

Construction of Nursing Management Protocol for Promoting Tolerance of Enteral Nutrition in Tube-Fed Perioperative Cancer Patients

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Purpose: Enteral nutrition tolerance studies are mainly focused on critical care, while perioperative nursing research for cancer patients is relatively scattered. This study aims to develop an evidence-based management protocol for promoting tolerance to perioperative enteral nutrition (tube feeding) for cancer patients in general surgical wards, to provide a practical reference for nursing staff.

Methods: The study used evidence-based methods. Domestic and international databases were searched for articles on perioperative enteral nutrition (tube feeding) in surgical oncology patients. An initial draft of a generalizable management protocol for enteral nutrition tolerance was formulated through evidence synthesis and research team discussion. Two rounds of Delphi consultation were subsequently conducted with experts in enteral nutrition.

Results: A management protocol for perioperative enteral nutrition tolerance in surgical oncology patients was established. The protocol comprised 5 first-level, 12 second-level, and 28 third-level indicators. The expert authority coefficients (Cr) for the two Delphi rounds were 0.857 and 0.880, respectively. The Kendall's concordance coefficients (W) were 0.215 and 0.257 (both $P < 0.05$), indicating significant consensus.

Conclusion: The management protocol for tube-fed enteral nutrition tolerance in perioperative surgical oncology patients showed strong expert consensus and high authority indices. The protocol provides a robust evidence-based foundation for the management of enteral nutrition tolerance in this patient population.

Keywords: enteral nutrition, tube feeding, tolerance management, feeding intolerance, perioperative cancer patients

Introduction

The prevalence of malnutrition among surgical patients is estimated to be between 20 and 60%.¹ A significant proportion of patients with gastrointestinal cancer present with preoperative nutritional risk, and 50–80% are malnourished.² Malnutrition is associated with suboptimal treatment outcomes and adverse clinical consequences (including infections and mortality),³ and represents an independent risk factor for various postoperative complications in patients with cancer.^{4,5} Advancements in Enhanced Recovery After Surgery (ERAS) protocols and nutritional therapy have led to increased attention on early perioperative enteral nutrition (EN) for oncology patients. However, feeding intolerance (FI) remains a frequent complication associated with EN administration, often occurring during the initial feeding phase and being particularly pronounced in tube-fed patients.

Current literature on EN tolerance has focused predominantly on the critical care setting. In 2016, the American Society for Parenteral and Enteral Nutrition (ASPEN) recommended daily assessment of EN tolerance in critically ill



patients.⁶ However, many assessment strategies are unsuited for perioperative patients in general wards. Critically ill patients often have conditions such as endotracheal intubation, high aspiration risk, and inability to express themselves. Healthcare professionals must utilize monitoring devices for precise assessments, such as using ultrasound equipment for gastric residual volume (GRV) monitoring and intra-abdominal pressure measurement,^{7,8} to avoid high incidences of aspiration, elevated GRV, increased intra-abdominal pressure, and diarrhea. However, in surgical wards, perioperative tumor patients generally have stabilized conditions, the goal of enteral nutrition here is to meet their perioperative metabolic needs, reduce stress on the digestive system, promote rapid recovery and discharge.^{3,9} Patients are conscious, able to express discomfort such as abdominal pain, bloating, choking, or diarrhea, and nurse-patient communication is better than in the ICU. Additionally, surgical wards typically lack ultrasound equipment, and nursing staff have not received systematic training for its use. Therefore, based on these considerations, we need to develop a specialized and standardized enteral nutrition protocol for perioperative cancer patients.

Nurses are the primary executors and observers of EN delivery. Currently, there is no standardized protocol for proactive assessment, judgment, and intervention regarding EN tolerance in surgical wards, and nurses often rely on experience. Compounding this issue is the implementation of “hospital-wide unified bed management systems” in many Chinese hospitals. This aims to enhance operational efficiency, enabling rapid turnover in surgical wards. However, some departments (such as orthopedics), due to their specialty characteristics, have historically had limited experience in tube feeding for enteral nutrition and insufficient training in nutritional support, making it difficult to manage feeding tolerance. This poses significant challenges when admitting patients across specialties. Therefore, the surgical nursing system requires a universal and homogeneous enteral nutrition tolerance management protocol to guide clinical practice.

To address this gap, the study used an evidence-based approach combined with the Delphi method. Published studies on perioperative EN in surgical oncology patients were systematically analyzed and the evidence synthesized, leading to the identification and refinement of key nursing interventions for managing EN tolerance. Subsequent application of the Delphi consensus technique resulted in the development of a management protocol for EN tolerance specifically designed for use by nursing staff in surgical wards. This protocol aims to provide systematic, evidence-based guidance for nurses across various wards, ultimately seeking to improve EN tolerance in perioperative surgical patients and hasten patient recovery.

Methods

Establishment of the Evidence-Based Research Team

The team responsible for searching the literature comprised four members: one head nurse (master’s degree, trained in evidence-based practice), one master’s degree candidate (trained in evidence-based practice), and two staff nurses (with bachelor’s degrees, who had received evidence-based training). This team was responsible for the retrieval and appraisal of suitable publications, development of the expert consultation questionnaire, and the collation and analysis of the expert feedback. Additionally, a seven-member appraisal panel was formed, comprising two clinicians (PhD, Associate Chief Physicians), one dietitian (master’s degree, Attending Physician), two head nurses (one Deputy Chief Nurse and one Nurse-in-Charge), and two staff nurses (both Nurses-in-Charge, and certified Clinical Nutrition Specialty Nurses). This panel was responsible for refining the expert consultation questionnaire.

Evidence-Based Research Process

Evidence Retrieval

The “6S” evidence pyramid model was used to guide searches of relevant guideline repositories,¹⁰ published in either Chinese or English, as well as professional society websites, and the UpToDate clinical decision support resource. The databases searched included the Joanna Briggs Institute (JBI) Evidence-Based Practice Database, the Cochrane Library, DynaMed (evidence-based clinical decision support), the Registered Nurses’ Association of Ontario (RNAO) Guidelines, WHO Guidelines, the National Guideline Clearinghouse (NGC), Web of Science, PubMed, CNKI, Wanfang, VIP, and the Chinese Biomedical Literature Database (CBM). A combination of Medical Subject Headings (MeSH) and free-text terms was used. The references of the identified articles were also searched manually. Search strategy: (“neoplasm*” OR “tumor”

OR “cancer”) AND (“enteral nutrition” OR “nutrition therapy” OR “nutritional support” OR “artificial feeding” OR “enteral feeding” OR “force feeding*” OR “tube feeding” OR “gastric feeding tube*”) AND (“feeding intolerance” OR “feeding tolerance”). The search period covered the interval between database inception to December 10, 2023.

Inclusion and Exclusion Criteria

The evidence-based question was formulated using the PIPPOST framework:¹¹

P (Population): Surgical oncology patients receiving tube-fed EN support during the perioperative period.

I (Intervention): Evidence related to improving tube-fed EN tolerance in oncology patients.

P (Professionals): Nursing staff, physicians, and dietitians, among others.

O (Outcomes): Nutritional status, EN tolerance.

S (Setting): Surgical wards.

T (Type of evidence): Guidelines, systematic reviews, evidence summaries, expert consensus statements, randomized controlled trials (RCTs).

Inclusion Criteria: Studies involving adult oncology patients in the perioperative period, interventions related to assessment of EN tolerance and nursing management; evidence types including clinical practice guidelines, evidence summaries, expert consensus statements, systematic reviews, and RCTs; publications in Chinese or English.

Exclusion Criteria: Studies involving ICU patients; Studies not involving tube-fed EN; Duplicate publications or translations; Low-quality studies; No available full text.

Evidence Quality Appraisal

Two members of the research team, systematically trained in evidence-based nursing, independently appraised the quality of the included publications. The appraisal methods were selected based on the Johns Hopkins evidence level and quality guide and the JBI methodology (2016), utilizing specific critical appraisal tools appropriate to the evidence type:

Guidelines: AGREE II instrument.¹²

RCTs, systematic reviews, expert consensus: Corresponding JBI critical appraisal tools.¹³

Evidence summaries: Evidence from the original sources cited in the summaries was retrieved and appraised using the relevant tool based on the appropriate study design.

Following this process, 19 articles were ultimately included: 6 guidelines/expert consensus statements,^{9,14–18} 2 systematic reviews,^{19,20} 1 evidence summary,²¹ and 10 original research studies.^{22–31}

Evidence Synthesis, Grading, and Determination of Recommendation Strength

Two graduate student members of the research team independently extracted the evidence. The extracted evidence was graded using the JBI Evidence-Based Practice Center (2014) evidence level system.⁷ Discrepancies in grading were resolved by consultation with a senior researcher from the appraisal panel to reach a consensus. Evidence levels were assigned from Level 1 (highest) to Level 5 (lowest). Based on the JBI (2014) system, the recommendation strength for each piece of evidence was categorized as Grade A (strong recommendation) or Grade B (weak recommendation).

Development of the Draft Tolerance Management Protocol

To enhance clinical relevance, the research team conducted group discussions. Drawing upon established clinical nursing workflows and the steps involved in nutritional diagnosis and therapy,^{3,4} a tolerance management framework was constructed. This process resulted in an initial draft protocol comprising 5 first-level (nursing workflows), 11 second-level (main work contents), and 31 third-level indicators (specific implementation contents), the detailed breakdown is provided in [Table 1](#).

Delphi Expert Consultation

Selection of Delphi Consultation Experts

The inclusion criteria for consultation experts were as follows: (1) Engagement in EN-related work in surgical wards or nutrition departments for ≥ 10 years; (2) Holding an associate senior or higher clinical medical or nursing technical title;

Table 1 Nursing Management Program for Promoting Tolerance of Enteral Nutrition in Tube-Fed Perioperative Cancer Patients (Draft)

Primary Indicator	Secondary Indicator	Tertiary Indicator
1. Enteral Nutrition Assessment	1.1 Clinical Assessment	Assess primary disease status, gastrointestinal function, aspiration risk, medications, physiological contraindications, and hemodynamic stability before initiating EN.
	1.2 Nutritional Risk Assessment	Assess patient's nutritional risk using the Nutritional Risk Screening 2002 (NRS 2002) scale.
	1.3 Nutrition Timing and Requirements	1.3.1 Provide oral nutritional supplements to malnourished oncology patients prior to major abdominal surgery.
		1.3.2 Initiate EN within 24 hours postoperatively for patients unable to resume adequate oral intake (<50% of needs) lasting >7 days.
1.3.3 Recommended total energy intake is approximately 25–30 kcal/(kg d).		
2. Safety Nursing Measures	2.1 Basic Requirements	2.1.1 Equipment Preparation: Ensure nutritional pump, warming device, and nutrition formula are ready and functional.
		2.1.2 Nurse Preparation: Verify medical orders, prepare all items, maintain aseptic technique, prepare formula immediately before use, and communicate the importance of enteral nutrition and tube care to the patient or family.
		2.1.3 Prepared enteral nutrition formula should not be stored at room temperature for more than 4 hours. If storage exceeds 4 hours, refrigerate. Discard any unused formula after 24 hours. Ready-to-use formulas should be stored according to product instructions.
		2.1.4 Use dedicated EN administration sets and medication labels. Hang separately to distinguish from intravenous therapy.
		2.1.5 Secure and maintain feeding tubes per protocol; record insertion depth, flushing, and clamping.
		2.1.6 Replace nutrition bags, feeding tubes, and formula containers every 24 hours.
	2.2 Quality Control	2.2.1 Establish a multidisciplinary nutritional support team and adopt a process-oriented management approach.
		2.2.2 Establish EN-sensitive indicators for quality control throughout the process.
3. Nursing Measures to Improve Tolerance	3.1 Infusion Care	3.1.1 The osmolality of the nutrition formula is recommended to be <330 mOsm/L. For patients with advanced gastrointestinal tumors, short-peptide-based EN suspensions are preferred.
		3.1.2 Select the appropriate feeding route based on the surgical procedure and duration of nutrition. Post-pyloric feeding is recommended for patients with feeding intolerance or high aspiration risk.
		3.1.3 Perform gastric residual volume measurements regularly.
		3.1.4 Use an EN pump for continuous infusion. Start at a low rate, generally 20–50 mL/h, for post-pancreatic surgery, start at 10–20 mL/h. Gradually increase based on tolerance, max rate ≤150 mL/h.
		3.1.5 For intermittent bolus feeding, increase the enteral nutrition volume incrementally. Start with an initial volume of 100 mL, then gradually increase by 25 mL every 2–3 hours.
		3.1.6 Use a dedicated EN pump warmer to maintain formula temperature at 37–40°C.
	3.2 Positioning Care	3.2.1 When feasible postoperatively, position patients with head elevated 30°–45°. Maintain semi-upright position for 30–60 minutes after infusion stops.
		3.2.2 If lowering the head of bed is necessary for procedures, return it to the elevated position as soon as possible afterward.
	3.3 Other Measures	3.3.1 Encourage early postoperative gum chewing to promote bowel function recovery.
		3.3.2 Implement multi-modal health education to improve patient and family adherence to EN.
		3.3.3 Administer nebulized medication as prescribed.
		3.3.4 Provide oral care with normal saline every 6 hours to maintain oral moisture.

(Continued)

Table 1 (Continued).

Primary Indicator	Secondary Indicator	Tertiary Indicator
4. Tolerance Diagnosis and Management	4.1 Tolerance Diagnosis	Primary nurses should assess tolerance every 4–6 hours using the Enteral Nutrition Tolerance Assessment Scale once EN is initiated.
	4.2 Graded Management	4.2.1 Tolerant: Continue EN. Increase rate by 10 mL/h or maintain current rate. Provide symptomatic treatment.
		4.2.2 Partially Tolerant: Continue EN. Decrease rate by 10 mL/h or reduce the infusion speed by 50%. Reassess after 2 hours.
		4.2.3 Intolerant: Temporarily stop EN. Reassess or change the feeding route. Provide symptomatic treatment.
		4.2.4 For patients with intolerance but no gastrointestinal obstruction, prokinetic agents (eg, erythromycin, metoclopramide) may be used.
5. Evaluation	Evaluation Indicators	72-hour enteral nutrition attainment rate, time to achieve target feeding volume, incidence of intolerance symptoms, anthropometric measurements, and laboratory test parameters.

(3) Possession of research experience; (4) Motivated and willing to participate in the consultation process. Fifteen experts meeting these criteria were enrolled in the study.

Development of Consultation Questionnaire

The Delphi consultation questionnaire was based on the preliminary protocol, included the following sections: (1) Instructions for completion; (2) Main section: Assessment of the importance of each indicator at all levels using a 5-point Likert scale (1 = “not important” to 5 = “very important”). Experts could offer comments and suggestions in the remarks column for each indicator; (3) Expert demographic information, including education, position, professional title, and years of experience; (4) Expert familiarity was assessed using a 5-level scale ranging from “Unfamiliar” to “Very Familiar,” with corresponding scores of 0.1, 0.3, 0.5, 0.7, and 0.9. The basis for expert judgment was categorized into four aspects: theoretical analysis, practical experience, intuitive perception, and reference to domestic and international literature. The degree of influence from each aspect was further classified into three levels: low, medium, and high. The specific scoring assignments are detailed in [Table 2](#).

Implementation of the Delphi Consultation

Two rounds of electronic Delphi consultations were conducted, the response period was 2 weeks. After the return of the first-round questionnaires, the indicators were screened and modified based on the expert feedback. The screening criteria used were an importance score ≥ 4 and a coefficient of variation (CV) < 0.25 . The second-round questionnaire was developed after integration of the feedback. The expert opinions in the second round tended to converge, concluding the consultation process.

Table 2 Expert Judgment Basis Assignment Table

Judgment Basis	Degree of Influence		
	High	Medium	Low
Practical Experience	0.5	0.4	0.3
Theoretical Analysis	0.3	0.2	0.1
Reference to Domestic/International Literature	0.1	0.1	0.1
Intuitive Perception	0.1	0.1	0.1

Statistical Analysis

Statistical analyses were performed using SPSS software version 26.0 (IBM Corp., Armonk, NY, USA). Categorical data are presented as frequencies and percentages, while continuous data are shown as mean \pm standard deviation (SD). Expert enthusiasm was measured by the questionnaire response rate and the expert opinion proposal rate. A response rate $\geq 70\%$ was considered indicative of high enthusiasm. Expert authority was assessed based on the experts' self-reported judgment basis and familiarity with the consultation content. The expert authority coefficient (Cr) was calculated as: $Cr = (Ca + Cs)/2$, where Ca represents the judgment coefficient and Cs represents the familiarity coefficient. Generally, $Cr \geq 0.70$ is considered acceptable. The degree of consensus among experts was assessed using Kendall's coefficient of concordance (W) and coefficient of variation (CV). *P*-values < 0.05 obtained in statistical tests were considered statistically significant.

Results

Expert Demographics

The 15 experts consulted in this study included surgeons (4), surgical nursing administrators (6), nurses specialized in clinical nutrition (3), and dietitians (2). The mean age of the experts was 41.40 ± 4.58 years, with a mean working experience of 22.33 ± 6.07 years. The professional and educational backgrounds of the experts are shown in Table 3.

Expert Enthusiasm and Authority Level

The experts demonstrated high enthusiasm, with a 100% response rate for both rounds of the consultation questionnaire. In the first round, the expert opinion submission rate was 46.67%. The experts' basis of judgment was 0.973, their familiarity level was 0.740, resulting in an expert authority coefficient of 0.857. In the second round, the expert opinion submission rate was 26.67%, the basis of judgment remained at 0.973, the familiarity level increased to 0.786, leading to an expert authority coefficient of 0.880. The experts had a high level of certainty regarding the items, with the authority coefficient for both rounds of consultation being ≥ 0.7 , indicating that the results are reliable.

Degree of Coordination in Expert Opinions

The results from the first round of expert consultation showed that the third-level indicator "Administer nebulization as prescribed" had a CV of 0.25, while the CV values for all other indicators were from 0 to 0.24, and the Kendall's coefficient of concordance W for the importance of expert opinions was 0.215 ($P = 0.001$). After the second round of expert consultation, the CV ranged from 0 to 0.19, and the Kendall's W for the importance of expert opinions was 0.257 ($P = 0.000$). As the number of consultation rounds increased, the degree of coordination among expert opinions improved.

Table 3 Distribution of Participants by Education Level and Professional Title

Category	Subcategory	Number (n)	Percentage (%)
Education Level	Bachelor's Degree	4	26.67
	Master's Degree	5	33.33
	Doctoral Degree	6	40.00
Professional Title	Associate Chief Nurse	6	40.00
	Chief Nurse	3	20.00
	Associate Chief Physician	4	26.67
	Chief Physician	2	13.33

Expert Consultation Outcomes

First Round

Following the first round of expert consultation, the mean importance scores and CV for each indicator are presented in Table 4. According to the predetermined indicator selection criteria, two third-level indicators were deleted: “3.3.3 Administer nebulization as prescribed” (Mean score=4.00, CV=0.25) and “3.3.4 Provide saline mouth rinses every 6 hours to maintain oral moisture” (Mean score=3.86). Experts also commented that these two indicators had weak

Table 4 Nursing Management Program for Promoting Tolerance of Enteral Nutrition in Tube-Fed Perioperative Cancer Patients (First Round of Delphi Survey)

Indicator	Importance Score	Coefficient of Variation	Expert Opinion	Handling of Expert Opinion
1. Enteral Nutrition Assessment	4.93 ± 0.26	0.05		
1.1 Clinical Assessment	5	<0.01		
Assess primary disease condition, gastrointestinal function, aspiration risk, medications, physiological contraindications, hemodynamics, etc., before initiating EN.	5			
1.2 Nutritional Risk Assessment	4.93 ± 0.26	0.05	Subjective and objective indicators should be used for a comprehensive nutritional status assessment.	Accepted
Assess patient's nutritional risk using the Nutritional Risk Screening 2002 (NRS 2002) scale.	4.71 ± 0.61	0.13		
1.3 Nutrition Timing and Requirements	4.80 ± 0.41	0.09		
1.3.1 Provide oral nutritional supplements to malnourished oncology patients prior to major abdominal surgery.	5	<0.01		
1.3.2 Initiate EN within 24 hours postoperatively for patients unable to resume adequate oral intake (<50% of needs) lasting >7 days.	5	<0.01		
1.3.3 The recommended total energy intake is approximately 25–30 kcal/(kg·d).	4.54 ± 0.88	0.19	The expression is inaccurate; refer to guidelines.	Accepted
2. Safety Nursing Measures	4.86 ± 0.35	0.07		
2.1 Basic Requirements	4.93 ± 0.26	0.05	The readiness of the formula and suggestions for its storage can be integrated; content can be streamlined, focusing mainly on tolerance-related factors.	Accepted
2.1.1 Equipment preparation: Ensure the feeding pump, warming device, and nutritional formula are ready for use.	4.93 ± 0.26	0.05		
2.1.2 Nurse preparation: Verify medical orders, prepare all items, maintain aseptic technique, prepare immediately before use, and explain the importance of enteral nutrition and tube care to the patient or family.	4.93 ± 0.26	0.05		
2.1.3 Prepared enteral nutrition formula should not be stored at room temperature for more than 4 hours. If exceeding 4 hours, refrigerate. Discard if not used within 24 hours. Commercial enteral nutrition formulas should be stored according to product instructions.	4.93 ± 0.26	0.05		
2.1.4 Use dedicated EN administration sets and medication labels. Hang separately to distinguish from intravenous therapy.	4.93 ± 0.26	0.05		
2.1.5 Secure and maintain feeding tubes per protocol; record insertion depth, flushing, and clamping.	4.80 ± 0.41	0.09		
2.1.6 Replace nutrition bags, feeding tubes, and formula containers every 24 hours.	4.93 ± 0.26	0.05		
2.2 Quality Control	4.86 ± 0.35	0.07		
2.2.1 Establish a multidisciplinary nutrition support team and adopt a process management approach.	4.87 ± 0.65	0.13		
2.2.2 Establish enteral nutrition-sensitive indicators for whole-process quality control.	4.87 ± 0.65	0.13		

(Continued)

Table 4 (Continued).

Indicator	Importance Score	Coefficient of Variation	Expert Opinion	Handling of Expert Opinion
3. Nursing Measures to Improve Tolerance	4.86 ± 0.35	0.07		
3.1 Infusion Care	5	<0.01		
3.1.1 The osmolality of the nutritional formula is recommended to be <330 mOsm/L. For patients with advanced digestive system tumors, short-peptide enteral nutrition suspensions are preferable.	4.93 ± 0.26	0.05	It is suggested to add: select an appropriate nutritional formula for the patient.	Accepted
3.1.2 Select the appropriate feeding route based on the surgical procedure and expected duration of nutrition. Post-pyloric feeding is recommended for patients with feeding intolerance or high aspiration risk.	5	<0.01		
3.1.3 Perform gastric residual volume measurements regularly.	4.33 ± 1.05	0.24	Gastric residual volume monitoring is currently controversial and can be considered based on ward conditions.	Recommended for deletion after group discussion, not as a routine operation.
3.1.4 Use an EN pump for continuous infusion. Start at a low rate, generally 20–50 mL/h, for post-pancreatic surgery patients, start at 10–20 mL/h. Gradually increase based on tolerance, max rate ≤150 mL/h.	4.73 ± 0.46	0.10	Change to: Recommend using a feeding pump for continuous infusion.	Accepted
3.1.5 For intermittent bolus feeding, increase the enteral nutrition infusion volume incrementally. The initial infusion volume is 100 mL, gradually increasing by 25 mL every 2–3 hours.	4.33 ± 1.05	0.24		
3.1.6 Use a dedicated EN pump warmer to maintain formula temperature at 37–40°C.	4.47 ± 1.08	0.24		
3.2 Positioning Care	4.80 ± 0.41	0.09		
3.2.1 When feasible postoperatively, position patients with head elevated 30°–45°. Maintain semi-upright position for 30–60 minutes after infusion stops.	5	<0.01		
3.2.2 If lowering the head of bed is necessary for procedures, return it to the elevated position as soon as possible afterward.	4.87 ± 0.35	0.07		
3.3 Other	4.73 ± 0.46	0.10		
3.3.1 Encourage early postoperative gum chewing to promote bowel function recovery.	4.87 ± 0.35	0.07		
3.3.2 Implement multi-modal health education to improve patient and family adherence to EN.	4.80 ± 0.41	0.09		
3.3.3 Administer nebulized inhalation as prescribed.	4.00 ± 1.00	0.25	The nebulization and mouth rinse care procedures have a weak association with enteral nutrition tolerance; suggested deletion.	Accepted
3.3.4 Provide oral care with normal saline every 6 hours to maintain oral moisture.	3.86 ± 0.74	0.19		
4. Tolerance Diagnosis and Management	4.67 ± 0.49	0.10		
4.1 Tolerance Diagnosis	4.93 ± 0.26	0.05		
Primary nurses should assess tolerance every 4–6 hours using the Enteral Nutrition Tolerance Assessment Scale once EN is initiated.	4.93 ± 0.26	0.05		
4.2 Graded Management	4.62 ± 0.65	0.14		
4.2.1 Tolerant: Continue EN, increase rate by 10 mL/h or maintain current rate. Provide symptomatic treatment.	4.80 ± 0.41	0.09		
4.2.2 Partially tolerant: Continue EN, decrease rate by 10 mL/h or reduce the infusion speed by 50%, and reassess after 2 hours.	4.73 ± 0.59	0.12		
4.2.3 Intolerant: Temporarily stop enteral nutrition, reassess or change the feeding route. Provide symptomatic treatment.	4.67 ± 0.82	0.18		

(Continued)

Table 4 (Continued).

Indicator	Importance Score	Coefficient of Variation	Expert Opinion	Handling of Expert Opinion
4.2.4 For patients with intolerance but no gastrointestinal obstruction, prokinetic agents (eg, erythromycin, metoclopramide) may be used.	5	<0.01	Suggest adding content on Traditional Chinese Medicine treatment.	Accepted
5. Evaluation	5	<0.01		
Evaluation indicators	4.80 ± 0.41	0.09		
72-hour enteral nutrition attainment rate, time to achieve target feeding volume, incidence of intolerance symptoms, anthropometric measurements, and laboratory test parameters.	4.80 ± 0.41	0.09	Suggest listing evaluation indicators in detail.	Accepted

relevance to enteral nutrition tolerance management and recommended their removal. This recommendation was adopted after discussion by the research team. Two experts disagreed on “Gastric residual volume measurement” routines in general wards, after further review of the literature and consultation with physicians, this recommendation was modified.

A total of six items were revised. For instance, experts noted that “1.2 nutritional risk assessment” alone could not comprehensively reflect a patient’s nutritional status. They recommended adopting a holistic assessment utilizing both subjective and objective indicators. After reviewing the literature and group discussion, this item was revised to “Assess nutritional status using tools such as the Nutritional Risk Screening 2002 (NRS-2002), the Patient-Generated Subjective Global Assessment (PG-SGA), anthropometric measurements, and laboratory tests”. Three experts suggested that the description “Energy assessment 25–30 kcal/kg/day” was inaccurate, further literature review led to its revision to reflect the recommendations of guidelines “Postoperative total energy requirement 25–30 kcal/kg/day; Protein target 1.5–2.0 g/kg/day; Achieved gradually over 5–7 days based on tolerance.”^{3,6,9}

Three items (2.1.1–2.1.3) were suggested for consolidation, retaining only content directly relevant to tolerance management. This suggestion was adopted following group discussion. The consolidated item was revised to “Store EN formula correctly, use aseptic technique, and prepare immediately before use.”¹⁸ For the final item of the protocol, experts recommended detailing the specific evaluation indicators. This recommendation was adopted after the research team conducted a literature review and discussion. The supplemented details are provided in the evaluation section of Table 5.

Three items were added during this revision. Four experts suggested incorporating “select an appropriate nutritional formula” into the infusion care content. Following a literature review, this was refined to “Select an appropriate nutritional formula with an osmolality less than 330 mOsm/L.”^{6,22} Separately, two experts recommended adding “blood glucose monitoring” to the “other measures” section, citing its relevance to tolerance management. Another expert noted the beneficial effects of Traditional Chinese Medicine (TCM) in improving tolerance. These suggestions were incorporated after team discussion.

Second Round

Following these adjustments, we developed a management protocol comprising 5 first-level indicators, 12 second-level indicators, and 28 third-level indicators. This revised program was then subjected to a second round of Delphi consultation, the results of which are presented in Table 5. All indicators met the retention criteria, with importance scores greater than 4 and CV less than 0.25, therefore, all were retained.

One expert suggested changing “Nursing measures to improve tolerance” to the more concise “Preventive measures” which was adopted. Another expert pointed out the critical importance of selecting appropriate nutritional formulas, suggesting specific recommendations regarding concentration and type. Therefore, item 3.1.1 was revised to “select appropriate enteral formulations according to clinical indications, with an osmolality less than 330 mOsm/L. For individuals with gastrointestinal impairment or malabsorption, amino acid-based or short peptide-based nutritional formulas can be considered, cancer patients may benefit from immunonutrition”.^{6,14}

Table 5 Nursing Management Program for Promoting Tolerance of Enteral Nutrition in Tube-Fed Perioperative Cancer Patients (Second Round of Delphi Survey)

Level 1 Indicator	Level 2 Indicator	Level 3 Indicator	Importance Assignment (mean±SD, points)	Coefficient of Variation (CV)	Evidence Level	Recommendation Grade
1. EN Assessment	1.1 Clinical Assessment	Assess primary disease status, gastrointestinal function, aspiration risk, medications, physiological contraindications, and hemodynamic stability before initiating EN.	4.93±0.26	0.05	5b	A
	1.2 Nutritional Assessment	Assess nutritional status using tools such as NRS-2002, PG-SGA, anthropometric measurements, and laboratory tests.	4.73±0.46	0.10	1a	A
	1.3 Nutrition Timing & Requirements	1.3.1 Provide oral nutritional supplements to malnourished oncology patients prior to major abdominal surgery.	4.53±0.52	0.11	5a	A
		1.3.2 Initiate EN within 24 hours postoperatively for patients unable to resume adequate oral intake (<50% of needs) lasting >7 days.	5.00	<0.01	5a	A
		1.3.3 Postoperative target total energy intake: 25–30 kcal/(kg·d); Protein target: 1.5–2.0 g/kg/d. Achieve targets gradually over 5–7 days based on tolerance.	4.53±0.52	0.11	5a	A
2. Safety Nursing Measures	2.1 Basic Requirements	2.1.1 Store EN formula correctly; use aseptic technique; prepare immediately before use.	5.00	<0.01	5b	A
		2.1.2 Use dedicated EN administration sets and medication labels; hang separately to distinguish from intravenous therapy.	4.87±0.35	0.07	5b	A
		2.1.3 Secure and maintain feeding tubes per protocol; record insertion depth, flushing, and clamping.	4.87±0.35	0.07	5b	A
		2.1.4 Change EN feeding bags, tubing, and containers every 24 hours.	5.00	<0.01	5b	A
	2.2 Quality Control	2.2.1 Establish a multidisciplinary Nutrition Support Team (NST) and implement process-oriented management.	5.00	<0.01	1d	A
		2.2.2 Establish EN-sensitive indicators for quality control throughout the process.	4.93±0.26	0.05	5c	B
3. Preventive Nursing Measures	3.1 Infusion Care	3.1.1 select appropriate enteral formulations according to clinical indications, with an osmolality less than 330 mOsm/L. For individuals with gastrointestinal impairment or malabsorption, amino acid-based or short peptide-based nutritional formulas can be considered; cancer patients may benefit from immunonutrition.	4.73±0.52	0.11	5a	A
		3.1.2 Select the appropriate feeding route based on surgery type and expected duration. Post-pyloric feeding is recommended for patients with feeding intolerance or high aspiration risk.	5.00	<0.01	5a	A
		3.1.3 Use an infusion pump for continuous administration. Start infusion at a low rate (generally 20–50 mL/h; 10–20 mL/h post-pancreatic surgery). Increase gradually based on tolerance; max rate ≤150 mL/h.	4.87±0.35	0.07	5a	A

(Continued)

Table 5 (Continued).

Level 1 Indicator	Level 2 Indicator	Level 3 Indicator	Importance Assignment (mean±SD, points)	Coefficient of Variation (CV)	Evidence Level	Recommendation Grade	
		3.1.4 For intermittent bolus feeding, increase infusion volume incrementally. Initial bolus: 100 mL; increase by 25 mL every 2–3 hours.	4.47±0.83	0.19	2c	B	
		3.1.5 Use an EN-specific feeding tube warmer; maintain formula temperature at 37–40°C.	4.47±0.83	0.19	5b	A	
	3.2 Positioning Care	3.2.1 When feasible postoperatively, position patients with head elevated 30°–45°. Maintain semi-upright position for 30–60 minutes after infusion stops.	5.00	<0.01	1a	A	
		3.2.2 If lowering the head of bed is necessary for procedures, return it to the elevated position as soon as possible afterward.	4.87±0.35	0.19	5c	B	
	3.3 Other Measures	3.3.1 Blood Glucose Monitoring: Monitor blood glucose and maintain ≤10 mmol/L before starting or during EN; monitor every 4–6 hours.	4.53±0.52	0.11	3c	B	
		3.3.2 Encourage early postoperative gum chewing to promote bowel function recovery.	4.67±0.49	0.10	1c	A	
		3.3.3 Implement multi-modal health education to improve patient and family adherence to EN.	4.93±0.26	0.05	2c	B	
	4. Tolerance Diagnosis & Graded Management	4.1 Tolerance Diagnosis	Primary nurses should assess tolerance every 4–6 hours using the Enteral Nutrition Tolerance Assessment Scale once EN is initiated.	4.80±0.41	0.09	1d	A
		4.2 Graded Management	4.2.1 Tolerant: Continue EN; increase rate by 10 mL/h or maintain current rate; provide symptomatic treatment.	4.93±0.26	0.05	5b	B
4.2.2 Partially Tolerant: Continue EN; decrease rate by 10 mL/h or reduce current rate by 50%; re-assess tolerance after 2 hours.			5.00	<0.01	5b	B	
4.2.3 Intolerant: Temporarily suspend EN; re-assess or change feeding route; provide symptomatic treatment.			5.00	<0.01	5b	B	
4.3 Comprehensive Therapy		4.3.1 Pharmacotherapy: For intolerance without gastrointestinal obstruction, consider prokinetic agents (eg, Erythromycin, Metoclopramide).	5.00	<0.01	1a	A	
		4.3.2 Traditional Chinese Medicine (TCM) Care: Apply TCM modalities based on individual patient pattern differentiation (eg, herbal decoctions like Si Jun Zi Tang or Bu Yi San Jie Yin, holographic scraping therapy, moxibustion at Zusanli ST36).	4.80±0.41	0.09	1b	B	
5. Evaluation	Evaluation Indicators	Use the following to evaluate: 72-hour EN adequacy rate; time to achieve target feeding volume; incidence of intolerance symptoms (eg, abdominal pain, diarrhea, bloating); anthropometric measurements (eg, BMI, calf circumference, mid-upper arm circumference, triceps skinfold thickness); laboratory parameters (eg, albumin, prealbumin, transferrin).	4.80±0.41	0.09	1d	B	

After two rounds of expert consultation and subsequent data analysis, a management protocol for perioperative tube-fed EN tolerance in surgical oncology patients was finalized.

Discussion

FI is a common complication encountered during enteral nutrition. The concept of the feeding intolerance syndrome was explicitly defined in 2012 by the Abdominal Problems Working Group of the European Society of Intensive Care Medicine.³² While research on tolerance-related issues has focused predominantly on critically ill patients in the ICU, these procedures are not fully applicable to general wards. In surgical nursing practice, oncology patients face a high risk of perioperative malnutrition. Early initiation of EN is recommended particularly for patients undergoing major surgery,^{3,9} with tube feeding being the primary route. The effective management of tolerance is therefore crucial for this population. However, there have been no standardized, homogeneous tolerance management protocol for use in surgical settings. This gap is especially significant in the context of the “hospital-wide unified bed management systems” model, in which nurses care for patients across various surgical specialties. The development of a universal management protocol for EN tolerance would provide an evidence-based foundation for cross-specialty nursing care within the surgical system.

Evidence-Based Nature of the Management Protocol for Perioperative Enteral Nutrition Tolerance in Surgical Oncology Patients

This study aimed to address the issue of tube-fed EN tolerance in perioperative surgical oncology patients. Following evidence-based research steps, the PIPOST framework was used for formulation of the clinical question.¹¹ This was followed by the systematic analysis and synthesis of clinical practice guidelines and the primary literature. The quality of evidence was rigorously appraised using established standards, namely, the AGREE II instrument for guidelines,¹² JBI critical appraisal tools for systematic reviews and randomized controlled trials, and the JBI tool for expert consensus.¹³ This process ensured the inclusion of high-quality evidence, thereby guaranteeing the scientific rigor of the research content.

Clinical Applicability of the Management Protocol for Perioperative Enteral Nutrition Tolerance in Surgical Oncology Patients

Expert Selection and Validation

The clinical relevance of the protocol was ensured through the selection of a multidisciplinary expert panel intimately involved in clinical EN delivery, including surgeons, surgical nursing managers, nutrition physicians, and clinical nurses, representing a well-balanced composition. Two rounds of Delphi consultation and multidisciplinary validation were used for further refinement of the protocol to align closely with clinical practice. The results indicated questionnaire response rates of 100% for both rounds, with authority coefficients (Cr) of 0.857 and 0.880, respectively. The Kendall's harmony coefficients for the importance of each item were statistically significant ($P < 0.05$), demonstrating high expert engagement, authority, and consensus consistency, thereby affirming the reliability of the program.

Structure and Process

The development of first-level indicators integrated established clinical nursing workflows and clinical nutrition care pathways.^{3,4} This approach aligns with clinical reasoning^{3,4} and enables planned, continuous, and systematic management of enteral nutrition tolerance in perioperative patients.

The Tolerance Management Protocol Demonstrates Both Generalizability and Professional Value for Surgical Nurses

This tolerance management protocol is specifically designed for perioperative oncology patients within the surgical system. The primary objectives of enteral nutrition in this context are to meet perioperative metabolic demands, reduce gastrointestinal stress, provide nutritional supplementation, and facilitate enhanced recovery, as outlined in key surgical and nutritional guidelines.^{3,9,16} These goals differ from those guiding enteral nutrition management in critically ill patients. Furthermore, within the surgical system itself, heterogeneity exists across different specialties. Variations in the

knowledge training nurses receive, daily operational protocols, and levels of interprofessional collaboration regarding enteral nutrition care can pose significant limitations to the implementation of hospital-wide unified bed management and the provision of cross-specialty nursing. This study addresses this gap by developing a universally applicable and operationally practical tolerance management protocol.

First, assessment is fundamental to clinical decision-making and patient care safety.³³ During the first Delphi round, the initial focus on nutritional risk screening was broadened to include a comprehensive nutritional assessment. This expansion encompassed indicators such as NRS2002, PG-SGA, BMI and laboratory parameters.^{3,9} The aim was to facilitate nurses' effective participation in Nutrition Support Team (NST) activities by ensuring they thoroughly understand the patient's baseline condition prior to initiating enteral nutrition. By the second Delphi round, the importance score for this revised indicator increased to 4.73, with a decreased CV of 0.10.

Regarding GRV measurement, although elevated GRV can be an indicator of enteral feeding intolerance, its application in routine intolerance assessment remains controversial. A systematic review incorporating five randomized controlled trials found no statistically significant difference in the incidence of pneumonia or aspiration between critically ill patients managed with or without GRV monitoring.³⁴ Consequently, following team discussion, routine GRV monitoring was not included as a standard indicator for tolerance assessment in general wards and was removed from the protocol. It is acknowledged that GRV measurement remains necessary in clinical practice for critically ill patients or those requiring long-term enteral nutrition.¹⁷

Concerning preventive measures for intolerance, the indicator "blood glucose monitoring" was added after the first Delphi round. This addition reflects expert recognition of glycemic control as a significant factor influencing tolerance. A literature review revealed that blood glucose levels >10 mmol/L or persistent hyperglycemia are associated with a higher risk of

Intolerance.³⁵ The Canadian Critical Care Nutrition Guidelines further recommend avoiding hyperglycemia (>10 mmol/L) during nutritional support.³⁶ Given its clinical significance and ease of implementation, this measure was incorporated into the protocol following team discussion, thereby enriching the management strategies compared to traditional approaches for post-upper gastrointestinal or esophageal cancer surgery patients.^{19,21}

Regarding tolerance diagnosis, the use of a standardized tolerance scale has been shown to improve EN adequacy rates.^{37,38} The management protocol adopted the Enteral Nutrition Tolerance Assessment Scale from the group standard of the Chinese Nursing Association.¹⁸ This scale recommends proactive nurse assessment every 4–6 hours, facilitating the early detection of intolerance. Critically, the protocol can guide nurses to implement tolerance-level-specific interventions, which is key to promoting tolerance and ensuring the successful delivery of nutrition. For example, a tolerance score of 4 represents partial intolerance, indicating that nurses should decrease the infusion rate by 10 mL/h or reduce the current rate by 50%, with reassessment after 2 hours, accompanied by dynamic adjustments, rather than discontinuing EN.

Regarding the treatment of feeding intolerance, experts suggested incorporating TCM for alleviating intolerance symptoms. Our findings indicate that TCM modalities such as the herbal decoctions Si Jun Zi Tang and Bu Yi San Jie Yin, holographic scraping therapy, and moxibustion at Zusanli (ST36) have demonstrated efficacy.^{20,26,39,40} Following the second round of consultation, this indicator achieved an importance score of 4.8 with a coefficient of variation (CV) of 0.02, indicating its reliability. These findings are more comprehensive than those of previous studies and highlight the unique strengths of TCM.

Conclusions

The management protocol for tube-fed enteral nutrition tolerance in perioperative surgical oncology patients showed strong expert consensus and high authority indices. The conceptual framework and indicators of this management protocol offer practical utility for surgical nursing staff. It can actively guide clinical nurses in the provision of structured tolerance management for tube-fed patients, promoting the appropriate implementation of nutritional therapy within the surgical system and contributing significantly to patient recovery.

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Disclosure

The authors declare no conflicts of interest in this work.

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