acknowledgments.



Citation: Deftereos, I.; Yeung, J.M.C.; Arslan, J.; Carter, V.M.; Isenring, E.; Kiss, N.; on behalf of The NOURISH Point Prevalence Study Group.

Assessment of Nutritional Status and Nutrition Impact Symptoms in Patients Undergoing Resection for Upper Gastrointestinal Cancer: Results from the Multi-Centre NOURISH Point Prevalence Study. *Nutrients* **2021**, *13*, 3349. <https://doi.org/10.3390/nu13103349>

Academic Editor: Riccardo Caccialanza

Received: 23 August 2021

Accepted: 21 September 2021

Published: 24 September 2021

Publisher's Note: MDPI stays neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Copyright: © 2021 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<https://creativecommons.org/licenses/by/4.0/>).

Abstract: Background: Identification and treatment of malnutrition are essential in upper gastrointestinal (UGI) cancer. However, there is limited understanding of the nutritional status of UGI cancer patients at the time of curative surgery. This prospective point prevalence study involving 27 Australian tertiary hospitals investigated nutritional status at the time of curative UGI cancer resection, as well as presence of preoperative nutrition impact symptoms, and associations with length of stay (LOS) and surgical complications. Methods: Subjective global assessment, hand grip strength (HGS) and weight were performed within 7 days of admission. Data on preoperative weight changes, nutrition impact symptoms, and dietary intake were collected using a purpose-built data collection tool. Surgical LOS and complications were also recorded. Multivariate regression models were developed for nutritional status, unintentional weight loss, LOS and complications. Results: This study included 200 patients undergoing oesophageal, gastric and pancreatic surgery. Malnutrition prevalence was 42% (95% confidence interval (CI) 35%, 49%), 49% lost $\geq 5\%$ weight in 6 months, and 47% of those who completed HGS assessment had low muscle strength with no differences between surgical procedures ($p = 0.864$, $p = 0.943$, $p = 0.075$, respectively). The overall prevalence of reporting at least one preoperative nutrition impact symptom was 55%, with poor appetite (37%) and early satiety (23%) the most frequently reported. Age (odds ratio (OR) 4.1, 95% CI 1.5, 11.5, $p = 0.008$), unintentional weight loss of $\geq 5\%$ in 6 months (OR 28.7, 95% CI 10.5, 78.6, $p < 0.001$), vomiting (OR 17.1, 95% CI 1.4, 207.8, 0.025), reduced food intake lasting 2–4 weeks (OR 7.4, 95% CI 1.3, 43.5, $p = 0.026$) and ≥ 1 month (OR 7.7, 95% CI 2.7, 22.0, $p < 0.001$) were independently associated with preoperative malnutrition. Factors independently associated with unintentional weight loss were poor appetite (OR 3.7, 95% CI 1.6, 8.4, $p = 0.002$) and degree of solid food reduction of $< 75\%$ (OR 3.3, 95% CI 1.2, 9.2, $p = 0.02$) and $< 50\%$ (OR 4.9, 95% CI 1.5, 15.6, $p = 0.008$) of usual intake. Malnutrition (regression coefficient 3.6, 95% CI 0.1, 7.2, $p = 0.048$) and unintentional weight loss (regression coefficient 4.1, 95% CI 0.5, 7.6, $p = 0.026$) were independently associated with LOS, but no associations were found for complications. Conclusions: Despite increasing recognition of the importance of preoperative nutritional intervention, a high proportion of patients present with malnutrition or clinically significant weight loss, which are associated with increased LOS. Factors associated with malnutrition and weight loss should be incorporated into routine preoperative

screening. Further investigation is required of current practice for dietetics interventions received prior to UGI surgery and if this mitigates the impact on clinical outcomes.

Keywords: gastrointestinal cancer; gastrointestinal surgery; subjective global assessment; malnutrition; nutrition impact symptoms

1. Introduction

Upper gastrointestinal (UGI) cancers carry some of the highest mortality rates of all cancers worldwide [1]. Although there have been advances in treatments and associated survival rates, surgery for UGI cancer carries a high morbidity rate and is considered a major procedure [2–4]. Optimisation of nutritional status is recognised as a key component of perioperative care in major abdominal surgery, as malnutrition and unintentional weight loss can increase the risk of surgical complications, length of stay (LOS) and mortality [5,6].

Patients with UGI cancer are one of the highest-risk groups for malnutrition [7]. Although oncological care pathways advocate for the dietitian to be an essential member of the multi-disciplinary treatment team [8], often there is a lack of resourcing to support adequate nutritional screening and intervention prior to gastrointestinal cancer surgery [9,10]. Furthermore, there is a lack of high-quality evidence regarding optimal methods of nutrition support prior to UGI cancer surgery and in neoadjuvant therapy [11], and UGI cancer-specific evidence-based nutrition guidelines do not exist [12]. Previous studies have investigated the nutritional status of UGI cancer patients in the ambulatory and inpatient settings, with reported malnutrition prevalence between 48 and 52% depending on the assessment method utilised [13–15]. More recently, the INFORM study has investigated the nutritional status of oesophageal and head and neck cancer populations [16]. However, there is a lack of data regarding the nutritional status of UGI cancer patients at the time of curative surgery. Furthermore, little is known regarding preoperative factors that are associated with malnutrition or clinically significant weight loss on presentation for surgery.

The Nutritional Outcomes of patients Undergoing Resection for upper gastroIntestinal cancer in AuStralian Hospitals study (The NOURISH Point Prevalence Study) was conducted to investigate nutritional status and nutritional interventions received by patients undergoing UGI cancer surgery and associations with clinical outcomes, as well as health-service-level practices [17]. NOURISH is the largest study internationally to assess nutritional status in UGI cancer patients at the time of surgery using a validated assessment method [17]. The aims of this study were to determine the prevalence of malnutrition, clinically important weight loss, low muscle strength and nutrition impact symptoms. Secondary aims were to determine factors associated with malnutrition and unintentional weight loss, and to investigate associations between nutritional status and surgical outcomes (complications and length of stay (LOS)). Further outcomes of the NOURISH point prevalence study, including site-specific nutritional practices, perioperative nutritional intervention and nutritional adequacy post-surgery, will be reported in subsequent publications.

2. Materials and Methods

2.1. Study Design and Setting

A prospective, observational point prevalence study was conducted at 27 tertiary hospitals across six states in Australia. Ethics approval was obtained from The Peter MacCallum Cancer Centre Ethics Committee (Melbourne VIC, Australia) prior to commencement (LNR/51107/PMCC-2019). Further details of study design, participating sites and methods are reported in the study protocol [17]. The target sample size was set at a minimum of 200 participants to obtain a minimum precision of $\pm 7\%$ for the 95% confidence interval (CI) of malnutrition prevalence [17].

2.2. Participants

Study dietitians identified and approached eligible patients prior to, or during their surgical admission. Participants were eligible if they were ≥ 18 years, a hospital inpatient having received curative intent surgery for UGI cancer including gastrectomy (total, subtotal, distal, partial), pancreatectomy (total, distal, partial, pancreatico-duodenectomy), oesophagectomy or gastro-oesophagectomy), able to consent to participation by English language communication or with the presence of an interpreter, and if they received assessment of nutritional status with subjective global assessment (SGA) by a dietitian within seven days of surgery. Participants were excluded if they had received palliative surgery, or non-oncological UGI surgery, unable to participate in SGA, unable to provide consent including if they were on intravenous opioids at time of consent, or they were unaware of diagnosis of malignancy. Recruitment commenced on 2 September 2019 and ended on 30 May 2020. Data collection was completed on 30 June 2020. All participants provided verbal consent to participate according to the approved ethics statement [17] and received the standard dietetics care of the participating health service throughout this study.

2.3. Data Collection

Site investigator dietitians conducted a nutritional assessment within seven days of surgery. Patients were also asked to recall information regarding preoperative weight loss, symptoms and prior dietetics or nutritional intervention. The remainder of data were extracted from the medical records. Baseline characteristics included age, sex, primary language spoken, usual social situation, and postcode of residence to determine metropolitan or regional/rural locality. Clinical data included date of diagnosis, tumour type, site and pathological tumour staging (T stage) [18], details of neoadjuvant chemo/radiotherapy if relevant, surgical procedure and surgical technique.

2.4. Outcome Measures

The primary outcome of nutritional status was measured according to the validated SGA, which includes a comprehensive assessment of recent weight loss, symptoms, dietary intake, metabolic requirements, physical and functional status [19]. Current weight was measured using calibrated scales, or was patient reported. Weight on discharge was measured where possible. Patients were asked to recall their weight history from 2 weeks, 1, 3, 6 and >6 months prior to surgery, and if they had lost weight unintentionally prior to surgery. Percentage weight loss was calculated, with clinically important weight loss defined as $\geq 5\%$ in 6 months [20]. Body mass index (BMI) (in kg/m^2) was calculated from weight and height and categorised as underweight (<18.5 if age < 65 or <22 if age ≥ 65), healthy weight ($18.5\text{--}24.9$ if age < 65 or $22\text{--}27$ if age ≥ 65) and overweight/obese (≥ 25 if age < 65 or >27 if age ≥ 65). Upper body muscle strength was measured using hand grip strength (HGS) dynamometry as per the methodology of the American Society of Hand Therapists (ASHT) [21]. Thresholds for diagnosis of low muscle strength were <20 kg for women and <30 kg for men, or <18 kg for women and <26 kg for men for participants of Asian background [20]. Participants were asked to report prevalence of gastrointestinal symptoms persisting > 2 weeks prior to surgery that have been impacting their ability to eat (poor appetite, nausea, vomiting, diarrhoea, constipation, pain when eating, taste changes, dry mouth, problems swallowing or early satiety), as well as any reduction in food intake, degree and length of time of reduction. Discharge destination and LOS (days) were recorded from the medical records. Surgical complications were recorded as documented by the medical team. These included sepsis, anastomotic leak, pancreatic fistula, pneumonia/respiratory tract infection, pneumothorax, pressure injury, wound infection, return to theatre, abdominal collection, ileus, chyle leak, gastroparesis and pleural effusion. Total number of complications were recorded and transformed into a binary variable of 'no complication' or ' ≥ 1 complication' for analysis.

2.5. Statistical Analysis

Statistical analyses were conducted using Stata/IC 16.0 software (StataCorp, College Station, TX, USA, 2019). Descriptive statistics included measurements such as frequencies, percentages, mean and standard deviation (SD), and median and interquartile range (IQR). Differences in nutritional outcomes between surgical procedures were determined using Fisher's exact test. Univariate logistic regression analysis was used to determine demographic and clinical factors associated with malnutrition and clinically significant weight loss prior to surgery. Multivariate logistic regression models were developed to determine factors independently associated with malnutrition and unintentional weight loss. Best models were selected using statistical significance threshold of $p < 0.05$ and goodness of fit R^2 . Multivariate models adjusting for age, surgical procedure, tumour location and tumour stage were developed to determine associations between malnutrition and unintentional weight loss with LOS (continuous outcome, linear regression), and surgical complications (binary outcome, logistic regression).

3. Results

Of the 240 patients screened, 217 met the eligibility criteria and of those, 200 consented to participate. Figure 1 reports participant flow.

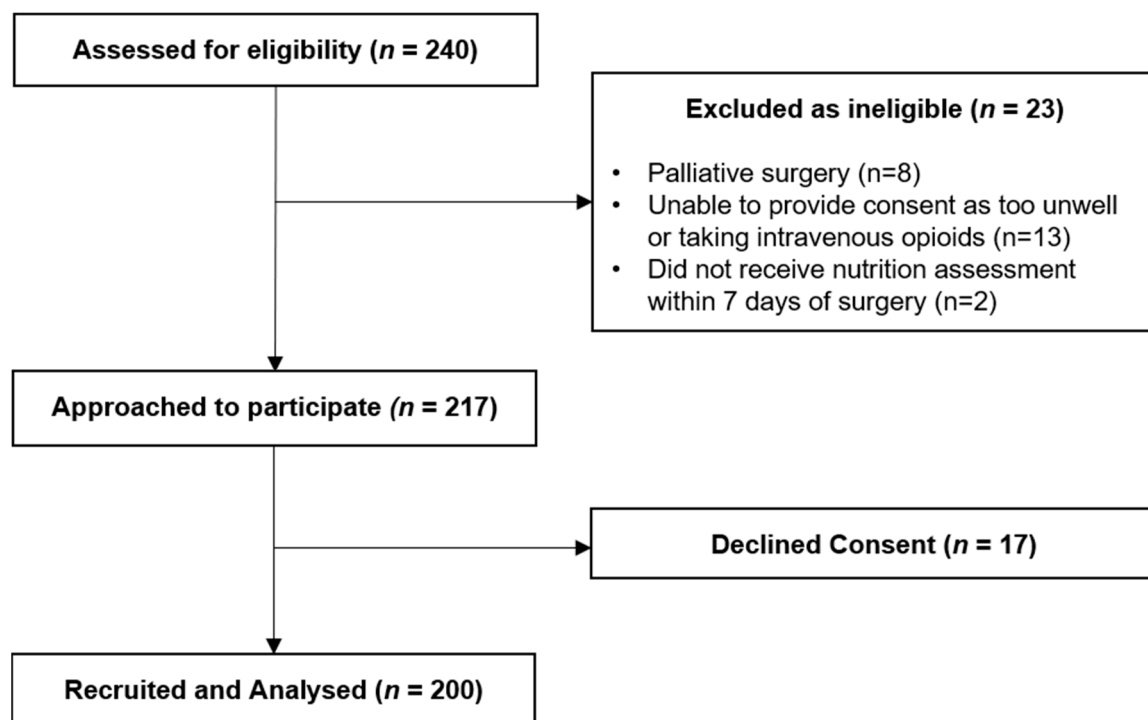


Figure 1. Participant flow diagram.

3.1. Baseline and Clinical Characteristics

Baseline and clinical characteristics of the overall cohort, as well as for each surgery type (oesophagectomy, gastrectomy and pancreatectomy procedures) are presented in Table 1. There were no significant differences for surgery type in age or sociodemographic factors. A higher proportion of males underwent gastrectomy than oesophagectomy and pancreatectomy procedures ($p = 0.002$), while a higher proportion of patients undergoing oesophagectomy and gastrectomy received neoadjuvant therapy than those who underwent pancreatectomy ($p < 0.001$).

Table 1. Demographic, tumour and surgical characteristics by surgery type.

Variables	Overall <i>n</i> = 200	Oesophagectomy <i>n</i> = 66	Gastrectomy <i>n</i> = 50 ^a	Pancreatectomy <i>n</i> = 84 ^b	<i>p</i> Value				
Age (Mean, SD)	67	10	66	9	70	11	66	10	0.111
Sex (<i>n</i> , %)									0.002
Male	117	58.5%	50	75.8%	27	54.0%	40	47.6%	
Female	83	41.5%	16	24.2%	23	46.0%	44	52.4%	
Tumour Location (<i>n</i> , %)									NA
Gastric	50	25.0%	1	1.5%	48	96.0%	1	1.2%	
Oesophageal	60	30.0%	60	90.9%	NA	NA	NA	NA	
Pancreatic	55	27.5%	NA	NA	NA	NA	55	65.5%	
Ampullary	17	8.5%	NA	NA	NA	NA	17	20%	
GOJ	7	3.5%	5	7.6%	2	4.0%	NA	NA	
Bile Duct	6	3.0%	NA	NA	NA	NA	6	7.1%	
Duodenal	5	2.5%	NA	NA	NA	NA	5	6.0%	
Tumour Type (<i>n</i> , %)									NA
Adenocarcinoma	170	85.0%	54	81.8%	49	98.0%	67	79.8%	
SCC	11	5.5%	11	16.7%	0	0.0%	NA	NA	
GIST	2	1.0%	NA	NA	1	2.0%	1	1.2%	
NET	11	5.5%	NA	NA	0	0.0%	11	13.1%	
Other	6	3.0%	1	1.5%	0	0.0%	5	6.0%	
Intraoperative Tumour Stage (<i>n</i> , %)									<0.001
T0	15	7.5%	12	18.2%	2	4.0%	1	1.2%	
T1	44	22.0%	20	30.3%	11	22.0%	13	15.5%	
T2	49	24.5%	10	15.2%	8	16.0%	31	36.9%	
T3	63	31.5%	21	31.8%	20	40.0%	22	26.2%	
T4	14	7.0%	0	0.0%	9	18.0%	5	6.0%	
TX	2	1.0%	0	0.0%	0	0.0%	2	2.4%	
Unknown/Unassessed ^c	13	6.5%	3	4.5%	0	0.0%	10	11.9%	
Received Neoadjuvant Therapy (<i>n</i> , %)									<0.001
No	106	53.3%	17	25.8%	15	30.6%	74	88.1%	
Yes	93	46.7%	49	74.2%	34	69.4%	10	11.9%	
Type of Neoadjuvant Therapy (<i>n</i> , %) ^d									<0.001
Chemotherapy	52	55.9%	10	20.4%	32	94.1%	10	100.0%	
Chemotherapy and Radiotherapy	41	44.1%	39	79.6%	2	5.9%	0	0.0%	
Completed Neoadjuvant Therapy (<i>n</i> , %) ^d									
No	3	3.2%	0	0.0%	3	8.9%	0	0.0%	
Yes	88	94.6%	49	100.0%	29	85.3%	10	100.0%	

Table 1. Cont.

Variables	Overall <i>n</i> = 200		Oesophagectomy <i>n</i> = 66		Gastrectomy <i>n</i> = 50 ^a		Pancreatectomy <i>n</i> = 84 ^b		<i>p</i> Value
Unknown	2	2.2%	0	0.0%	2	5.8%	0	0.0%	0.482
Location of Residence (<i>n</i> , %)									
Metropolitan	143	71.5%	45	68.2%	39	78.0%	59	70.2%	0.872
Rural/Regional	57	28.5%	21	31.8%	11	22.0%	25	29.8%	
Social Situation (<i>n</i> , %)									<0.001
Lives Alone	45	22.5%	13	19.7%	10	20.0%	22	26.2%	
Lives with Family/Carer	152	76.0%	52	78.8%	39	78.0%	61	72.6%	
Lives in Shared Accommodation	0	0.0%	0	0.0%	0	0.0%	0	0.0%	
Lives in Residential Care	3	1.5%	1	1.5%	1	2.0%	1	1.2%	
Surgical Technique (<i>n</i> , %)									
Open	187	93.5%	53	80.3%	50	100.0%	84	100.0%	
Laparoscopic/Minimally Invasive	13	6.5%	13	19.7%	0	0.0%	0	0.0%	

^a Includes total, subtotal, partial and distal gastrectomy. ^b Includes total, distal, partial, pancreatico-duodenectomy. ^c Incomplete information from medical records. ^d Presented as a proportion of participants who responded 'yes' to neoadjuvant therapy (*n* = 93). SCC = squamous cell carcinoma, GIST = gastrointestinal stromal tumour, NET = neuroendocrine tumour, and GOJ = gastro-oesophageal. Bolded *p* values indicate statistical significance.

3.2. Nutritional Status, Weight Loss and Muscle Strength

The overall prevalence of malnutrition according to SGA was 42% ($n = 84$), 95% CI (35%,49%). The prevalence of any degree of unintentional weight loss was 65% ($n = 129$), 95% CI (57%,71%), whilst 47% ($n = 48$), 95% CI (37%,57%) of those who completed the HGS had low muscle strength (Table 2). The HGS test was unable to be completed in 49% ($n = 98$) of the cohort due to unavailability of equipment or participants being too unwell to complete this assessment within seven days of surgery. There were no differences in nutritional status, unintentional weight loss or muscle strength between surgery types (Table 2).

Table 2. Nutritional status and nutrition impact on symptoms by surgery type.

Variable	Overall $n = 200$	Oesophagectomy $n = 66$	Gastrectomy $n = 50$	Pancreatectomy $n = 84$	p Value			
SGA Rating ($n, \%$)					0.689			
A No Malnutrition	116	58.0%	40	60.6%	28	56.0%	48	57.1%
B Mild/Moderate Malnutrition	79	39.5%	24	36.4%	22	44.0%	33	39.3%
C Severe Malnutrition	5	2.5%	2	3.0%	0	0.0%	3	3.6%
Overall Nutritional Status ($n, \%$)					0.864			
Well Nourished (SGA A)	116	58.0%	40	60.6%	28	56.0%	48	57.1%
Malnourished (SGA B/C)	84	42.0%	26	39.4%	22	44.0%	36	42.9%
Unintentional Weight Loss ($n, \%$)					0.645			
No	71	35.5%	24	36.4%	20	40.0%	27	32.1%
Yes	129	64.5%	42	63.6%	30	60.0%	57	67.9%
Low Muscle Strength ($n, \%$)					0.075			
No	54	27.0%	23	34.8%	9	1.08%	22	26.2%
Yes	48	24.0%	12	18.2%	16	32.0%	20	23.8%
Not Completed	98	49.0%	31	47.0%	25	50.0%	42	50.0%
BMI (kg/m^2) (Mean, SD)	27.3	5.6	27.3	5.1	27.5	6.1	27.2	5.7
BMI Categories ($n, \%$)					0.917			
Underweight	20	10.0%	3	4.5%	8	16.0%	9	10.7%
Normal Weight	75	37.5%	30	45.5%	17	34.0%	28	33.3%
Overweight/Obese	105	52.5%	33	50.0%	25	50.0%	47	56.0%
Reduced Dietary Intake before Surgical Admission ($n, \%$)					0.099			
No	101	50.5%	34	51.5%	31	62.0%	36	42.9%
Yes	99	49.5%	32	48.5%	19	38.0%	48	57.1%
Degree of Reduction in Solid Food Intake ($n, \%$)					0.602			
>75% of Usual Intake	27	13.5%	10	15.2%	6	12.0%	11	13.1%
≤75% of Usual Intake	34	17.0%	11	16.7%	6	12.0%	17	20.2%
≤50% of Usual Intake	31	15.5%	8	12.1%	6	12.0%	17	20.2%
≤25% of Usual Intake	7	3.5%	3	4.5%	1	2.0%	3	3.6%
No Reduction in Intake	101	50.5%	34	51.5%	31	62.0%	36	42.9%
Length of Time of Reduced Dietary Intake ($n, \%$)					0.021			
<1 Week	2	1.0%	2	1.5%	0	0.0%	1	1.2%
1–2 Weeks	11	5.5%	2	1.5%	3	6.0%	7	8.3%
2–4 Weeks	13	6.5%	2	1.5%	1	2.0%	11	13.1%
≥1 Month	69	34.5%	29	43.9%	14	28.0%	26	31.0%
No Reduction in Intake	105	52.5%	34	51.5%	32	64.0%	39	46.4%
Symptoms Persisting >2 Weeks Prior to Surgery Impacting								
Ability to Eat								
Poor Appetite								
No	126	63.0%	49	74.2%	30	60.0%	47	56.0%
Yes	74	37.0%	17	25.8%	20	40.0%	37	44.0%
Nausea								
No	172	86.0%	60	90.9%	43	86.0%	69	82.1%
Yes	28	14.0%	6	9.1%	7	14.0%	15	17.9%
Vomiting								
No	187	93.5%	62	93.9%	47	94.0%	78	92.9%
Yes	13	6.5%	4	6.1%	3	6.0%	6	7.1%
Diarrhoea								
No	181	90.5%	66	100.0%	46	92.0%	69	82.1%
Yes	19	9.5%	0	0.0%	4	8.0%	15	17.9%
Constipation								
No	193	96.5%	62	93.9%	49	98.0%	82	97.6%
Yes	7	3.5%	4	6.1%	1	2.0%	2	2.4%
Pain When Eating								
No	178	89.0%	59	89.0%	44	88.0%	75	89.3%
Yes	22	11.0%	7	11.0%	6	12.0%	9	10.7%
Taste Changes								
No	172	86.0%	51	77.3%	41	82.0%	80	95.2%
Yes	28	14.0%	15	22.7%	9	18.0%	4	4.8%
Dry Mouth								
No	191	95.5%	64	97.0%	47	94.0%	80	95.2%
Yes	9	4.5%	2	3.0%	3	6.0%	4	4.8%
Problems Swallowing								
No	185	92.5%	53	80.3%	50	100.0%	82	97.6%
Yes	15	7.5%	13	19.7%	0	0.0%	2	2.4%

Table 2. Cont.

Variable	Overall n = 200		Oesophagectomy n = 66		Gastrectomy n = 50		Pancreatectomy n = 84		p Value
Early Satiety									0.034
No	155	77.5%	57	86.4%	33	66.0%	65	77.4%	
Yes	45	22.5%	9	13.6%	17	34.0%	19	22.6%	
No Problems Reported									0.972
No	110	55.0%	36	54.5%	27	54.0%	47	56.0%	
Yes	90	45.0%	30	45.5%	23	46.0%	37	44.0%	

SGA = subjective global assessment, SD = standard deviation, and BMI = body mass index. Bolded *p* values indicate statistical significance.

3.3. Prevalence of Clinically Significant Weight Loss

Median (IQR) percentage weight loss in 1, 3, 6 and >6 months was 1.2% (0%, 3.2%), 4.8% (1.8%, 8.2%), 7.4% (3.8%, 11.6%) and 8.3% (4.3%, 12.5%), respectively. There were no differences in median weight loss between the surgery types for any timeframe (Supplementary Table S1), except for 1 month before surgery where patients undergoing pancreatectomy procedures lost more weight than those undergoing oesophagectomy and gastrectomy procedures (2.2% (0.0%, 4.5%) vs. 0.5% (0.0%, 2.5%) and 0.0% (0.0%, 1.9%) $p = 0.021$). Figure 2 demonstrates the proportion of patients with clinically significant weight loss of $\geq 5\%$ in the 3 months and 6 months before surgery. Twenty-seven percent of participants reported $\geq 10\%$ weight loss in >6 months. There were no differences in prevalence of clinically significant weight loss at any timepoint before surgery between the surgery groups (Supplementary Table S1).

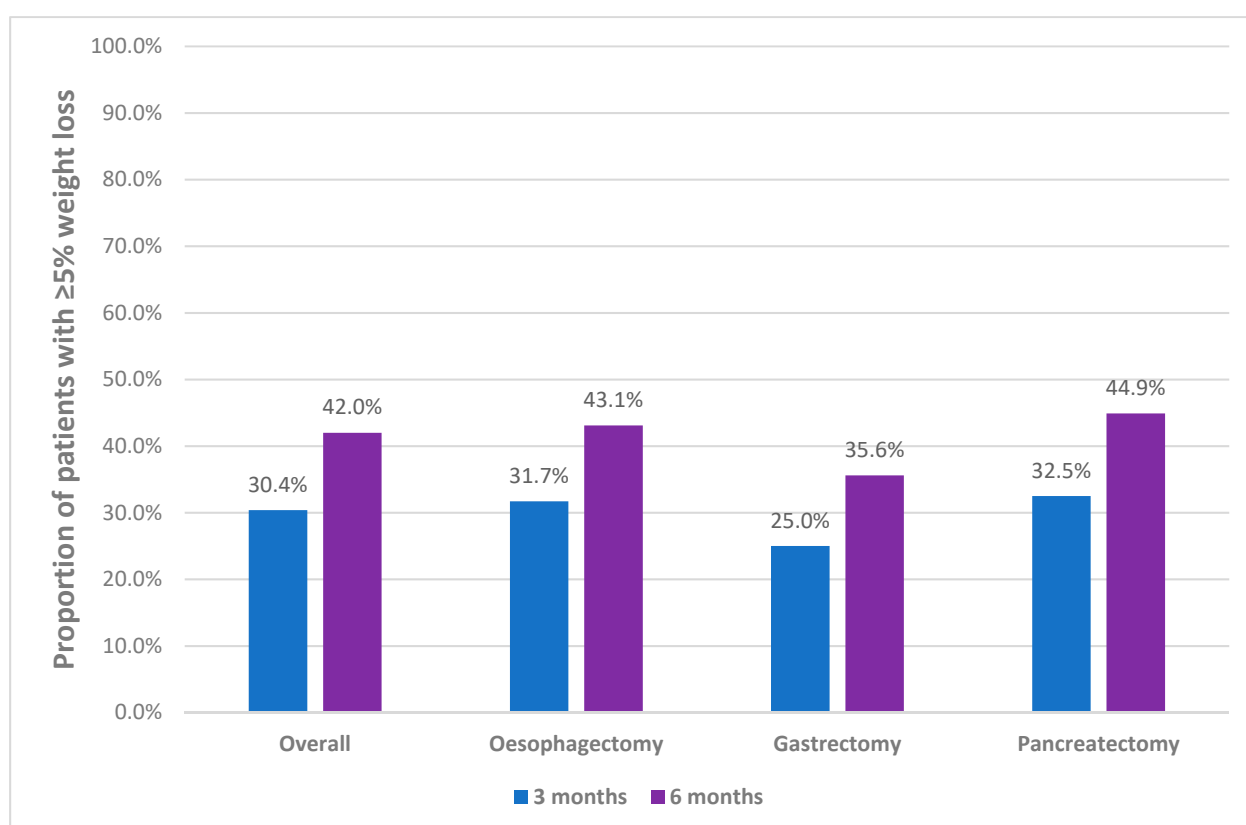


Figure 2. Prevalence of $\geq 5\%$ weight loss in 3 and 6 months.

3.4. Prevalence of Preoperative Nutrition Impact Symptoms

The overall prevalence of reporting at least one preoperative gastrointestinal nutrition impact symptom (Table 2) was 55%, with poor appetite and early satiety being the most reported (37% and 23%, respectively). Patients undergoing pancreatic surgery were more

likely to report preoperative diarrhoea (18% vs. 0% oesophagectomy and 8% gastrectomy, $p = 0.001$), whilst patients undergoing oesophagectomy and gastrectomy were more likely to report taste changes (23% and 18% vs. 5%, $p = 0.005$). Problems swallowing were reported in a significantly higher proportion of oesophagectomy patients (20% versus 0% gastrectomy and 2% pancreatectomy, $p < 0.001$), whilst early satiety was reported in 34% of gastrectomy and 23% of pancreatectomy, compared to 14% oesophagectomy procedures ($p = 0.034$). Reduced dietary intake was reported in 49% of the cohort, with the majority (35%) having reduced intake for ≥ 1 month. Patients undergoing oesophagectomy were more likely to report reduced intake for ≥ 1 month (44% vs. 31% gastrectomy and 28% pancreatectomy, $p = 0.021$).

3.5. Factors Associated with Preoperative Malnutrition and Clinically Significant Weight Loss

On univariate model analysis, factors associated with malnutrition included age, T4 stage, unintentional weight loss of any amount, unintentional weight loss of $\geq 5\%$ and $\geq 10\%$ in six months, reduced dietary intake of all degrees and timeframes, and the presence of any GI symptoms (Tables 3 and 4). On multivariate analysis, age, unintentional weight loss of $\geq 5\%$ in 6 months, vomiting, reduced solid food intake lasting 2–4 weeks and >1 month were independently associated with preoperative malnutrition (Table 5). Factors associated with clinically significant weight loss on univariate analysis included poor appetite, nausea, vomiting, pain when eating, problems swallowing, early satiety, reduced dietary intake, and degree of solid food intake reduction of $<75\%$, 50%, 25% of usual intake for all timeframes (Table 3). Factors independently associated with unintentional weight loss were poor appetite, and degree of solid food reductions of $<75\%$ and $<50\%$ of usual intake (Table 5).

Table 3. Demographic and clinical factors associated with malnutrition and clinically significant weight loss ($\geq 5\%$ in 6 months) prior to surgery ($n = 200$).

Variable	Malnutrition			Unintentional Weight Loss $\geq 5\%$ in 6 Months						
	Bivariate Analysis (Fisher's Exact)		Univariate Logistic Model	Bivariate Analysis (Fisher's Exact)		Univariate Logistic Model				
	WN <i>n</i> (%)	MN <i>n</i> (%)	<i>p</i> Value	OR (95% CI)	<i>p</i> Value	No <i>n</i> (%)	Yes <i>n</i> (%)	<i>p</i> Value	OR (95% CI)	<i>p</i> Value
Age			0.012					0.549		
<65	54 (69.2)	24 (30.8)		1.0 (ref)		45 (60.8)	29 (39.2)		1.0 (ref)	
≥ 65	62 (50.8)	60 (49.2)		2.2 (1.2, 4.0)	0.011	64 (56.1)	50 (43.9)		1.2 (0.7, 2.2)	0.526
Sex (<i>n</i> , %)			0.082					0.654		
Male	74 (63.2)	43 (36.8)		1.0 (ref)		66 (59.5)	45 (40.5)		1.0 (ref)	
Female	42 (50.6)	41 (49.4)		1.7 (0.9, 3.0)	0.075	43 (55.8)	34 (44.2)		1.2 (0.6, 2.1)	0.621
Surgery Type			0.864					0.588		
Oesophagectomy	40 (60.6)	26 (39.4)		1.0 (ref)		37 (56.9)	28 (43.1)		1.0 (ref)	
Gastrectomy	28 (56.0)	22 (44.0)		1.2 (0.6, 2.5)	0.618	29 (64.4)	16 (35.6)		0.7 (0.3, 1.6)	0.429
Pancreatectomy	48 (57.1)	36 (42.9)		1.2 (0.6, 2.2)	0.669	43 (55.1)	35 (44.9)		1.1 (0.6, 2.1)	0.830
Tumour Location			0.926					0.380		
Bile Duct	4 (66.7)	2 (33.3)		1.0 (ref)		4 (66.7)	2 (33.3)		1.0 (ref)	
Gastric	28 (56.0)	22 (44.0)		1.6 (0.3, 9.4)	0.620	29 (63.0)	17 (37.0)		1.2 (0.2, 7.1)	0.862
Oesophageal	37 (61.7)	23 (38.3)		1.2 (0.2, 7.3)	0.810	34 (56.7)	26 (43.3)		1.5 (0.3, 9.0)	0.638
Pancreatic	32 (58.2)	23 (41.8)		1.4 (0.2, 8.5)	0.689	29 (56.9)	22 (43.1)		1.5 (0.3, 9.1)	0.657
Ampullary	10 (58.8)	7 (41.2)		1.4 (0.2, 9.9)	0.736	10 (62.5)	6 (37.5)		1.2 (0.2, 8.7)	0.857
Duodenal	2 (40.0)	3 (60.0)		3 (0.3, 35.3)	0.383	0 (0)	4 (100)		Empty	
GOJ	3 (42.9)	4 (57.1)		2.7 (0.3, 25.6)	0.396	3 (60)	2 (40)		1.3 (0.1, 15.7)	0.819
Tumour Type			0.335					0.912		
Adenocarcinoma	95 (55.9)	75 (44.1)		1.6 (0.3, 8.9)	0.604	91 (57.2)	68 (42.8)		0.8 (0.2, 3.8)	0.726
SCC	6 (54.5)	5 (45.5)		1.7 (0.2, 13.2)	0.629	7 (63.6)	4 (36.4)		0.6 (0.1, 4.3)	0.587
GIST	2 (100)	0 (0)		Empty		1 (50)	1 (50)		1.0 (0.1, 24.6)	1.00
NET	9 (81.8)	2 (18.2)		0.4 (0.0, 4.4)	0.487	7 (70)	3 (30)		0.4 (0.1, 3.5)	0.428
Other	4 (66.7)	2 (33.3)		1.0 (ref)		3 (50)	3 (50)		1.0 (ref)	
Tumour Stage			0.022					0.509		
T0	10 (66.7)	5 (33.3)		1.0 (ref)		9 (64.3)	5 (35.7)		1.0 (ref)	
T1	31 (70.5)	13 (29.5)		0.8 (0.2, 2.9)	0.783	28 (65.1)	15 (34.9)		1.0 (0.3, 3.4)	0.955
T2	29 (59.2)	20 (40.8)		1.4 (0.4, 4.7)	0.604	25 (55.6)	20 (44.4)		1.4 (0.4, 5.0)	0.565
T3	29 (46.0)	34 (54.0)		2.3 (0.7, 7.6)	0.158	31 (51.7)	29 (48.3)		1.7 (0.5, 5.6)	0.397
T4	4 (28.6)	10 (71.4)		5.0 (1.0, 24.3)	0.046	5 (41.7)	7 (58.3)		2.5 (0.5, 12.3)	0.253
Neoadjuvant Therapy			0.774					0.882		
No	60 (56.6)	46 (43.4)		1.0 (ref)		57 (58.8)	40 (41.2)		1.0 (ref)	
Yes	55 (59.1)	38 (40.9)		0.9 (0.5, 1.6)	0.718	51 (56.7)	39 (43.3)		1.1 (0.6, 2.0)	0.772
Type of Neoadjuvant			0.205					0.089		
Chemotherapy	34 (65.4)	18 (34.6)		1.0 (ref)		32 (65.3)	17 (34.7)		1.0 (ref)	
Chemotherapy and Radiotherapy	21 (51.2)	20 (48.8)		0.6 (−0.3, 1.4)	0.169	19 (46.3)	22 (53.7)		2.2 (0.9, 5.1)	0.072
Completed Neoadjuvant			1.000					1.000		
No	2 (66.7)	1 (33.3)		1.0 (ref)		2 (66.7)	1 (33.3)		1.0 (ref)	
Yes	51 (58.0)	37 (42.0)		1.5 (0.1, 16.6)	0.765	47 (55.3)	38 (44.7)		1.6 (0.1, 18.5)	0.699
Location of Residence			0.428					0.137		
Rural/Regional	36 (63.2)	21 (36.8)		1.0 (ref)		35 (67.3)	17 (32.7)		1.0 (ref)	
Metropolitan	80 (55.9)	63 (44.1)		1.3 (0.7, 2.5)	0.352	74 (54.4)	62 (45.6)		1.7 (0.9, 3.4)	0.111
Social Situation			0.132					0.863		
Lives with Family or Carer	94 (61.8)	58 (38.2)		1.0 (ref)		85 (59)	59 (41)		1.0 (ref)	
Lives Alone	21 (46.7)	24 (53.3)		1.8 (0.9, 3.6)	0.07	23 (54.8)	19 (45.2)		1.2 (0.6, 2.4)	0.622
Lives in Residential Care	1 (33.3)	2 (66.7)		3.2 (0.3, 36.5)	0.341	1 (50)	1 (50)		1.4 (0.1, 23.5)	0.798

WN = well nourished, MN = malnourished, OR = odds ratio, CI = confidence interval, SCC = squamous cell carcinoma, GIST = gastrointestinal stromal tumour, NET = neuroendocrine tumour, and GOJ = gastro-oesophgeal. Bolded *p* values indicate statistical significance. Ref = reference value used in the model.

Table 4. Nutrition-related factors associated with malnutrition and clinically significant weight loss ($\geq 5\%$ in 6 months) prior to surgery ($n = 200$).

Variable	Malnutrition			Univariate Logistic Model		Unintentional Weight Loss $\geq 5\%$ in 6 Months				
	Bivariate Analysis (Fisher's Exact)		<i>p</i> Value	OR (95% CI)	<i>p</i> Value	Bivariate Analysis (Fisher's Exact)		Univariate Logistic Model		
	No <i>n</i> (%)	Yes <i>n</i> (%)				No <i>n</i> (%)	Yes <i>n</i> (%)	<i>p</i> Value	OR (95% CI)	<i>p</i> Value
BMI			<0.001					<0.001		
Normal Weight	33 (44.0)	42 (56.0)		1.0 (ref)		37 (53.6)	32 (46.4)		1.0 (ref)	
Underweight	3 (85.0)	17 (85.0)		4.5 (1.2, 16.5)	0.025	3 (16.7)	15 (83.3)		5.8 (1.5, 21.8)	0.010
Overweight/Obese	80 (76.2)	25 (23.8)		0.2 (0.1, 0.5)	<0.001	69 (68.3)	21 (31.7)		0.5 (0.3, 1.1)	0.053
GI Symptoms										
Unchecked				1.0 (ref)					1.0 (ref)	
Poor Appetite	17 (23.0)	57 (77.0)	<0.001	12.3 (6.2, 24.5)	<0.001	18 (26.9)	49 (73.1)	<0.001	8.3 (4.2, 16.3)	<0.001
Nausea	8 (28.6)	20 (71.4)	0.001	4.2 (1.8, 10.1)	0.001	7 (26.9)	19 (73.1)	0.001	4.6 (1.8, 11.6)	0.001
Vomiting	1 (7.7)	12 (92.3)	<0.001	19.2 (2.4, 150.6)	0.005	3 (23.1)	10 (76.9)	0.016	5.1 (1.4, 19.3)	0.016
Diarrhoea	4 (21.1)	15 (78.9)	0.001	6.1 (1.9, 19.1)	0.002	6 (35.3)	11 (54.7)	0.069	2.8 (1.0, 7.9)	0.054
Constipation	1 (14.3)	6 (85.7)	0.043	8.8 (1.0, 74.9)	0.046	2 (28.6)	5 (71.4)	0.133	3.6 (0.7, 19.1)	0.131
Pain When Eating	8 (36.4)	14 (63.6)	0.039	2.7 (1.1, 6.8)	0.034	4 (21.1)	15 (78.9)	0.001	6.2 (2.0, 19.4)	0.002
Taste Changes	10 (35.7)	18 (63.3)	0.013	2.9 (1.3, 6.6)	0.012	13 (48.1)	14 (51.9)	0.296	1.6 (0.7, 3.6)	0.266
Dry Mouth	2 (22.2)	7 (77.8)	0.037	5.2 (1.0, 25.6)	0.044	4 (50)	4 (50)	0.722	1.4 (0.3, 5.8)	0.642
Problems Swallowing	3 (20.0)	12 (80.0)	0.002	6.3 (1.7, 23.0)	0.006	4 (28.6)	10 (71.4)	0.025	3.8 (1.2, 12.6)	0.029
Early Satiety	16 (35.6)	29 (64.4)	0.001	3.3 (1.6, 6.6)	0.001	17 (42.5)	23 (57.5)	0.031	2.2 (1.1, 4.5)	0.027
Any LOW			<0.001			NA			NA	
No	67 (94.4)	4 (5.6)		1.0 (ref)	<0.001					
Yes	49 (38.0)	80 (62.0)		27.3 (9.4, 79.7)	<0.001	NA			NA	
LOW $\geq 5\%$ in 6 Months			<0.001							
No	29 (74.4)	10 (25.6)		1.0 (ref)						
Yes	14 (21.5)	51 (78.5)		29.1 (13.1, 64.6)	<0.001					
LOW $\geq 10\%$ in 6 Months			<0.001			NA			NA	
No	40 (58.0)	29 (42.0)		1.0 (ref)						
Yes	3 (8.6)	32 (91.4)		40.0 (11.6, 138.1)	<0.001					
Reduced Dietary Intake			<0.001					<0.001		
No	84 (83.2)	17 (16.8)		1.0 (ref)		75 (77.3)	22 (22.7)		1.0 (ref)	
Yes	32 (32.3)	67 (67.7)		10.3 (5.3, 20.2)	<0.001	34 (37.4)	57 (62.6)		5.7 (3.0, 10.8)	<0.001
Degree of Reduction in Solid Food Intake			<0.001					<0.001		
No Reduction in Intake	84 (83.2)	17 (16.8)		1.0 (ref)		75 (77.3)	22 (22.7)		1.0 (ref)	
>75% of Usual Intake	17 (63.0)	10 (37.0)		2.9 (1.1, 7.4)	0.026	16 (69.6)	7 (30.4)		1.5 (0.6, 4.1)	0.437
$\leq 75\%$ of Usual Intake	8 (23.5)	26 (76.5)		16.1 (6.2, 41.5)	<0.001	10 (32.3)	21 (67.7)		7.2 (2.9, 17.4)	<0.001
$\leq 50\%$ of Usual Intake	5 (16.1)	26 (83.9)		25.7 (8.6, 76.4)	<0.001	6 (20)	24 (80)		13.6 (5.0, 37.6)	<0.001
$\leq 25\%$ of Usual Intake	2 (28.6)	5 (71.4)		12.4 (2.2, 69.1)	0.004	2 (28.6)	5 (71.4)		8.5 (1.6, 47.0)	0.014
Length of Time of Reduced Dietary Intake			<0.001					<0.001		
No Reduction in Intake	88 (83.8)	17 (16.2)		1.0 (ref)		78 (77.2)	23 (22.8)		1.0 (ref)	
<1 Week	2 (100.0)	0 (0)		Empty		1 (100)	0 (0)		(empty)	
1–2 Weeks	5 (45.5)	6 (54.5)		6.2 (1.7, 22.7)	0.006	4 (36.4)	7 (63.6)		5.9 (1.6, 22.1)	0.008
2–4 Weeks	5 (38.5)	8 (61.5)		8.3 (2.4, 28.4)	0.001	6 (46.2)	7 (53.8)		4.0 (1.2, 13.0)	0.023
≥ 1 Month	16 (23.2)	53 (76.8)		17.1 (8.0, 36.8)	<0.001	20 (32.3)	42 (67.7)		7.1 (3.5, 14.4)	<0.001

OR = odds ratio, CI = confidence interval, GI = Gastrointestinal and LOW = loss of weight. Bolded *p* values indicate statistical significance. Ref = reference value used in the model.

Table 5. Factors independently associated with malnutrition and weight loss $\geq 5\%$ by multivariate analysis ($n = 200$).

Variable	Malnutrition OR (95% CI)	<i>p</i> Value	Unintentional Weight Loss $\geq 5\%$ OR (95% CI)	<i>p</i> Value
Age ≥ 65	4.1 (1.5, 11.5)	0.008		
LOW $\geq 5\%$ in 6 Months	28.7 (10.5, 78.6)	<0.001		
Length of Time of Reduced Intake				
2–4 Weeks	7.4 (1.3, 43.5)	0.026		
≥ 1 Month	7.7 (2.7, 22.0)	<0.001		
Degree of reduction in solid food intake				
$\leq 75\%$ of Usual Intake			3.3 (1.2, 9.2)	0.02
$\leq 50\%$ of Usual Intake			4.9 (1.5, 15.6)	0.008
Nutrition Impact Symptoms				
Vomiting	17.1 (1.4, 207.6)	0.025		
Poor Appetite			3.7 (1.6, 8.4)	0.002

OR = odds ratio, CI = confidence interval, and LOW = loss of weight. Bolded *p* values indicate statistical significance.

3.6. Associations between Preoperative Nutritional Status and Clinically Significant Weight Loss, with Surgical Length of Stay and Complications

Using bivariate analysis, patients who were malnourished had an increased surgical LOS of 4 days compared with well-nourished participants (14 days, IQR 8,18 versus 10 days, IQR 8,14, $p = 0.046$). Patients with unintentional weight loss of $\geq 5\%$ in 6 months also had an increased LOS of 4 days compared with those with $\leq 5\%$ (14 days, IQR 8, 19.3 versus 10 days IQR 8,14, $p = 0.007$). On multivariate model analysis (adjusting for age, surgical procedure, tumour location and stage), malnutrition (Coefficient 3.6, 95% CI 0.1, 7.2, $p = 0.048$) was independently associated with increased LOS, as was unintentional weight loss (Coefficient 4.1, 95% CI 0.5,7.6, $p = 0.026$). There were no differences in prevalence of surgical complications between malnourished or well-nourished participants, or patients with clinically significant weight loss on bivariate, univariate, or multivariate model analysis.

4. Discussion

NOURISH is the largest study to assess nutritional status in UGI cancer patients at the time of curative intent surgery using a validated assessment method. The majority of other studies of nutritional status in UGI cancer have been undertaken in ambulatory populations or have been included as part of mixed cancer or gastrointestinal surgical populations [14,22–27]. This study is unique as patients were recruited from 27 tertiary health services at the time of curative intent surgery, only major UGI surgical oncology procedures were included, and the validated SGA tool was used to diagnose malnutrition.

4.1. Nutritional Status, Weight Loss and Muscle Strength

Overall, 42% of participants were malnourished which is relatively consistent with previous studies investigating nutritional status in UGI cancer cohorts, despite some differences in methodology, population or setting [14,23,24]. Interestingly, there were no differences between malnutrition prevalence rates between surgical procedures or tumour types, indicating that all patients presenting for major UGI cancer surgeries are equally at high risk. In this study, almost one-third (30.4%) of patients had clinically significant weight loss within the three months prior to surgery. This is particularly relevant for patients with oesophageal and gastric tumours undergoing neoadjuvant therapy (which typically spans three months including the post-treatment washout period), as weight loss during chemo/radiotherapy is associated with treatment intolerance and decreased survival [24]. It appears that nutritional interventions provided to participants of this study may not be adequate given the high rates of ongoing weight loss and malnutrition.

A lower proportion of patients undergoing pancreatic procedures received preoperative chemotherapy treatment, which is consistent with current clinical practice [28]. However, this group presented with the highest proportion of preoperative weight loss for the 2 week, 1 month and 3 month periods prior to surgery, indicating rapid weight loss that is more likely to be associated with tumour progression and associated symptoms, rather than the effects of chemotherapy. This indicates the need for early dietetics intervention in the hospital outpatient clinic setting for patients undergoing pancreatic surgery, which is typically less resourced than the chemoradiotherapy setting [9].

Although 47% of participants had low muscle strength according to HGS, only 51% of participants were able to complete the test within the seven-day post-operative period, limiting interpretation of these results. Whilst other studies have demonstrated that using HGS for GI surgical patients is a valid method of muscle strength assessment [22,29], these studies have undertaken the test within two–five days of surgery in cohorts consisting largely of general or colorectal GI surgical procedures. This was not often possible in this study due to patients being too unwell post-major UGI surgery. Future studies should attempt to complete the HGS on admission for surgery in UGI patients. However, this was unable to be completed in our study due to funding and staff limitations.

4.2. Prevalence of Preoperative Nutrition Impact Symptoms

Over half of the NOURISH cohort experienced nutrition impact symptoms persisting for two or more weeks preoperatively, with symptoms consistent with clinical expectations for each cancer type. This highlights the importance of targeted screening for GI symptoms known to be related to each cancer type and ensuring appropriate intervention. For example, pancreatic enzyme replacement therapy is recommended to improve symptoms of early satiety and diarrhoea in patients with pancreatic cancer [30]. In oesophageal cancer, previous studies have also demonstrated that patients commonly report dysphagia and poor appetite prior to the commencement of preoperative chemo/radiotherapy [31,32]. However, these symptoms are typically thought to improve after neoadjuvant therapy and its associated side effects subside [33]. In this study, dysphagia and pain when eating were still present in 31% of patients with oesophageal cancer at the time of surgery, indicating that symptoms continue for many patients after the cessation of neoadjuvant treatment. This has implications on the provision of nutrition support, as patients may not have as much contact with the dietitian or treating team in between neoadjuvant treatment completion and surgery.

4.3. Factors Associated with Malnutrition and Unintentional Weight Loss

Whilst a number of factors were associated with malnutrition on univariate analysis, age, unintentional weight loss of $\geq 5\%$ in 6 months, vomiting and reduced dietary intake remained independently associated on multivariate analysis. Unintentional weight loss of $\geq 5\%$ in 6 months is well recognised as a key component of a malnutrition diagnosis across all clinical populations and is a phenotypical criterion of the recently developed Global Leadership Initiative on Malnutrition (GLIM) criteria [20]. Our study confirms that it is associated with higher risk of malnutrition in people with UGI cancer. Previous studies in both surgical and oncological cohorts have identified age as an independent factor for malnutrition [14,34], which can impact on clinical outcomes. A recent study demonstrated that malnutrition in older adults with gastrointestinal cancer was associated with impairments in geriatric assessment measures [35]. This study reinforces recommendations that age should be considered when performing surgical risk stratification, and nutritional intervention and exercise should be provided to improve clinical outcomes [6]. Vomiting and loss of appetite were also independently associated with malnutrition in a study of 4783 cancer patients [34], whilst a higher symptom burden has also been associated with poorer nutritional status in a longitudinal study of oesophago-gastric cancer patients undergoing radical treatment [32]. However, this is the first study, to our knowledge, to investigate symptoms persisting at the time of surgery. Similarly, poor appetite and

a reduced food intake were associated with clinically significant weight loss. Current oncological guidelines recommend that gastrointestinal symptoms should be monitored as risk factors for malnutrition [36]. The results of this study further highlight the importance of monitoring symptoms and dietary intake, which may not be captured in some malnutrition screening tools. Given that nutritional intervention should be provided for at least 7–10 days before surgery to have an effect on surgical outcomes [5], screening of symptoms and dietary intake should be conducted as early as possible, and ideally should be repeated at frequent intervals until one week prior to surgery.

4.4. Associations between Nutritional Status and Surgical Outcomes

Malnutrition and unintentional weight loss of $\geq 5\%$ in 6 months were both independently associated with a longer surgical LOS. This finding consolidates previous research demonstrating that malnutrition and unintentional weight loss are key modifiable risk factors for post-operative outcomes and that preoperative nutrition intervention is essential [5]. Malnutrition has been linked to an increased risk of surgical complications previously [23]; however, this was not demonstrated in our study. Although we recorded complications known to be affected by malnutrition, site investigators did not classify complications into severe/non-severe as per the Clavien Dindo grading system [37] due to funding and time constraints. However, analysis of each complication separately did not reveal any significant differences between malnourished and well-nourished participants. Patients who were malnourished may have been more likely to receive perioperative enteral or parenteral nutrition support, which has not been accounted for in this analysis.

4.5. Strengths and Limitations

Strengths of this study include the large sample size for a UGI surgical oncology cohort, with representation from public and private hospital settings across six Australian states. The SGA is a validated nutritional assessment tool in both surgical and oncology populations. Limitations include the inability to assess muscle mass using an objective tool, such as computed tomography (CT) analysis, as well as the low number of participants who were able to perform the HGS test post-surgery. Although standardised training was conducted to ensure inter-rater reliability, and all data collectors were experienced clinical dietitians, data collection bias due to multiple assessors cannot be excluded.

5. Conclusions

The present study demonstrates that a high proportion of patients undergoing major UGI oncological resections still present with malnutrition or clinically significant weight loss at the time of curative surgery, despite increasing recognition of the importance of preoperative nutritional intervention. Risk factors including age, presence of GI symptoms and decreased food intake prior to surgery should be considered during preoperative nutritional risk stratification in clinical practice and appropriate nutritional intervention should be provided, as outlined by current oncology guidelines. Further research is required to determine if current practice for dietetics intervention prior to UGI surgery has a positive impact on clinical outcomes. This could support prioritization of service delivery improvement initiatives and nutrition research trials in this high-risk group.

Supplementary Materials: The following are available online at <https://www.mdpi.com/article/10.3390/nu13103349/s1>. Table S1: Unintentional Weight Loss by Surgery Type.

Author Contributions: I.D. is the coordinating investigator and conceptualized this study. N.K., J.M.C.Y., E.I. and V.M.C. contributed to the study conception and design. I.D. drafted and edited the study protocol and manuscript. N.K., J.M.C.Y., V.M.C. and E.I. assisted with the study protocol and edited the manuscript. J.A. and I.D. performed the statistical analysis. Members of the NOURISH Point Prevalence Study Group provided feedback on the study protocol and manuscript, carried out the study recruitment and data collection. All authors have read and agreed to the published version of the manuscript.

Funding: An unrestricted project grant has been received from Nestlé Health Science to support out of pocket expenses (ethics application administration/processing fees at participating sites) for the operation of this study. The funder was not involved in the study design, data collection, results analysis, interpretation or dissemination. Publication costs are supported by a grant from the Australian Society of Enteral and Parenteral Nutrition (AuSPEN). All study investigators were unfunded.

Institutional Review Board Statement: This study was conducted according to the guidelines of the Declaration of Helsinki, and approved by The Peter MacCallum Cancer Centre Ethics Committee prior to commencement (LNR/51107/PMCC-2019) in May 2019.

Informed Consent Statement: Informed consent was obtained from all subjects involved in the study.

Data Availability Statement: Not applicable.

Acknowledgments: Members of the NOURISH Point Prevalence Study Group: Anna Cardamis (Eastern Health, Melbourne), Annika Dorey (Cabrini Hospital, Melbourne), Aurora Ottaway (Western Health, Melbourne), Brook Maguire (Westmead Hospital, Sydney), Brydie Cleeve (Epworth HealthCare, Melbourne), Caitlin Davis (Flinders Medical Centre, Adelaide), Carmel Zoanetti (Epworth HealthCare, Melbourne), Catrina Gray (Epworth HealthCare, Melbourne) Christine Choong (Austin Health, Melbourne), Claire Douglas (Prince of Wales Hospital, Sydney), Claire Nixon (The Royal Adelaide Hospital, Adelaide), Daniel Platt (Cabrini Hospital, Melbourne) Eleanor Quinn (John Hunter Hospital, Newcastle), Eliza Simpson (The Royal Adelaide Hospital, Adelaide), Emma Hamdorf (The Queen Elizabeth Hospital, Adelaide), Emma McNamara (John Hunter Hospital, Newcastle), Emma Whelan (Charles Gairdner Hospital, Perth), Gayathri Jegendran (Liverpool Hospital, Sydney), Georgia Moore (Eastern Health, Melbourne), Georgina Lockwood (Sunshine Coast Hospital and Health Service, Sunshine Coast), Jacqueline McNamara (Western Health, Melbourne), Jemma Corriگان (Canberra Hospital, Canberra), Karina Haaksma (Prince of Wales Hospital, Sydney), Kate Fox (St Vincent’s Hospital Melbourne, Melbourne), Kate Furness (Monash Health, Melbourne), Kiah Witney Cochrane (Royal Prince Alfred Hospital, Sydney), Kieu Huynh (Bankstown Lidcombe Hospital, Sydney), Kai Chyi Lee (Fiona Stanley Hospital, Perth), Nadia Hames (Westmead Hospital, Sydney), Nadia Hendricks (St Vincent’s Hospital Melbourne, Melbourne), Naomi Page (St George Hospital, Sydney), Natalie Brooks (Fiona Stanley Hospital, Perth), Lauren Nevin (Barwon Health), Lindy Parfrey (The Queen Elizabeth Hospital, Adelaide), Emma Putrus (The Royal Adelaide Hospital, Adelaide), Rachel Pons (St George Hospital, Sydney), Roy Hoevenaars (Barwon Health, Geelong), Sheena Singh (Concord Hospital, Sydney), Simone McCoy (Princess Alexandra Hospital, Brisbane), Siobhan Wallin (The Royal Brisbane and Women’s Hospital, Brisbane), Stella Mexias (Northern Health, Melbourne), Suzie Daniells (Prince of Wales Hospital, Sydney), Tayla Storr (The Townsville Hospital and Health Service, Townsville), Tayla Robertson (Princess Alexandra Hospital, Brisbane), and Teresa Brown (The Royal Brisbane and Women’s Hospital, Brisbane).

Conflicts of Interest: The authors have nil conflicts of interest to declare.

References

1. Bray, F.; Ferlay, J.; Soerjomataram, I.; Siegel, R.L.; Torre, L.A.; Jemal, A. Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J. Clin.* **2018**, *68*, 394–424. [\[CrossRef\]](#)
2. Lee, J.H.; Nam, B.; Ryu, K.W.; Ryu, S.Y.; Park, Y.K.; Kim, S.; Kim, Y.-W. Comparison of outcomes after laparoscopy-assisted and open total gastrectomy for early gastric cancer. *BJS* **2015**, *102*, 1500–1505. [\[CrossRef\]](#) [\[PubMed\]](#)
3. McGuigan, A.; Kelly, P.; Turkington, R.; Jones, C.; Coleman, H.G.; McCain, R.S. Pancreatic cancer: A review of clinical diagnosis, epidemiology, treatment and outcomes. *World J. Gastroenterol.* **2018**, *24*, 4846–4861. [\[CrossRef\]](#)
4. Fransen, L.F.C.; Luyer, M.D.P. Effects of improving outcomes after esophagectomy on the short- and long-term: A review of literature. *J. Thorac. Dis.* **2019**, *11*, S845–S850. [\[CrossRef\]](#) [\[PubMed\]](#)
5. Weimann, A.; Braga, M.; Carli, F.; Higashiguchi, T.; Hübner, M.; Klek, S.; Laviano, A.; Ljungqvist, O.; Lobo, D.N.; Martindale, R.; et al. ESPEN guideline: Clinical nutrition in surgery. *Clin. Nutr.* **2017**, *36*, 623–650. [\[CrossRef\]](#) [\[PubMed\]](#)
6. Lobo, D.N.; Gianotti, L.; Adiamah, A.; Barazzoni, R.; Deutz, N.E.; Dhatriya, K.; Greenhaff, P.L.; Hiesmayr, M.; Jakobsen, D.H.; Klek, S.; et al. Perioperative nutrition: Recommendations from the ESPEN expert group. *Clin. Nutr.* **2020**, *39*, 3211–3227. [\[CrossRef\]](#) [\[PubMed\]](#)
7. Arends, J.; Baracos, V.; Bertz, H.; Bozzetti, F.; Calder, P.; Deutz, N.; Erickson, N.; Laviano, A.; Lisanti, M.; Lobo, D.; et al. ESPEN expert group recommendations for action against cancer-related malnutrition. *Clin. Nutr.* **2017**, *36*, 1187–1196. [\[CrossRef\]](#)
8. Muscaritoli, M.; Arends, J.; Aapro, M. From guidelines to clinical practice: A roadmap for oncologists for nutrition therapy for cancer patients. *Ther. Adv. Med. Oncol.* **2019**, *11*, 1–14. [\[CrossRef\]](#)

9. Carey, S.; He, L.; Ferrie, S. Nutritional management of patients undergoing major upper gastrointestinal surgery: A survey of current practice in Australia. *Nutr. Diet.* **2010**, *67*, 219–223. [[CrossRef](#)]
10. Williams, J.; Wischmeyer, P.E. Assessment of perioperative nutrition practices and attitudes—A national survey of colorectal and GI surgical oncology programs. *Am. J. Surg.* **2017**, *213*, 1010–1018. [[CrossRef](#)]
11. Deftereos, I.; Kiss, N.; Isenring, E.; Carter, V.M.; Yeung, J.M. A systematic review of the effect of preoperative nutrition support on nutritional status and treatment outcomes in upper gastrointestinal cancer resection. *Eur. J. Surg. Oncol.* **2020**, *46*, 1423–1434. [[CrossRef](#)] [[PubMed](#)]
12. Lu, Y.; Carey, S. Translating Evidence-Based Practice Guidelines into a Summary of Recommendations for the Nutrition Management of Upper Gastrointestinal Cancers. *Nutr. Clin. Pract.* **2014**, *29*, 518–525. [[CrossRef](#)] [[PubMed](#)]
13. Bozzetti, F.; Mariani, L.; Lo Vullo, S.; Amerio, M.L.; Biffi, R.; Caccialanza, G.; Correja, I.; Cozzaglio, L.; Di Leo, A.; Di Cosmo, L.; et al. The nutritional risk in oncology: A study of 1453 cancer outpatients. *Support. Care Cancer* **2012**, *20*, 1919–1928. [[CrossRef](#)]
14. Marshall, K.M.; Loeliger, J.; Nolte, L.; Kelaart, A.; Kiss, N.K. Prevalence of malnutrition and impact on clinical outcomes in cancer services: A comparison of two time points. *Clin. Nutr.* **2019**, *38*, 644–651. [[CrossRef](#)] [[PubMed](#)]
15. Hébuterne, X.; Lemarié, E.; Michallet, M.; De Montreuil, C.B.; Schneider, S.; Goldwasser, F. Prevalence of Malnutrition and Current Use of Nutrition Support in Patients with Cancer. *J. Parenter. Enter. Nutr.* **2014**, *38*, 196–204. [[CrossRef](#)] [[PubMed](#)]
16. Findlay, M.; Bauer, J.D.; Dhaliwal, R.; De Van Der Schueren, M.; Laviano, A.; Widaman, A.; Martin, L.; Day, A.G.; Gramlich, L.M. Translating Evidence-Based Guidelines into Practice—Are We Getting It Right? A Multi-Centre Prospective International Audit of Nutrition Care in Patients with Foregut Tumors (INFORM). *Nutrients* **2020**, *12*, 3808. [[CrossRef](#)]
17. Deftereos, I.; Yeung, J.M.C.; Carter, V.M.; Isenring, E.; Kiss, N.K. Nutritional Outcomes of patients Undergoing Resection for upper gastroIntestinal cancer in AuStralian Hospitals (NOURISH): Protocol for a multicentre point prevalence study. *BMJ Open* **2020**, *10*, e035824. [[CrossRef](#)] [[PubMed](#)]
18. Brody, T. *Clinical Trials: Study Design, Endpoints and Biomarkers, Drug Safety, and FDA and ICH Guidelines*; Elsevier Science & Technology: St. Louis, MO, USA, 2011.
19. Detsky, A.S.; McLaughlin, J.R.; Baker, J.P.; Johnston, N.; Whittaker, S.; Mendelson, R.A.; Jeejeebhoy, K.N. What is subjective global assessment of nutritional status? *JPEN J. Parenter. Enter. Nutr.* **1987**, *11*, 8–13. [[CrossRef](#)]
20. Jensen, G.L.; Cederholm, T.; Correia, M.; Gonzalez, M.C.; Fukushima, R.; Higashiguchi, T.; de Baptista, G.A.; Barazzoni, R.; Blaauw, R.; Coats, A.J.S.; et al. GLIM Criteria for the Diagnosis of Malnutrition: A Consensus Report from the Global Clinical Nutrition Community. *JPEN J. Parenter. Enter. Nutr.* **2019**, *43*, 32–40. [[CrossRef](#)]
21. Roberts, H.C.; Denison, H.; Martin, H.J.; Patel, H.P.; Syddall, H.; Cooper, C.; Sayer, A.A. A review of the measurement of grip strength in clinical and epidemiological studies: Towards a standardised approach. *Age Ageing* **2011**, *40*, 423–429. [[CrossRef](#)]
22. Narendra, K.; Kiss, N.; Margerison, C.; Johnston, B.; Chapman, B. Impact of nutritional status/risk and post-operative nutritional management on clinical outcomes in patients undergoing gastrointestinal surgery: A prospective observational study. *J. Hum. Nutr. Diet.* **2020**, *33*, 587–597. [[CrossRef](#)]
23. Garth, A.K.; Newsome, C.M.; Simmance, N.; Crowe, T.C. Nutritional status, nutrition practices and post-operative complications in patients with gastrointestinal cancer. *J. Hum. Nutr. Diet.* **2010**, *23*, 393–401. [[CrossRef](#)] [[PubMed](#)]
24. Hill, A.; Kiss, N.; Hodgson, B.; Crowe, T.C.; Walsh, A.D. Associations between nutritional status, weight loss, radiotherapy treatment toxicity and treatment outcomes in gastrointestinal cancer patients. *Clin. Nutr.* **2011**, *30*, 92–98. [[CrossRef](#)]
25. Attar, A.; Malka, D.; Sabaté, J.M.; Bonnetain, F.; LeComte, T.; Aparicio, T.; Locher, C.; Laharie, D.; Ezenfis, J.; Taieb, J. Malnutrition Is High and Underestimated During Chemotherapy in Gastrointestinal Cancer: An AGEO Prospective Cross-Sectional Multicenter Study. *Nutr. Cancer* **2012**, *64*, 535–542. [[CrossRef](#)]
26. Cao, J.; Xu, H.; Li, W.; Guo, Z.; Lin, Y.; Shi, Y.; Hu, W.; Ba, Y.; Li, S.; Li, Z.; et al. Nutritional assessment and risk factors associated to malnutrition in patients with esophageal cancer. *Curr. Probl. Cancer* **2021**, *45*, 100638. [[CrossRef](#)] [[PubMed](#)]
27. Guo, Z.Q.; Yu, J.M.; Li, W.; Fu, Z.; Lin, Y.; Shi, Y.Y.; Hu, W.; Ba, Y.; Li, S.Y.; Li, Z.N.; et al. Survey and analysis of the nutritional status in hospitalized patients with malignant gastric tumors and its influence on the quality of life. *Support. Care Cancer* **2020**, *28*, 373–380. [[CrossRef](#)]
28. Lambert, A.; Schwarz, L.; Borbath, I.; Henry, A.; Van Laethem, J.-L.; Malka, D.; Ducreux, M.; Conroy, T. An update on treatment options for pancreatic adenocarcinoma. *Ther. Adv. Med. Oncol.* **2019**, *11*, 1–43. [[CrossRef](#)] [[PubMed](#)]
29. Byrnes, A.; Mudge, A.; Young, A.; Banks, M.; Bauer, J. Use of hand grip strength in nutrition risk screening of older patients admitted to general surgical wards. *Nutr. Diet.* **2018**, *75*, 520–526. [[CrossRef](#)] [[PubMed](#)]
30. Powell-Brett, S.; Carino, N.D.L.; Roberts, K. Understanding pancreatic exocrine insufficiency and replacement therapy in pancreatic cancer. *Eur. J. Surg. Oncol.* **2020**, *47*, 539–544. [[CrossRef](#)]
31. Movahed, S.; Tabrizi, F.V.; Pahlavani, N.; Toussi, M.S.; Motlagh, A.G.; Eslami, S.; Ghayour-Mobarhan, M.; Nematy, M.; Ferns, G.A.; Emadzadeh, M.; et al. Comprehensive assessment of nutritional status and nutritional-related complications in newly diagnosed esophageal cancer patients: A cross-sectional study. *Clin. Nutr.* **2021**, in press. [[CrossRef](#)]
32. Grace, E.M.; Shaw, C.; Lalji, A.; Mohammed, K.; Andreyev, H.J.N.; Whelan, K. Nutritional status, the development and persistence of malnutrition and dietary intake in oesophago-gastric cancer: A longitudinal cohort study. *J. Hum. Nutr. Diet.* **2018**, *31*, 785–792. [[CrossRef](#)] [[PubMed](#)]
33. Bower, M.; Jones, W.; Vessels, B.; Scoggins, C.; Martin, R. Nutritional Support with Endoluminal Stenting During Neoadjuvant Therapy for Esophageal Malignancy. *Ann. Surg. Oncol.* **2009**, *16*, 3161–3168. [[CrossRef](#)] [[PubMed](#)]

34. De Pinho, N.B.; Martucci, R.B.; Rodrigues, V.D.; D'Almeida, C.A.; Thuler, L.C.S.; Saunders, C.; Jager-Wittenaar, H.; Peres, W.A.F. Malnutrition associated with nutrition impact symptoms and localization of the disease: Results of a multicentric research on oncological nutrition. *Clin. Nutr.* **2019**, *38*, 1274–1279. [[CrossRef](#)] [[PubMed](#)]
35. Williams, G.R.; Al-Obaidi, M.; Dai, C.; Mir, N.; Challa, S.A.; Ms, M.D.; Patel, H.; Barlow, B.; Young-Smith, C.; Gbolahan, O.; et al. Association of malnutrition with geriatric assessment impairments and health-related quality of life among older adults with gastrointestinal malignancies. *Cancer* **2020**, *126*, 5147–5155. [[CrossRef](#)]
36. Arends, J.; Bachmann, P.; Baracos, V.; Barthelemy, N.; Bertz, H.; Bozzetti, F.; Fearon, K.; Hütterer, E.; Isenring, E.; Kaasa, S.; et al. ESPEN guidelines on nutrition in cancer patients. *Clin. Nutr.* **2017**, *36*, 11–48. [[CrossRef](#)]
37. Dindo, D.; Demartines, N.; Clavien, P.A. Classification of surgical complications: A new proposal with evaluation in a cohort of 6336 patients and results of a survey. *Ann. Surg.* **2004**, *240*, 205–213. [[CrossRef](#)]