

Journal of Parenteral and Enteral Nutrition

<http://pen.sagepub.com/>

A.S.P.E.N. Clinical Guidelines: Nutrition Support of Hospitalized Adult Patients With Obesity

Patricia Choban, Roland Dickerson, Ainsley Malone, Patricia Worthington, Charlene Compher and the American Society for Parenteral and Enteral Nutrition

JPEN J Parenter Enteral Nutr 2013 37: 714 originally published online 23 August 2013

DOI: 10.1177/0148607113499374

The online version of this article can be found at:

<http://pen.sagepub.com/content/37/6/714>

Published by:



<http://www.sagepublications.com>

On behalf of:



American Society for Parenteral
and Enteral Nutrition

The American Society for Parenteral & Enteral Nutrition

Additional services and information for *Journal of Parenteral and Enteral Nutrition* can be found at:

Email Alerts: <http://pen.sagepub.com/cgi/alerts>

Subscriptions: <http://pen.sagepub.com/subscriptions>

Reprints: <http://www.sagepub.com/journalsReprints.nav>

Permissions: <http://www.sagepub.com/journalsPermissions.nav>

>> [Version of Record](#) - Nov 1, 2013

[OnlineFirst Version of Record](#) - Aug 23, 2013

[What is This?](#)

A.S.P.E.N. Clinical Guidelines: Nutrition Support of Hospitalized Adult Patients With Obesity

Patricia Choban, MD¹; Roland Dickerson, PharmD, BCNSP²; Ainsley Malone, MS, RD, CNSC³; Patricia Worthington, MSN, RN⁴; Charlene Compher, PhD, RD, CNSC, LDN, FADA, FASPEN⁵; and the American Society for Parenteral and Enteral Nutrition

Journal of Parenteral and Enteral Nutrition
 Volume 37 Number 6
 November 2013 714–744
 © 2013 American Society for Parenteral and Enteral Nutrition
 DOI: 10.1177/0148607113499374
 jpen.sagepub.com
 hosted at
 online.sagepub.com



Abstract

Background: Due to the high prevalence of obesity in adults, nutrition support clinicians are encountering greater numbers of obese patients who require nutrition support during hospitalization. The purpose of this clinical guideline is to serve as a framework for the nutrition support care of adult patients with obesity. **Method:** A systematic review of the best available evidence to answer a series of questions regarding management of nutrition support in patients with obesity was undertaken and evaluated using concepts adopted from the Grading of Recommendations, Assessment, Development and Evaluation working group. A consensus process, that includes consideration of the strength of the evidence together with the risks and benefits to the patient, was used to develop the clinical guideline recommendations prior to multiple levels of external and internal review and approval by the A.S.P.E.N. Board of Directors. **Questions:** (1) Do clinical outcomes vary across levels of obesity in critically ill or hospitalized non-intensive care unit (ICU) patients? (2) How should energy requirements be determined in obese critically ill or hospitalized non-ICU patients? (3) Are clinical outcomes improved with hypocaloric, high protein diets in hospitalized patients? (4) In obese patients who have had a malabsorptive or restrictive surgical procedure, what micronutrients should be evaluated? (*JPEN J Parenter Enteral Nutr.* 2013;37:714-744)

Keywords

adult; life cycle; calorimetry; nutrition; assessment; outcomes; research/quality; support practice; obesity

Background

As of June 2013, the American Medical Association recognized obesity as a disease that requires medical treatment.^{1,2} Based on the National Health and Nutrition Examination Survey 2009-2010, the prevalence of obesity in the United States is 35.5% in adult men, 35.8% in adult women, including 4.4% and 8.2% respectively with body mass index (BMI) ≥ 40 kg/m².³ Thus, nutrition support clinicians are likely to care for obese patients, particularly during hospital admissions. While nutrition support clinicians care for patients across a broad range of clinical settings, the bulk of publications available for this clinical guideline have come from hospitalized patients. Furthermore, since the clinical acuity of patients admitted to intensive care units (ICUs) is much higher than those who are not critically ill, for this guideline most recommendations have been made separately for these 2 groups of obese hospitalized patients when data were available.

Bariatric surgery is a common treatment for patients who have severe obesity, with estimates of approximately 200,000 adults treated with bariatric surgery annually in the United States.⁴ Since these procedures are designed to limit the patient's nutrient intake as a strategy to promote significant and durable weight loss, patients treated with these procedures may require nutrition care. Thus, the purpose of this clinical

guideline is to guide clinicians on the nutrition support care of hospitalized adult patients who have obesity.

From ¹Mt Carmel Hospital, Central Ohio Surgical Associates, Columbus, OH, USA; ²University of Tennessee Health Science Center, Memphis, TN, USA; ³Department of Pharmacy, Mt Carmel West Hospital, Columbus, OH, USA; ⁴Thomas Jefferson University Hospital, Philadelphia, PA, USA; and ⁵University of Pennsylvania School of Nursing, Philadelphia, PA, USA.

The A.S.P.E.N. Clinical Guidelines Editorial Board guided the development of and review of these guidelines using the GRADE system. The A.S.P.E.N. Board of Directors approved the guidelines on June 26, 2013.

Financial disclosure: None declared.

Speaker's Bureau: Nestlé (RND); Abbott (AM)

Received for publication July 5, 2013; accepted for publication July 5, 2013.

This article originally appeared online on August 23, 2013.

Corresponding Author:

Charlene Compher, PhD, RD, CNSD, LDN, FADA, FASPEN, Professor of Nutrition Science, University of Pennsylvania School of Nursing, Claire M. Fagin Hall, 418 Curie Blvd, Philadelphia, PA 19104-4217, USA.

Email: compher@nursing.upenn.edu

Method

The American Society for Parenteral and Enteral Nutrition (A.S.P.E.N.) is an organization comprised of healthcare professionals representing the disciplines of medicine, nursing, pharmacy, dietetics, and nutrition science. The mission of A.S.P.E.N. is to improve patient care by advancing the science and practice of clinical nutrition and metabolism. A.S.P.E.N. vigorously works to support quality patient care, education, and research in the fields of nutrition and metabolic support in all healthcare settings. These clinical guidelines were developed under the guidance of the A.S.P.E.N. Board of Directors. Promotion of safe and effective patient care by nutrition support practitioners is a critical role of the A.S.P.E.N. organization. A.S.P.E.N. has been publishing clinical guidelines since 1986.⁵⁻¹⁵

These A.S.P.E.N. clinical guidelines are based on general conclusions of health professionals who, in developing such guidelines, have balanced potential benefits to be derived from a particular mode of medical therapy against certain risks inherent with such therapy. However, the professional judgment of the attending health professional is the primary component of quality medical care. Because guidelines cannot account for every variation in circumstances, the practitioner must always exercise professional judgment in their application. These clinical guidelines are intended to supplement, but not replace, professional training and judgment.

A.S.P.E.N. clinical guidelines has adopted concepts of the Grading of Recommendations, Assessment, Development and Evaluation (GRADE) working group.¹⁶⁻¹⁹ A full description of the methodology has been published.²⁰ Briefly, specific clinical questions where nutrition support is a relevant mode of therapy are developed and key clinical outcomes are identified. A rigorous search of the published literature is conducted, each included study assessed for research quality, tables of findings developed, the body of evidence for the question evaluated and graded. Randomized controlled clinical trials are initially graded as strong evidence, but may be downgraded in quality based on study limitations. Controlled observational studies are initially graded as weak evidence, but may be graded down further based on study limitations or upgraded based on study design strengths. In a consensus process, the authors make recommendations for clinical practice that are based on the evidence review assessed against consideration of the risks and benefits to patients. Recommendations are graded as strong when the evidence is strong and/or the risk vs benefit analysis is strong. Weak recommendations may be based on weaker evidence and/or weaker trade-offs to the patient. When limited research is available to answer a question, the recommendation is for further research to be conducted.

The guideline authors represent a range of academic and clinical expertise (medicine, dietetics, nursing, pharmacy). The external and internal expert reviewers, including the A.S.P.E.N. Board of Directors, have a similar breadth of professional expertise. This clinical guideline is planned for revision in 2018.

The questions are summarized in Table 1. With the assistance of a reference librarian a search was conducted in PubMed, EMBASE, and CINAHL on August 1, 2012, and updated May 2, 2013, using inclusion criteria of adult subjects, English language, randomized controlled trials, observational studies, and publications over the past 10 years. Search terms “obesity,” “clinical outcomes,” “mortality,” “infection,” “parenteral nutrition,” and “enteral nutrition” were applied in various combinations for questions 1-3. For question 1, 31 articles met the inclusion criteria. For question 2, 9 articles that described measures in hospitalized or clinical populations of obese patients and that reported data with accuracy and bias rates were included. For question 3, the time limitation was relaxed to obtain all published information on the topic, yielding 8 articles. For question 4, search terms of “copper,” “zinc,” “iron,” “selenium,” “vitamin deficiency,” “nutrient deficiency,” “gastric bypass,” “biliopancreatic diversion,” “vitamin D,” and “bariatric surgery” were used in various combinations with a time limitation of the past 10 years, which yielded 22 articles.

Results

Question 1: Do Clinical Outcomes Vary Across Levels of Obesity in Critically Ill or Hospitalized Non-ICU Patients? (Tables 2-3)

Recommendation

1a. Critically ill patients with obesity experience more complications than patients with optimal BMI levels. Nutrition assessment and development of a nutrition support plan is recommended within 48 hours of ICU admission (strong).

Evidence Grade: Low.

1b. All hospitalized patients, regardless of BMI, should be screened for nutrition risk within 48 hours of admission, with nutrition assessment for patients who are considered at risk (strong).

Evidence Grade: Low.

Rationale. Clinical outcomes in patients with obesity may be impacted by numerous factors, including comorbid conditions, associated metabolic changes and any modifications in clinical care (including nutrition support) that are made on behalf of the obese patient. The available studies comparing outcomes of mortality, length of stay (LOS), and complications in obese ICU and non-ICU patients are limited by their retrospective database evaluation,²¹⁻³⁵ by a relatively small number of obese subjects,^{24-28,36-41} or by overall small sample size.^{22,24-28,31,34,39-43} In particular, mortality outcomes are varied, depending on these factors. To address concerns about limitations in statistical power for the outcome of mortality, we considered the evidence from 8 studies with more than 300 obese subjects. One found increased mortality in obese trauma patients,²¹ 5 reported reduced mortality in mixed ICU types,^{23,35,42,44,45} and 3 reported no difference in mortality.^{29,32,46} LOS in the ICU was not

Table 1. Nutrition Support Clinical Guideline Recommendations in Adult Patients With Obesity.

| Question | Recommendation | Recommendation Grade and Evidence Quality |
|---|---|--|
| 1. Do clinical outcomes vary across levels of obesity in critically ill or hospitalized non-ICU patients? | 1a. Critically ill patients with obesity experience more complications than patients with optimal BMI levels. Nutrition assessment and development of a nutrition support plan is recommended within 48 hours of ICU admission. | Recommendation: Strong Evidence: Low |
| | 1b. All hospitalized patients, regardless of BMI, should be screened for nutrition risk within 48 hours of admission, with nutrition assessment for patients who are considered at risk. | Recommendation: Strong Evidence: Low |
| 2. How should energy requirements be determined in obese critically ill or hospitalized non-ICU patients? | 2a. In the critically ill obese patient, if indirect calorimetry is unavailable, energy requirements should be based on the Penn State University 2010 predictive equation, or the modified Penn State equation if the patient is over the age of 60 years. | Recommendation: Strong Evidence: High |
| | 2b. In the hospitalized obese patient, if indirect calorimetry is unavailable and the Penn State University equations cannot be used, energy requirements may be based on the Mifflin–St Jeor equation using actual body weight. | Recommendation: Weak Evidence: Moderate |
| 3. Are clinical outcomes improved with hypocaloric, high protein diets in hospitalized patients with obesity? | 3a. Clinical outcomes are at least equivalent in patients supported with high protein, hypocaloric feeding to those supported with high protein, eucaloric feeding. A trial of hypocaloric, high protein feeding is suggested in patients who do not have severe renal or hepatic dysfunction. Hypocaloric feeding may be started with 50%-70% of estimated energy needs or < 14 kcal/kg actual weight. High protein feeding may be started with 1.2 g/kg actual weight or 2-2.5 g/kg ideal body weight, with adjustment of goal protein intake by the results of nitrogen balance studies. | Recommendation: Weak Evidence: Low |
| | 3b. Hypocaloric, low protein feedings are associated with unfavorable outcomes. Clinical vigilance for adequate protein provision is suggested in patients who do not have severe renal or hepatic dysfunction. | Recommendation: Weak Evidence: Low |
| 4. In obese patients who have had a malabsorptive or restrictive surgical procedure, what micronutrients should be evaluated? | 4. Patients who have undergone sleeve gastrectomy, gastric bypass, or biliopancreatic diversion ± duodenal switch have increased risk of nutrient deficiency. In acutely ill hospitalized patients with history of these procedures, evaluation for evidence of depletion of iron, copper, zinc, selenium, thiamine, folate, and vitamins B ₁₂ and D is suggested as well as repletion of deficiency states. | Recommendation: Weak Evidence: Low |

ICU, intensive care unit.

significantly different in obese than nonobese subjects in the single large study reporting this outcome.⁴⁵ Studies with more than 300 obese patients reported more complications in obese than nonobese patients,^{25,47} as did 3 smaller studies in trauma patients.^{33,37,48} One large study in patients admitted to the medical ICU observed no difference in complications in obese than nonobese patients.³² These complications may impact adjunctive nutrition care and thus support our consensus that an early nutrition assessment (as for all critically ill patients) and care plan is indicated.

In the hospitalized, non-critically ill obese patient, 2 studies had more than 300 obese patients. One of these in surgical patients reported lower mortality and hospital

LOS,³⁰ while a study of patients with myocardial infarction reported higher mortality and no difference in complications.⁴⁹ Further research is very likely to change our assessment of the outcomes associated with obesity in non-ICU patients. However, all patients should be screened for nutrition risk, and those who are at risk further assessed for nutrition status and potential development of a nutrition support care plan.¹⁵

Clearly, more prospective, adequately powered outcomes research is needed to clarify the risks associated with varying levels of obesity in hospitalized ICU and non-ICU patients. Studies that include measures of inflammation, body composition (with a focus on lean body mass), and micronutrient status

Table 2. Evidence Summary Question 1: Do Clinical Outcomes Vary Across Levels of Obesity in Critically Ill or Hospitalized Patients?

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|--------------------------------------|--|---|--|--|--|
| ICU patients | | | | | |
| Nelson et al, 2012 ⁸⁹ | Retrospective record review Small sample 90 obese patients | Single center trauma database of admissions 1996-present with Injury Severity Score ≥ 16 • BMI ≤ 18.5 kg/m ² , n = 30 • BMI = 18.5-24.9, n = 603 • BMI 25.0-29.9, n = 361 • BMI ≥ 30 , n = 90 Total N = 1084 | Compare resuscitation, treatment, and short-term outcomes by BMI group | Mortality: • BMI ≥ 30 vs normal BMI, OR 2.52 (95% CI, 1.3-4.9) Mortality on day 0: • BMI ≥ 30 vs normal BMI, 8.9% vs 2.8%, $P = .023$ Uncontrolled hemorrhage most common cause | |
| Abhyankar et al, 2012 ⁴⁴ | Retrospective record review Large sample 5287 obese patients | Admissions to single hospital MICU, SICU, or CCU, 2001-2008 • BMI ≤ 18.5 kg/m ² , n = 786 • BMI = 18.5-24.9, n = 5463 • BMI 25.0-29.9, n = 5276 • BMI 30-39.9, n = 4168 • BMI ≥ 40 , n = 1119 Total N = 16,812 | Examine BMI vs 30-day and 1-year mortality | 30-day Mortality: • BMI ≤ 18.5 kg/m ² , OR 1.41 (95% CI, 1.13-1.76) • BMI = 18.5-24.9, reference group • BMI 25.0-29.9, OR 0.81 (95% CI, 0.7-0.93) • BMI ≥ 30 , OR 0.74 (95% CI, 0.64-0.86) 1-year Mortality: • BMI ≤ 18.5 kg/m ² , OR 1.51 (95% CI, 1.18-1.94) • BMI = 18.5-24.9, reference group • BMI 25.0-29.9, OR 0.68 (95% CI, 0.59-0.79) • BMI ≥ 30 , OR 0.57 (95% CI, 0.49-0.67) • BMI ≥ 40 kg/m ² , OR 0.70 (95% CI, 0.54-0.90) | Lower mortality in obese than normal weight patients |
| Hoffmann et al, 2012 ²¹ | Retrospective record review 760 obese subjects Multivariate analysis adjusted for age, new injury severity score, head injury, Glasgow Coma Scale, base excess, coagulation, severe bleeding, cardiac arrest | Trauma patients with Injury Severity Score > 16 , years 2004-2008 in German Society for Trauma Registry • BMI ≤ 20 kg/m ² , n = 269 • BMI = 20-24.9, n = 2617 • BMI 25.0-29.9, n = 2120 • BMI ≥ 30 , n = 760 Total N = 5766 | Determine whether low or high BMI is linked with worse outcomes | Hospital Mortality: • BMI 25.0-29.9 vs normal BMI, OR = 0.99, (95%, CI = 0.76-1.29) • BMI ≥ 30 vs normal BMI, OR 1.6 (95% CI, 1.1-2.3, $P = .009$) Time to Death: • BMI 25.0-29.9 vs normal BMI, 16.6 vs 10.1 days, $P < .001$ • BMI ≥ 30 vs normal BMI, 16.6 vs 10.1 days, $P < .001$ | Mortality increased, and time to death longer |
| Westerly et al, 2011 ²² | Retrospective record review Diagnostic similarity 545 obese patients No adjustment for comorbidities or acuity | Admissions to single hospital 2000-2008 Quartiles of BMI ≥ 40 : • BMI 40-47.5 kg/m ² , n = 127 • BMI 47.6-54.6, n = 151 • BMI 54.7-65, n = 147 • BMI > 65 , n = 120 Total N = 545 | Evaluate outcomes of hospitalized morbidly obese patients | Across quartiles of BMI > 40 , mortality was not different. Hospital LOS increased, $P < .001$ Tracheostomy increased, $P = .001$ | |
| Hutagalung et al, 2011 ²³ | Retrospective record review HR adjusted for acuity measures 2245 obese patients Loss of 24% due to no height/weight | German surgical ICU patients, 2004-2009 • BMI ≤ 18.5 kg/m ² , n = 186 • BMI 18.6-24.9, n = 2633 • BMI 25.0-29.9, n = 4093 • BMI 30-39.9, n = 2066 • BMI ≥ 40 , n = 179 Total N = 9935 | Assess impact of obesity on 60-day hospital mortality | 60-day Mortality: • BMI 25.0-29.9 vs normal BMI, HR (lower HR in study indicates lower risk) 0.86 (95% CI, 0.74-0.99, $P = .047$) • BMI = 30-39.9 vs normal BMI, HR 0.83 (95% CI, 0.69-0.99, $P = .047$) • BMI ≥ 40 vs normal BMI, HR 1.14 (95% CI, 0.74-1.74) | BMI 30-39.9 with lower mortality than normal BMI |

(continued)

Table 2. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-----------------------------------|--|--|--|---|---|
| Evans et al, 2011 ²⁴ | Retrospective record review 154 obese patients, no power calculation Limited statistical analysis | US Level I Trauma Center registry, patients over age 45 years <ul style="list-style-type: none"> BMI < 18.5 kg/m², n = 22 BMI 18.6-24.9, n = 145 BMI 25.0-29.9, n = 140 BMI ≥ 30, n = 154 Total N = 461 | Assess impact of BMI on trauma outcomes, complications, injury distribution, n = 461 | <p>90-day Mortality:</p> <ul style="list-style-type: none"> No statistically significant differences across BMI groups in complications, ICU or hospital LOS, mortality or discharge to home | |
| Martino et al, 2011 ⁴⁵ | Multicenter international prospective observation study Large sample Data analysis adjusted for age, gender, APACHE II score, diagnosis category, geographic region, hospital type, ICU type, product of age and APACHE II score | Adults in 1 of 355 ICUs for more than 72 hours in 2007-2009 <ul style="list-style-type: none"> BMI < 18.5 kg/m², n = 423 BMI 18.5-24.9, n = 3490 BMI 25-29.9, n = 2604 BMI 30-39.9, n = 1772 BMI 40-49.9, n = 348 BMI 50-59.9, n = 118 BMI ≥ 60, n = 58 Total N = 8813 | Evaluate outcomes of severe obesity (BMI ≥ 40 kg/m ²) | <p>60-day Mortality:</p> <ul style="list-style-type: none"> BMI 25-29.9 vs normal BMI, OR 0.81 (95% CI, 0.71-0.91), <i>P</i> < .001 BMI 30-39.9 vs normal BMI, OR 0.74 (95% CI, 0.64-0.84), <i>P</i> < .001 BMI ≥ 40 vs normal BMI, OR 0.87 (95% CI, 0.69-1.09) <p>Ventilator Days:</p> <ul style="list-style-type: none"> BMI 25-29.9 vs normal BMI, HR (low hazard ratio in this study indicates higher risk) 0.97 (95% CI, 0.9-1.05) BMI 30-39.9 vs normal BMI, HR 0.85 (95% CI, 0.78-0.93), <i>P</i> < .001 BMI ≥ 40 vs normal BMI, HR 0.86 (95% CI, 0.77-0.97), <i>P</i> < .05) <p>ICU LOS:</p> <ul style="list-style-type: none"> BMI 25-29.9 vs normal BMI, HR 0.95 (95% CI, 0.88-1.03) BMI 30-39.9 vs normal BMI, HR 0.86 (95% CI, 0.79-0.94), <i>P</i> < .001 BMI ≥ 40 vs normal BMI, HR 0.82 (95% CI, 0.72-0.93), <i>P</i> < .05) <p>Hospital LOS:</p> <ul style="list-style-type: none"> BMI 25-29.9 vs normal BMI, HR 0.98 (95% CI, 0.91-1.05) BMI 30-39.9 vs normal BMI, HR 0.96 (95% CI, 0.89-1.04) BMI ≥ 40 vs normal BMI, HR 0.91 (95% CI, 0.80-1.04) | Obese patients (BMI 30-39.9) with lower mortality; all obese patients with longer ventilator intubation and ICU LOS |
| Serrano et al, 2010 ²⁵ | Retrospective record review 314 obese patients OR adjusted for potential confounders | Admissions to level I trauma center 2008 <ul style="list-style-type: none"> BMI 18.5-24.9, n = 382 BMI 25-29.9, n = 328 BMI 30-39.9, n = 250 BMI ≥ 40, n = 64 Total N = 1024 | Evaluate the importance of obesity as an independent risk factor for nosocomial infection in trauma patients | <p>Infection:</p> <ul style="list-style-type: none"> BMI 30-39.9 vs normal BMI, OR 4.69 (95% CI, 2.18-10.1) BMI ≥ 40 vs normal BMI, OR 5.91 (95% CI, 2.18-16.0) Most common types were pulmonary and wound infections | Obesity is independent risk factor for infection after trauma |

(continued)

Table 2. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-------------------------------------|---|--|--|--|--|
| Wurzinger et al, 2010 ²⁶ | Retrospective record review 66 obese patients, no power calculation | <ul style="list-style-type: none"> BMI ≤ 18.5 kg/m², n = 15 BMI 18.5-24.9, n = 125 BMI 25-29.9, n = 95 BMI 30-39.9, n = 66 Total N = 301 | Evaluate impact of BMI on mortality in patients with septic shock | In adjusted model, no difference in mortality by obesity SAPS II predicts mortality | |
| Duchesne et al, 2009 ⁴⁸ | Retrospective record review Very small sample 52 obese patients | All patients in Level I trauma center 2003-2006, total sample 12,759 patients Those with damage control laparotomy: <ul style="list-style-type: none"> BMI ≤ 18.5-29.9 kg/m², n = 52 BMI 30-39.9, n = 38 BMI ≥ 40, n = 15 Total N = 105 | Examine prevalence of surgical site infections in obese vs nonobese patients | Surgical Site Infections: <ul style="list-style-type: none"> Prevalence ratio in BMI ≥ 40 vs nonobese 4.42 (95% CI, 1.74-11.2) Intraabdominal Abscess: <ul style="list-style-type: none"> Prevalence ratio in BMI ≥ 40 vs nonobese 1.76 (95% CI, 0.73-4.28) Acute Renal Injury: <ul style="list-style-type: none"> Prevalence ratio in BMI 30-39.9 vs nonobese 2.07(95% CI, 1.9-4.7) Prevalence ratio in BMI ≥ 40 vs nonobese 3.07 (95% CI, 1.34-7.03) Multisystem Organ Failure: <ul style="list-style-type: none"> Prevalence ratio in BMI 30-39.9 vs nonobese 1.74 (95% CI, 1.14-2.66) Prevalence ratio in BMI ≥ 40 vs nonobese 1.82 (95% CI, 1.14-2.90) Prevalence ratios adjusted for age, gender, type of injury, blood pressure and base deficit Days on Ventilator: <ul style="list-style-type: none"> Nonobese vs obese vs severely obese, 9.8 ± 7 vs 14 ± 7 vs 24 ± 8, $P = .0001$ Hospital LOS: <ul style="list-style-type: none"> Nonobese vs obese vs severely obese, 14 ± 8 vs 14 ± 11 vs 27 ± 9, $P = .0001$ | |
| Dossett et al, 2009 ⁴⁷ | Prospective cohort observation OR adjusted for age, sex, APACHE II score 686 obese patients | Patients in ICU > 48 hr <ul style="list-style-type: none"> BMI ≤ 18.5 kg/m², n = 640 BMI 18.5-24.9, n = 672 BMI 25-29.9, n = 615 BMI 30-39.9, n = 494 BMI ≥ 40, n = 192 Total N = 2037 | Describe relationship between BMI and site-specific ICU-acquired infection risk | Catheter-related Bloodstream Infection Risk: <ul style="list-style-type: none"> BMI 30-39.9 vs normal BMI, OR 1.9 (95% CI, 1.2-2.9) BMI ≥ 40 vs normal BMI, OR 3.2 (95% CI, 1.9-5.3) | May be due to provider reluctance to pull established lines in patients with difficult venous access |
| Pieracci et al, 2008 ²⁷ | Retrospective record review BMI distribution of patients in ICU > 4 days not clear 232 obese patients | Patients admitted to ICU > 4 days <ul style="list-style-type: none"> BMI ≤ 18.5 kg/m², n = 53 BMI 18.5-24.9, n = 376 BMI 25-29.9, n = 285 BMI 30-39.9, n = 188 BMI ≥ 40, n = 44 Total N = 946 | Test hypothesis that BMI is associated with mortality from surgical critical illness | ROC analysis suggests BMI predicts mortality at level of chance alone Age and APACHE III were strongest predictors in all models, BMI was not significant | |

(continued)

Table 2. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|----------------------------------|--|---|---|--|--|
| Sakr et al, 2008 ⁴⁶ | Prospective observational cohort 505 obese patients Adjusted model | Multicenter study of epidemiology of sepsis in European countries, n = 198 ICUs <ul style="list-style-type: none"> BMI \leq 18.5 kg/m², n = 120 BMI 18.5-24.9, n = 1206 BMI 25-29.9, n = 1047 BMI 30-39.9, n = 424 BMI \geq 40, n = 81 Total N = 2878 | Investigate impact of obesity on morbidity and mortality in European sepsis in acutely ill patients study | BMI does not impact mortality or LOS ICU-acquired Infection: <ul style="list-style-type: none"> Obese vs optimal weight, 10.1% vs 9%, $P < .05$ Severely obese vs optimal weight, 12.3% vs 9.0%, $P < .01$ | |
| Frat et al, 2008 ³⁶ | Prospective case-control observation 82 obese patients Prognostic similarity | Patients matched for age, gender, center and SAPS II score <ul style="list-style-type: none"> BMI $<$ 30, n = 124 BMI \geq 35, n = 82 Total N = 206 | Evaluate influence of severe obesity on morbidity and mortality in mechanically ventilated patients | Only difference in morbidity was more frequent difficulty with tracheal intubation and postextubation stridor in obese No difference in mortality | |
| Morris et al, 2007 ²⁸ | Retrospective record review 165 obese patients OR adjusted for age, APACHE score, admission source, chronic health points, etiology of ALI | All ICU patients with ALI and BMI in 1999-2000 <ul style="list-style-type: none"> BMI $<$ 18.5 kg/m², n = 28 BMI 18.5-24.9, n = 179 BMI 25-29.9, n = 150 BMI 30-39, n = 125 BMI \geq 40, n = 40 Total N = 825 | Evaluate the association between BMI and outcomes in patients with ALI | Mortality: <ul style="list-style-type: none"> Not different by BMI group Discharge Disposition: <ul style="list-style-type: none"> To rehabilitation center BMI \geq 40 vs normal BMI, OR 6.0 (95% CI, 1.8-20.2) To skilled nursing facility BMI \geq 40 vs normal BMI, OR 4.3 (95% CI, 1.5-12.5) | |
| Newell et al, 2007 ³⁷ | Retrospective record review 264 obese patients, no power statement No adjustment of OR | Consecutive admissions to trauma center with Injury Severity Score \geq 16 and blunt trauma in 2001-2005 <ul style="list-style-type: none"> BMI missing n = 357 BMI $<$ 18.5 kg/m², n = 61 BMI 18.5-24.9, n = 554 BMI 25-29.9, n = 529 BMI 30-39, n = 271 BMI \geq 40, n = 93 Total N = 2108 | Evaluate clinical outcomes in blunt trauma patients stratified by BMI | Mortality: <ul style="list-style-type: none"> BMI \geq 40 vs normal BMI, OR 0.81 (95% CI, 0.35-1.86) Complications in BMI 30-39.9 vs normal BMI: <ul style="list-style-type: none"> Acute respiratory failure, OR 1.8 (95% CI, 1.3-2.4) Pneumonia, OR 1.7 (95% CI, 1.2-2.4) UTI, OR 1.8 (95% CI, 1.2-2.9) Complications in BMI \geq 40 vs normal BMI: <ul style="list-style-type: none"> ARDS, OR 3.68 (95% CI, 1.2-10.9) Acute respiratory failure, OR 2.79 (95% CI, 1.6-4.8) Acute renal failure, OR 13.5 (95% CI, 2.4-76.4) MEOF, OR 2.6 (95% CI, 1.09-6.4) Pneumonia, OR 2.5 (95% CI, 1.5-4.3) UTI, OR 2.3 (95% CI, 1.2-4.4) DVT, OR 4.1 (95% CI, 1.3-13.5) Decubitus ulcer, OR 2.8 (95% CI, 1.4-5.8) | Complications higher in severely obese than obese than normal BMI patients |

(continued)

Table 2. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|------------------------------------|---|---|---|---|--|
| Nasraway et al, 2006 ³⁰ | Retrospective record review 96 obese patients model adjusted for age, gender, acuity, renal failure, diabetes, vasopressor use, mechanical ventilation | Consecutive admissions to surgical ICU 1998-2001 • BMI ≤ 18.5 kg/m ² , n = 70 • BMI 18.5-24.9, n = 529 • BMI 25-29.9, n = 408 • BMI 30-39.9, n = 272 • BMI ≥ 40 , n = 94 Total N = 1373 Patients who stayed in ICU ≥ 4 d • BMI ≤ 18.5 kg/m ² , n = 26 • BMI 18.5-24.9, n = 164 • BMI 25-29.9, n = 119 • BMI 30-39.9, n = 74 • BMI ≥ 40 , n = 24 Total N = 406 | Determine whether BMI ≥ 40 is independent risk factor for death in ICU patients | Mortality, ICU LOS and hospital LOS not different in entire group of admissions | |
| Peake et al, 2006 ³⁸ | Prospective cohort observation 125 obese patients Model included age, APACHE II score, albumin Charlson comorbidity index | Patients admitted to medical-surgical ICU in 2001 • BMI < 18.5 kg/m ² , n = 24 • BMI 18.5-24.9, n = 129 • BMI 25-29.9, n = 151 • BMI 30-34.9, n = 75 • BMI ≥ 35 , n = 54 Total N = 433 | Evaluate effect of BMI on 30-day and 12-month survival | Increasing BMI associated with decreasing mortality TR > 1 is increased survival time: • 30-day TR for BMI = 1.85 (95% CI, 1.05, 3.26) • 12-month TR for BMI = 1.03 (95% CI, 1.005, 1.063) | |
| Duane et al, 2006 ³⁹ | Retrospective record review 115 obese patients, no power statement | Blunt trauma patients admitted 2004-2005 • BMI < 30 , n = 338 • BMI ≥ 30 , n = 115 Total N = 453 | Determine effect of obesity on morbidity and mortality in ICU and non-ICU population of blunt trauma patients | No difference in mortality or morbidity measures | |
| Alban et al, 2006 ⁴⁰ | Retrospective record review 135 obese patients, no power statement | Patients admitted to trauma ICU, 1999-2002 Nonobese, n = 783 Obese, n = 135 Total, n = 928 | Compare outcomes of obese vs nonobese patients after trauma | Mortality: • Obese vs nonobese, OR 0.8 (95% CI, 0.3-1.8) • Age > 55 y, OR 3.5 (95% CI, 1.8-6.6) • ISS > 20 , OR 8.9 (95% CI, 4.2-18.8) • APACHE II > 20 , OR 12.0 (95% CI, 4.7-30.6) • Blunt vs penetrating injury, OR 2.0 (95% CI, 1.1-3.9) | Severity of illness more predictive than obesity |
| O'Brien et al, 2006 ⁴² | Retrospective record review 457 obese patients Mortality adjusted for age, gender, race, SAPS II, team model, condition on admission, patient origin, diagnosis of skin or subcutaneous tissue disease, preexisting illness, use of pressors, ICU complications, number of preexisting diseases | Critically ill adults from 106 ICUs in 84 hospitals in acute lung injury IMPACT study • BMI < 18.5 kg/m ² , n = 88 • BMI 18.5-24.9, n = 544 • BMI 25-29.9, n = 399 • BMI 30-39.9, n = 326 • BMI ≥ 40 , n = 131 Total N = 1488 | Determine association between BMI and hospital mortality | Hospital Mortality: • BMI 30-39.9 vs normal BMI, OR 0.67 (95% CI, 0.46-0.97) • BMI ≥ 40 vs normal BMI, OR 0.78 (95% CI, 0.44-1.38) Unadjusted Differences in Care: • BMI ≥ 40 vs normal BMI • Heparin prophylaxis in 57% vs 44% • Tracheostomy, 26% vs 17% • Specialty bed, 29% vs 15% | |

(continued)

Table 2. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-------------------------------------|--|---|--|--|---|
| Aldawood et al, 2006 ³⁵ | Retrospective record review 540 obese patients Unadjusted OR | Critically ill adults from single ICU in Saudi Arabia, 2001-2004 <ul style="list-style-type: none"> • BMI < 18.5kg/m², n = 140 • BMI 18.5-24.9, n = 631 • BMI 25-29.9, n = 524 • BMI 30-34.9, n = 312 • BMI 35-39.9, n = 135 • BMI ≥ 40, n = 93 Total N = 1835 | Examine impact of obesity on hospital and ICU mortality, LOS, duration of mechanical ventilation | <p>Hospital Mortality:</p> <ul style="list-style-type: none"> • BMI ≥ 40 vs normal BMI, OR 0.51 (95% CI, 0.28-0.92, <i>P</i> = .025) Also predicted by chronic respiratory illness, age, medical vs surgical admission | Lowest mortality for BMI ≥ 40 |
| Ray et al, 2005 ³² | Retrospective record review 550 obese patients No adjustment for acuity | Medical ICU admissions 1997-2001 <ul style="list-style-type: none"> • BMI < 20 kg/m², n = 350 • BMI 20-24.9, n = 663 • BMI 25-29.9, n = 585 • BMI 30-39.9, n = 396 • BMI ≥ 40, n = 154 Total N = 2148 | Examine the effect of BMI on ICU outcome | <p>ICU Mortality:</p> APACHE II score predicts (<i>P</i> < .001) but BMI does not (<i>P</i> = .588) <p>Hospital Mortality:</p> APACHE II score predicts (<i>P</i> < .001) but BMI does not (<i>P</i> = .469) <p>Complications:</p> No difference by BMI group | Acuity score predicts mortality better than BMI |
| Winkelman et al, 2005 ⁴¹ | Prospective cohort observation Small sample | Critically ill patients with severe obesity BMI ≥ 40, n = 43 | Describe resources used by nurses to care of patients with severe obesity | <p>Most common equipment:</p> <ul style="list-style-type: none"> • Specialty bed or mattress • Large BP cuff • Large commodes • Large wheelchairs Assist of 2 to reposition patient Special skin care treatment | Nurses should anticipate these needs to avoid poor outcomes |
| Brown et al, 2005 ³³ | Retrospective record review 283 obese patients OR adjusted but factors used not reported | Trauma and ICU database <ul style="list-style-type: none"> • BMI < 30, n = 870 • BMI ≥ 30, n = 283 Total N = 1153 | Evaluate influence of obesity on outcomes after severe blunt trauma | Obesity independent risk factor for mortality: Adj OR 1.6 (95% CI, 1.0- 2.3, <i>P</i> = .03) ISS, GCS, hypotension on admission and age are stronger predictors Obese patients with more total complications, MSOF, ARDS, dialysis, MI | |
| O'Brien, 2004 ³⁴ | Retrospective record review 219 obese patients, no power statement 15% excluded due to missing variables Model not adjusted | Mechanically ventilated patients with ALI enrolled in RCT testing weaning protocols <ul style="list-style-type: none"> • BMI 18.5-24.9, n = 334 • BMI 25-29.9, n = 254 • BMI ≥ 30, n = 219 Total N = 807 | Examine association of obesity and outcome | <p>28-day Mortality:</p> <ul style="list-style-type: none"> • Overweight vs normal BMI, OR 1.09 (95% CI, 0.7-1.7) • Obese vs normal BMI, OR 1.1 (95% CI, 0.7-1.8) • Age, OR 1.04 (95% CI, 1.03-1.06) • APACHE III score, OR 1.02 (95% CI, 1.01-1.03) • Pao2:Fio2 ratio, OR 0.99 (95% CI, 0.99-0.99) • Assigned higher tidal volume, OR 1.7 (95% CI, 1.2-2.4) • Peak airway pressure, OR 1.03 (95% CI, 1.0-1.05) • Trauma diagnosis, OR 0.32 (95% CI, 0.12-0.86) | Acuity factors more important than BMI as predictors of outcome |

(continued)

Table 2. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|--|---|---|--|--|---|
| Garroutte-Orgeas et al, 2004 ⁴³ | Prospective cohort observation 227 obese patients | In 6 medical-surgical ICUs in France over 2 years ● BMI < 18.5, n = 189 ● BMI 18.5-24.9, n = 806 ● BMI 25-29.9, n = 476 ● BMI ≥ 30, n = 227 Total N = 1698 | Examine association between BMI and mortality in adult ICU patients | Mortality: Obese vs normal BMI, OR 0.6 (95% CI, 0.4-0.88) | |
| Tremblay et al, 2003 ²⁹ | Retrospective record review 18,221 obese patients Limited information on comorbid conditions | Project Impact Critical Care Data System, all patients with BMI and at least 1 severity score ● BMI < 18.5, n = 11,479 ● BMI 18.5-24.9, n = 24,332 ● BMI 25-29.9, n = 21,867 ● BMI 30-39.9, n = 13,952 ● BMI ≥ 40, n = 4269 Total N = 75,889 | | Mortality: ● Not significantly different in obese or severely obese from nonobese LOS: ● Not significantly different in obese or severely obese from nonobese | |
| Hospitalized non-ICU patients | | | | | |
| Nafiu et al, 2012 ³⁰ | Retrospective record review 49,761 obese patients Model adjusted for age, anesthesia status, racial group, elective vs emergent surgery | Racial/ethnic minority surgical patients 2005-2008 from 186 centers in National Surgical Quality Improvement Program ● Overall BMI = 30.3 ± 8.9 kg/m ² ● BMI < 18.5 kg/m ² , n = 3230 ● BMI = 18.6-24.9, n = 31,699 ● BMI 25.0-29.9, n = 34,929 ● BMI = 30-39.9, n = 34,450 ● BMI ≥ 40, n = 15,311 Total N = 119,619 | Evaluate contribution of BMI to 30-day postsurgical outcome | 30-day Mortality: ● BMI 18.6-24.9 vs BMI ≥ 40, OR 1.52 (95% CI, 1.23-1.87, P < .001) ● BMI 25.0-29.9 vs BMI ≥ 40, OR 1.33 (95% CI, 1.08-1.65, P = .009) ● BMI = 30-39.9 vs BMI ≥ 40, OR 1.2 (95% CI, 0.97-1.49) Hospital LOS: ● BMI 18.6-24.9, 8.9 ± 14.2 d ● BMI 25.0-29.9, 7.3 ± 12.2, P < .001 vs normal BMI ● BMI = 30-39.9, 6.7 ± 11.6, P < .001 vs normal BMI ● BMI ≥ 40, 5.3 ± 10.5, P < .001 vs normal BMI ● Most perioperative outcomes in obese subjects not different than normal weight | BMI ≥ 40 with lowest mortality & hospital LOS. Authors suggest that obese patients may have less severe disease or that they are monitored vigilantly and treated conservatively |
| Das et al, 2011 ⁴⁹ | Retrospective record review OR adjusted for age, prior PAD, BP, HR, shock, ECG findings, troponin ratio, creatinine 2558 patients with severe obesity | Patients in the National Cardiovascular Data Registry with diagnosis of MI ● BMI missing in 1831 (3.5%) ● BMI ≤ 18.5 kg/m ² , n = 344 ● BMI 18.5-24.9, n = 11,785 ● BMI 25-29.9, n = 19,408 ● BMI 30-39.9, n = 15,596 ● BMI ≥ 40, n = 2558 Total N = 50,149 | Evaluate impact of severe obesity on outcomes in patients with ST-segment MI | Mortality: ● BMI ≥ 40 vs BMI 30-35, Adjusted OR 1.64 (95% CI, 1.32-2.03) Major Bleeding: ● BMI ≥ 40 vs BMI 30-35, Adjusted OR 1.09 (95% CI, 0.94-1.26) | Mortality increased |
| Park et al, 2011 ³¹ | Retrospective record review No acuity scores No adjustment for confounders 147 obese patients | Surgical patients from single hospital 1999-2009 ● BMI 18.5-24.9, n = 469 ● BMI 30-39.9, n = 108 ● BMI ≥ 40, n = 39 Total N = 626 | Determine impact of obesity on perioperative and long-term clinical outcomes after open AAA repair or endovascular aneurysm repair | No difference in LOS, MI, ARF, wound infection, mortality ICU LOS: ● Obese vs normal BMI, P = .03 | |

Low HR indicates increased risk; low OR indicates reduced risk. AAA, abdominal aortic aneurysm; ALI, acute lung injury; APACHE, Acute Physiology and Chronic Health; ARDS, acute respiratory distress syndrome; ARF, acute renal failure; BMI, body mass index; BP, blood pressure; CCU, cardiac care unit; CI, confidence interval; DVT, deep vein thrombosis; GCS, Glasgow coma scale; HR, hazard ratio; ICU, intensive care unit; ISS, injury severity score; LOS, length of stay; MI, myocardial infarction; MICU, medical ICU; MSOF, multi-system organ failure; OR, odds ratio; PAD, peripheral artery disease; RCT, randomized controlled trial; ROC, receiver operator curve; SAPS, simplified acute physiology score; SICU, surgical ICU; TR, time ratio; UTI, urinary tract infection.

Table 3. GRADE Table Question 1: Do Clinical Outcomes Vary Across Levels of Obesity in Critically Ill or Hospitalized Non-ICU Patients?

| Comparison | Outcome | Quantity, Type of Evidence | Findings | Grade for Outcome | Overall Evidence GRADE |
|--|------------------------------|----------------------------|--|-------------------|------------------------|
| ICU patients | | | | | |
| Obese vs optimal BMI | Mortality (large studies) | 8 OBS | 1 increased ²¹ 5 decreased ^{23,35,42,44,45} 2 no difference ^{32,46} | Low | Low |
| | Hospital LOS (large studies) | 4 OBS | 3 increased ^{22,29,45} 1 no difference ⁴⁶ | Low | |
| | Complications | 6 OBS | 5 increased ^{25,37,46-48} 1 no difference ³² | Low | |
| BMI \geq 40 kg/m ² vs optimal BMI | Mortality (large studies) | 4 OBS | 1 decreased ⁴⁴ 3 no difference ^{22,23,45} | Low | |
| | Hospital LOS (large studies) | 4 OBS | 2 increased ^{22,29} 2 no difference ^{45,46} | Low | |
| Non-ICU patients | | | | | |
| Obese vs optimal BMI | Mortality | 2 OBS | 1 increased ⁴⁹ 1 no difference ⁹¹ | Low | |

ICU, intensive care unit; LOS, length of stay; OBS, observational study.

would be especially helpful. Finally, nutrition support interventions that aim to improve clinical outcomes are needed in this population.

Question 2: How Should Energy Requirements Be Determined in Obese Critically Ill or Hospitalized Non-ICU Patients? (Table 4)

Recommendation

2a. In the critically ill obese patient, if indirect calorimetry is unavailable, energy requirements should be based on the Penn State University 2010 predictive equation or the modified Penn State University equation if the patient is over the age of 60 years (strong).

Evidence Grade: High.

2b. In the hospitalized obese patient, if indirect calorimetry is unavailable and the Penn State University equations cannot be used, energy requirements may be based on the Mifflin–St Jeor equation using actual body weight (weak).

Evidence Grade: Moderate.

Rationale. Most studies recommend the use of indirect calorimetry to measure resting energy expenditure (REE); however, some patients do not meet valid testing criteria, and most facilities do not have indirect calorimeters. Avoiding energy overfeeding is an important goal; therefore either REE or use of a predictive equation to approximate REE is an essential part of nutrition assessment. In the critically ill, ventilator-dependent obese patient, the Penn State University (PSU) predictive equation most accurately predicts REE compared with others (including Harris–Benedict, Mifflin–St

Jeor, Swinamer, and Ireton-Jones). Frankenfield and colleagues compared multiple predictive equations with REE in patients with BMI \geq 30 kg/m² and found the PSU equation to have the highest prediction accuracy of 70% (\pm 10% of REE) with the least bias or the lowest likelihood of over or underestimation.⁵⁰ In another comparison study in critically ill patients with BMI \geq 45 kg/m², accuracy of the PSU equation was highest at 76% (\pm 10% of REE) compared with other equations studied.⁵¹ In the older critically ill obese patient (\geq 60 years) with BMI \geq 30, a modified PSU appears to be more accurate than the original PSU.⁵⁰ When compared with the unmodified version, the modified PSU was found to have an accuracy rate of 70% (\pm 10% of REE) vs 58% ($P = .04$).⁵⁰ Further, in a case series of 7 patients (including 2 obese patients) with REE measured continuously for 7 days, the prediction error using the PSU equation was only a total of -468 ± 642 kcal ($-3.7 \pm 5.1\%$) over 1 week.⁵²

The PSU equations⁵³ are as follows:

Younger obese patients:

- $RMR \text{ (kcal/d)} = MSJ(0.96) + Tmax(167) + VE(31) - 6212$

Older obese patients:

- $RMR \text{ (kcal/d)} = MSJ(0.71) + Tmax(85) + VE(64) - 3085$
- \circ Where MSJ = Mifflin–St Jeor equation (below); V_E = minute ventilation (L/minute); T_{max} = maximum temperature in prior 24 hours in degrees C

In the mixed ICU and non-ICU patients, the evidence is more difficult to assess due to several important variables. The

Table 4. Evidence Summary Question 2: How Should Energy Requirements Be Determined in Obese Critically Ill or Hospitalized Non-ICU Patients?

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|--|---|--|---|---|--|
| Frankenfield et al, 2012 ³¹ | Validation study Similar prognosis in obese group 55 obese patients | Critically ill patients at extremes of BMI BMI ≤ 21 kg/m ² , n = 56 BMI ≥ 45 kg/m ² , n = 55 | Validate the PSU prediction equation and test validity of II, ACCP, MSJ, HB | <p>ICU patients</p> <p>Accuracy within 10% REE (%):</p> <ul style="list-style-type: none"> PSU (76%) MSJ (55%) HB (60%) IJ (29%) ACCP (27%) <p>Bias in kcal/d (95% CI):</p> <ul style="list-style-type: none"> PSU (-33, +97) MSJ (-299, -82) HB (-105, +149) IJ (+283, +509) ACCP (-616, -403) | PSU valid in severely obese, critically ill patients |
| Kross et al, 2012 ⁹² | Retrospective validation study 401 obese patients | All mechanically ventilated patients with REE between 1998-2005 <ul style="list-style-type: none"> BMI 18.5-24.9, n = 254 BMI 25-29.9, n = 272 BMI 30-34.9, n = 176 BMI 35-39.9, n = 84 BMI ≥ 40, n = 141 Total N = 925 | Compare REE with HB, Owen, MSJ, IJ, ACCP | <p>BMI 30-34.9:</p> <p>Accuracy (%):</p> <ul style="list-style-type: none"> MSJ (18.8%) HB (34.1%) IJ (20.5%) ACCP (9.7%) Owen (9.7%) <p>Bias mean (95% CI):</p> <ul style="list-style-type: none"> MSJ, -177.8 (-203.9, -151.6) HB, -53.4 (-78.6, +10.1) IJ, -86.4 (-117.6, -55.2) ACCP, -218.7 (-245.3, -192.2) Owen, -205.6 (-233.1, +177.9) <p>BMI 35-39.9:</p> <p>Accuracy (%):</p> <ul style="list-style-type: none"> MSJ (18.8%) HB (27.4%) IJ (20.5%) ACCP (7.1%) Owen (14.3%) <p>Bias mean (95% CI)</p> <ul style="list-style-type: none"> MSJ, -166.6 (-209.4, -123.8) HB, -66.0 (-105.1, +27.3) IJ, -101.9 (-76.7, +23.8) ACCP, -243.7 (-285.5, -202.1) Owen, -198.9 (-240.2, -157) <p>BMI ≥ 40:</p> <p>Accuracy (%):</p> <ul style="list-style-type: none"> MSJ (33.3%) HB (28.4%) IJ (14.2%) ACCP (1.4%) Owen (20.6%) <p>Bias mean (95% CI):</p> <ul style="list-style-type: none"> MSJ, -91.8 (-119.5, -64.0) HB, -61.1 (-55.8, +19.5) IJ, -91.3 (-133.9, -48.7) ACCP, -243.7 (-319.1, -261.4) Owen, -145.2 (-174.1, -116.3) | Unable to evaluate PSU or Swinamer due to missing minute ventilation or tidal volume Equations are not adequate |

(continued)

Table 4. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|---|--|---|---|--|--|
| Frankenfield, 2011 ⁵³ | Validation study Included archived data in analysis, unclear prognostic similarity Precise measurement protocol | Obese, older ICU patients, n = 50 Age 70 ± 7 y BMI 38.4 ± 7.2 kg/m ² Data from previous studies: n = 79 | Test the validity of a modified PSU equation against Deltatrac REE measures | Accuracy: ● Modified PSU = 70% ● Original PSU = 66% Bias (95% CI): ● Modified PSU (-120, -12) kcal/d ● Original PSU (-90, +25) kcal/d | Both PSU equations include both body size and metabolic factors (temperature, minute ventilation) |
| Frankenfield et al, 2009 ⁵⁰ | Validation study Similar prognosis | REE measures in 202 critically ill patients in 2006-2007: Obese young: n = 47 Obese elderly: n = 51 | Compare REE measured by Deltatrac calorimeter with estimates by HB, MSJ, ACCP, Swinamer, IJ, PSU, Brandi, and Faisy equations | Accuracy: Young Obese: ● PSU (66%) ● MSJ (21%) ● HB (45%) ● IJ (49%) ● ACCP (53%) Elderly Obese: ● PSU (46%) ● MSJ (35%) ● HB (35%) ● IJ (51%) ● ACCP (12%) Bias (95% CI): Young Obese: ● PSU (-249, -31) ● MSJ (-544, -316) ● HB (-368, +89) ● IJ (-249, -31) ● ACCP (358, 874) Elderly Obese: ● PSU (-51, +133) ● MSJ (-440, -215) ● HB (-357, -126) ● IJ (-174, +31) ● ACCP (457, 749) | PSU equation unbiased and precise across all age and weight groups |

(continued)

Table 4. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-------------------------------------|--|---|--|--|---|
| Alves et al., 2009 ⁹³ | Validation study Dissimilar prognosis | Overweight or obese ICU patients Mean BMI 36.41 ± 9.03 kg/m ² Fasting, n = 42 Stable feeding, n = 29 | Compare REE measured by Deltatrac calorimeter with estimates by HB, IJ equations, and 21 kcal/kg of actual, average, and adjusted body weight | Accuracy (Concordance Coefficient): Fasted measures: • HB actual weight (0.767) • IJ actual weight (0.452) • 21 kcal/kg actual weight (0.446) Fed measures: • HB actual weight (0.829) • IJ actual weight (0.641) • 21 kcal/kg actual weight (0.490) Bias: Fasted measures: • HB actual weight -81.3 (-726.1, +563.4) • IJ actual weight -644.2 (-1369.8, +81.4) • 21 kcal/kg actual weight -413.3 (-1527.7, +701) Fed measures: • HB actual weight -63.7 (-658.3, +530.8) • IJ actual weight 461.9 (-172.7, +1096.5) • 21 kcal/kg actual weight +315.9 (-924.5, +1555.7) Use of adjusted body weight produced less accurate estimates | REE should be measured Bias with best equation could result in change in body weight if applied to energy delivery |
| Anderegg et al., 2009 ³⁵ | Validation study Dissimilar prognosis Different measuring devices Small sample | Hospitalized adult patients with BMI 38.2 ± 8 kg/m ² Ventilated, n = 27 Spontaneously breathing, n = 9 Total N = 36 | Identify which of 4 predictive equations gave estimates within 10% of measured energy expenditure by Deltatrac (ventilated) or Medgem (spontaneously breathing). | Accuracy: • HB actual weight (38.9%) • MSJ (19.4%) • IJ ventilator (38.9%) • 21 kcal/kg actual weight (41.5%) Bias (mean ± SD): • HB 110.1 ± 478.3 • MSJ 215.8 ± 470.7 • IJ 152.3 ± 399.1 • 21 kcal/kg actual weight -271 ± 641.7 Mean REE: • Ventilated 20.4 ± 5.1 kcal/kg/d • Spontaneously breathing, 15.5 ± .9 kcal/kg/d | Indirect calorimetry should be employed to measure energy expenditure in obese hospitalized patients |
| Boullata et al., 2007 ⁵⁴ | Retrospective record validation study Dissimilar prognosis Unclear how many obese patients are ventilator vs canopy measures | All patients with an REE in 1991, n = 395 Ventilator measures, n = 141 Canopy measures, n = 254 Obese, n = 51 | Evaluate the accuracy of 7 predictive equations against measured REE in hospitalized patients, including the critically ill and obese | Accuracy: • HB actual weight (62%) • IJ (32%) Bias: • HB +47 (-440, +534) | Data collection predates current level of obesity |

(continued)

Table 4. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|--|---|---|---|---|--|
| Dobratz et al, 2007 ⁵⁷ | Validation study Similar prognosis Small sample | Female pre-bariatric surgery patients, n = 14 BMI 49.8 ± 6.2, (range 41.3-65.3) kg/m ² | Identify which of 12 prediction equations is most accurate relative to measured REE using Deltatrac calorimeter | <p>Accuracy:</p> <ul style="list-style-type: none"> MSJ (86%) HB actual weight (69%) <p>Bias (mean difference):</p> <ul style="list-style-type: none"> MSJ -48 ± 191 kcal HB actual weight -89 ± 187 kcal/day <p>Use of adjusted body weight with HB equation made the underestimate worse</p> <p>Error for all predictive equations (including MSJ) ≥ 250 kcal</p> | Small sample Clinically stable prior to bariatric surgery Prediction error might result in change in body weight if applied to energy delivery |
| Frankenfield et al, 2003 ⁵⁶ | Validation study | Healthy volunteers and bariatric surgery patients in a hospital setting All canopy measures, BMI range up to 96.8 kg/m ² Nonobese, n = 83 BMI 30-39.9, n = 20 BMI ≥ 40, n = 27 | Evaluate equations for predicting resting metabolic rate against measured values in obese and nonobese people | <p>Accuracy of MSJ:</p> <ul style="list-style-type: none"> BMI 30-39.9 (70%), 10% underestimates, 20% overestimates BMI ≥ 40 (70%), 7% underestimates, 23% overestimates <p>Accuracy of HB:</p> <ul style="list-style-type: none"> BMI 30-39.9 (50%), 40% underestimates, 10% overestimates BMI ≥ 40 (74%), 22% underestimates, 4% overestimates | |

Bias is the 95% CI of difference between estimated and measured REE; precision is the percentage of measures ± 10% REE. ACCP, American College of Chest Physicians; CI, confidence interval; HB, Harris-Benedict; ICU, intensive care unit; IJ, Ireton-Jones; MSJ, Mifflin-St Jeor; PSU, Penn State University; REE, resting energy expenditure.

5 studies reviewed compared multiple predictive equations (Harris–Benedict, Schofield, Mifflin–St Jeor, and others) with REE but did not include all the same predictive equations in each. All included very small samples of obese patients, 1 reported on data collected in 1991,⁵⁴ and 1 used measures from 2 different calorimeter devices.⁵⁵ Accuracy ($\pm 10\%$ of REE) varied among the equations studied with Mifflin–St Jeor (MSJ) demonstrating the highest accuracy at 70%⁵⁶–86%⁵⁷ compared with 50% for Harris–Benedict with adjusted weight⁵⁵ and 50%,⁵⁶ 62%⁵⁴–69%⁵⁷ for Harris–Benedict using actual weight. In addition, significant bias⁵⁵ and prediction errors^{54,57} were measured that could result in undesired weight changes when applied to specific patients. The error for MSJ, however, was lower than that demonstrated with Harris–Benedict using actual weight.^{56,57}

The MSJ⁵⁸ equations are as follows:

- Men (kcal/day) = $5 + 10 \times \text{Weight (kg)} + 6.25 \times \text{Ht(cm)} - 5 \times \text{Age(y)}$
- Women (kcal/day) = $-161 + 10 \times \text{Weight (kg)} + 6.25 \times \text{Ht(cm)} - 5 \times \text{Age(y)}$

Whether provision of energy requirements based on REE provides superior clinical outcomes in hospitalized patients to those with energy needs estimated by a predictive equation has not yet been evaluated in patients with obese or optimal BMI.

Question 3: Are Clinical Outcomes Improved With Hypocaloric, High Protein Diets in Hospitalized Patients With Obesity? (Tables 5–6)

Recommendation

3a. Clinical outcomes are at least equivalent in patients supported with high protein hypocaloric feeding to those supported with high protein eucaloric feeding. A trial of hypocaloric high protein feeding is suggested in patients who do not have severe renal or hepatic dysfunction (weak). Hypocaloric feeding may be started with 50%–70% of estimated energy requirements or < 14 kcal/kg actual weight. High protein feeding may be started with 1.2 g/kg actual weight or 2–2.5 g/kg ideal body weight, with adjustment of goal protein intake by the results of nitrogen balance studies.

Evidence Grade: Low.

3b. Hypocaloric low protein feedings are associated with unfavorable outcomes. Clinical vigilance for adequate protein provision is suggested in patients who do not have severe renal or hepatic dysfunction (weak).

Evidence Grade: Low.

Rationale. Insulin resistance, glucose intolerance, hyperlipidemia, nonalcoholic fatty liver disease, and hypoventilation syndrome are more prevalent in patients with obesity than non-obese patients.⁵⁹ As a result, the hospitalized patient with

obesity is susceptible to experiencing complications associated with overfeeding. Because of these concerns, hypocaloric, high protein regimens have been designed by clinicians in an effort to minimize potential overfeeding complications while simultaneously achieving net protein anabolism.

Hypocaloric feeding is defined as providing a caloric intake less than measured or estimated energy expenditure whereas eucaloric feeding is intended to provide a caloric intake sufficient to meet caloric needs as assessed by measured energy expenditure. Hypercaloric feeding is the provision of a caloric intake greater than caloric requirements. Hypocaloric, high protein feeding is often mistaken for permissive underfeeding. Permissive underfeeding allows for both protein and caloric deficits whereas the intent of hypocaloric, high protein diets is to provide only a calorie deficit while ensuring adequate protein intake.

Four comparative studies^{59–62} and 2 case series^{63,64} examined the use of hypocaloric, high protein nutrition therapy for hospitalized patients with obesity. The hypocaloric, high protein diets contained average intakes ranging from 90 g to 140 g of protein and 900 kcals to 1300 kcals daily (Table 4). Significantly improved clinical outcomes, as evidenced by decreased LOS in the ICU, decreased duration of antibiotic therapy, and a trend toward decreased days of mechanical ventilation, were suggested in a single small observational study examining hypocaloric, high protein diets vs eucaloric, high protein diets for critically ill trauma patients with obesity.⁶¹ Positive clinical outcomes were also noted for use of hypocaloric, high protein feeding in 2 observational case series of surgical patients with obesity.^{63,64} In the only randomized controlled trial that examined clinical outcomes,⁵⁹ no difference in mortality or length of hospital stay was found for hospitalized patients with obesity who received hypocaloric high protein feeding when compared with eucaloric high protein diets. All 3 comparative studies^{59–61} indicated that nutrition outcomes, such as nitrogen balance and serum protein response, were similar between eucaloric and hypocaloric feeding in the presence of adequate protein intake. However, 1 large observational study indicated a worsened 60-day mortality rate when a hypocaloric diet was combined with a low protein intake (average daily caloric and protein intakes of 1000 kcals and 46 g, respectively) and given to hospitalized patients with Class II (BMI 35–39.9 kg/m²) obesity.⁶⁵

The current literature, which includes a total of 163 patients supported with hypocaloric, high protein regimens, indicates that clinical outcomes for hospitalized patients with obesity are at least equivalent, if not improved, by the provision of hypocaloric feeding when adequate protein intake is given to achieve net protein anabolism. A large randomized controlled trial is warranted to ascertain whether hypocaloric, high protein nutrition therapy offers a significant therapeutic advantage over eucaloric or hypercaloric feeding with respect to clinical outcomes and avoidance of complications from overfeeding for hospitalized patients with obesity.

Table 5. Evidence Summary Question 3: Are Clinical Outcomes Improved With Hypocaloric, High Protein Diets in Hospitalized Patients?

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-------------------------------------|---|---|--|--|----------|
| Dickerson et al, 2013 ⁶² | Retrospective cohort observation | Admissions to trauma center, 2009-2011 with BMI ≥ 30 kg/m ² BMI = 35 ± 6 kg/m ² Weight = 105 ± 26 kg Age 18-59 years, n = 41 Age ≥ 60 years, n = 33 | Examine whether older, critically ill trauma patients who are obese achieve nitrogen equilibrium and obtain similar clinical outcomes to younger obese patients during hypocaloric, high protein therapy | Daily Nutrient Delivery: <ul style="list-style-type: none"> Younger: 18 kcal/kg ideal weight, protein 1.9 g/kg ideal weight Older: 21 kcal/kg ideal weight, protein 2.1 g/kg ideal weight ($P < .05$) ICU LOS: 28 \pm 17 vs 30 \pm 13 days in younger vs older Hospital LOS: 45 \pm 30 vs 34 \pm 14 days in younger vs older, $P = .065$ Sepsis: 83% vs 76% in younger vs older, $P = .041$ Pneumonia: 39% vs 48% in younger vs older Antibiotic days adjusted for mortality: 10 \pm 3 vs 8 \pm 4 days in younger vs older, $P = .041$ | |
| Hamilton et al, 2011 ⁶³ | Retrospective record review No control Small sample | Bariatric surgery patients admitted for initiation of home PN to treat bowel obstruction or leak/fistula, 2000-2008 with follow-up data from home Baseline BMI = 39.8 (IQR 36.1, 48.1) Baseline weight = 113 kg (IQR 94.5, 134) N = 23 | Evaluate effect of hypocaloric PN on weight loss, albumin level, PN complications | Daily Nutrient Delivery: <ul style="list-style-type: none"> Energy 13.6 kcal/kg actual body weight Protein 132.6 \pm 6.6 g, 1.2 \pm 0.3 g/kg body weight Weight Loss: <ul style="list-style-type: none"> -7.0 \pm 5.1% in 1.5 months Complications: <ul style="list-style-type: none"> Readmission 52.5% | |

(continued)

Table 5. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-----------------------------------|--|---|---|--|---|
| Alberda et al, 2009 ⁶⁵ | Prospective cohort observation Some differences in cardiovascular dx at admission, similar APACHE II score OR adjusted for nutrition days, BMI, age, admission category, dx, APACHE II score 728 obese subjects, but < 200 in each of BMI 35-39.9 and > 40 groups | Adult patients admitted to 1 of 167 ICUs in 37 countries ● BMI < 20 kg/m ² , n = 289 ● BMI 20-24.9, n = 937 ● BMI 25-29.9, n = 818 ● BMI 30-34.9, n = 395 ● BMI 35-39.9, n = 162 ● BMI ≥ 40, n = 171 Total N = 2772 | Examine the relationship between amount of energy and protein provided to clinical outcomes, and the impact of preillness BMI on outcomes | <p>Daily Energy Intake:</p> <ul style="list-style-type: none"> ● BMI < 20 kg/m², 994 ± 469 kcal; 19.7 ± 9.6 kcal/kg ● BMI 20-24.9, 1024 ± 490; 15.7 ± 7.5 kcal/kg actual weight ● BMI 25-29.9, 1074 ± 536; 13.6 ± 6.7 kcal/kg ● BMI 30-34.9, 1008 ± 534 kcal; 11.2 ± 4.9 kcal/kg ● BMI 35-39.9, 1009 ± 532 kcal; 9.8 ± 5.1 kcal/kg ● BMI ≥ 40, 1048 ± 531 kcal; 8.1 ± 4.4 kcal/kg <p>Daily Protein Intake:</p> <ul style="list-style-type: none"> ● BMI < 20 kg/m², 44.7 ± 23.4 g; 0.9 ± 0.5 g/kg ● BMI 20-24.9, 46.7 ± 25.9 g; 0.7 ± 0.4 g/kg ● BMI 25-29.9, 47.5 ± 28.3 g; 0.6 ± 0.3 g/kg ● BMI 30-34.9, 47.9 ± 28.3 g; 0.5 ± 0.3 g/kg ● BMI 35-39.9, 45.8 ± 29.2 g; 0.4 ± 0.3 g/kg ● BMI ≥ 40, 50.3 ± 33.3 g; 0.4 ± 0.3 g/kg <p>60-day Mortality Per 1000 kcal/day Increase in Energy Intake:</p> <ul style="list-style-type: none"> ● BMI < 20 kg/m², OR 0.52 (95% CI, 0.29-0.95, P = .03) ● BMI 20-24.9, OR 0.62 (95% CI, 0.44-0.88, P = .007) ● BMI 25-29.9, OR 1.05 (95% CI, 0.75-1.49) ● BMI 30-34.9, OR 1.04 (95% CI, 0.64-1.68) ● BMI 35-39.9, OR 0.36 (95% CI, 0.16-0.80, P = .012) ● BMI ≥ 40, OR 0.63 (95% CI, 0.32-1.24) <p>60-day Mortality per 30 g Increase in Protein Intake:</p> <ul style="list-style-type: none"> ● BMI < 20 kg/m², OR 0.60 (95% CI, 0.41-0.87, P = .007) ● BMI 20-24.9, OR 0.81 (95% CI, 0.66-0.99, P = .036) ● BMI 25-29.9, OR 0.97 (95% CI, 0.79-1.19) ● BMI 30-34.9, OR 1.04 (95% CI, 0.79-1.37) ● BMI 35-39.9, OR 0.62 (95% CI, 0.39-0.98, P = .039) ● BMI ≥ 40, OR 0.72 (95% CI, 0.51-1.03) | Energy and protein targets for patients with obesity go down as BMI increases (20.2 kcal/kg and 1.0 g/kg; 17.9 kcal/kg and 0.9 g/kg; 15.0 kcal/kg and 0.8 g/kg; and for BMI 30-34.9, 35-39.9, ≥ 40 respectively) Increased energy and protein intake may be important for patients with BMI 35-39.9, not significantly so for BMI ≥ 40 |

(continued)

Table 5. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-------------------------------------|--|---|---|---|----------|
| Choban et al, 2005 ⁶⁶ | Retrospective record review | Obese adult patients from 2 sites BMI 30-39.9 kg/m ² , n = 48 BMI ≥ 40 kg/m ² , n = 22 | Evaluate protein requirements, using nitrogen balance, in hospitalized patients with obesity | <p>Protein Requirement:</p> <p>ICU Patients:</p> <ul style="list-style-type: none"> BMI 30-39.9 kg/m², 1.9 g/kg ideal body weight/day BMI ≥ 40 kg/m², 2.5 g/kg ideal body weight/day <p>Non-ICU Patients:</p> <ul style="list-style-type: none"> BMI 30-39.9 kg/m², 1.7 g/kg ideal body weight/day BMI ≥ 40 kg/m², 1.8 g/kg ideal body weight/day | |
| Dickerson et al, 2002 ⁶¹ | Retrospective record review Similar prognosis Small sample | Obese adult patients with > 7 days enteral tube feeding in surgical ICU Baseline BMI 41.3 ± 4.7 kg/m ² and weight 118 ± 41 kg 36 ± 12.4 kg/m ² and weight 102 ± 36 kg in eucaloric group Hypocaloric as energy intake < 20 kcal/kg adjusted body weight and protein intake 2 g/kg ideal body weight, n = 28 Eucaloric as energy intake ≥ 20 kcal/kg adjusted body weight and protein 2 g/kg ideal body weight, n = 12 Total N = 40 | Evaluate nutrition and clinical efficacy of eucaloric vs hypocaloric enteral feeding Daily feeding plan: <ul style="list-style-type: none"> Both groups with protein 2 g/kg ideal body weight (1.2 g/kg actual weight) Eucaloric goal 25-30 total kcal/kg adjusted body weight; actual intake 18.5-25.9 kcal/kg current body weight and 0.8-1.2 g protein/kg current body weight Hypocaloric goal < 20 kcal/kg adjusted body weight; actual intake 13.4-19.2 kcal/kg current body weight and 0.7-0.9 g protein/kg current body weight | <p>Actual intake:</p> <ul style="list-style-type: none"> Hypocaloric vs Eucaloric: 1285 ± 325 kcal, 90 ± 24 g protein vs 1841 ± 482 kcal, 111 ± 32 g protein daily <p>Length of ICU Stay:</p> <ul style="list-style-type: none"> Hypocaloric vs Eucaloric, 18.6 ± 9.9 vs 28.5 ± 16.1 days, <i>P</i> < .03 <p>Ventilator Days:</p> <ul style="list-style-type: none"> Hypocaloric vs Eucaloric, 15.9 ± 10.8 vs 23.7 ± 16.6 days, <i>P</i> = .09 <p>Duration Antibiotic Therapy:</p> <ul style="list-style-type: none"> Hypocaloric vs Eucaloric, 16.6 ± 11.7 vs 27.4 ± 17.3 days, <i>P</i> = .03 <p>Nutrition Measures:</p> <ul style="list-style-type: none"> No difference in nitrogen balance, change in prealbumin or albumin | |
| Choban et al, 1997 ⁵⁹ | RCT Balanced prognosis Blinded delivery of PN Indirect outcomes | Obese adult patients referred for PN, BMI 35 (range 26-46.5) kg/m ² Hypocaloric high protein PN, n = 16 Eucaloric high protein PN, n = 14 Total N = 30 | Evaluate efficacy of hypocaloric vs eucaloric PN with protein 2 gm/kg ideal body weight Daily feeding plan: <ul style="list-style-type: none"> Eucaloric goal with kcal/nitrogen 150:1, actual intake 1936 ± 198 kcal and 108 ± 14 g protein (1.2 g/kg actual weight, 2 g/kg ideal weight) Hypocaloric goal with kcal/nitrogen 75:1, actual intake 1293 ± 299 kcal and 120 ± 27 g protein | <p>Daily Nutrient Delivery:</p> <ul style="list-style-type: none"> Hypocaloric 1293 ± 298 nonprotein kcal, 120 ± 27 g protein Eucaloric 1936 ± 198 nonprotein kcal, 108 ± 14 g protein <p>Change in body weight</p> <ul style="list-style-type: none"> Hypocaloric vs Eucaloric: 0 ± 6.8 kg vs 2.7 ± 7kg <p>Change in Albumin:</p> <ul style="list-style-type: none"> Hypocaloric vs Eucaloric: -1 ± 2 g/L vs -2 ± 2 g/L <p>Nitrogen Balance:</p> <ul style="list-style-type: none"> Hypocaloric vs Eucaloric, 4.0 ± 4.2 vs 3.6 ± 4.1 g nitrogen | |

(continued)

Table 5. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-------------------------------------|--|---|--|---|----------|
| Burge et al, 1994 ⁶⁰ | RCT Unblinded PN delivery Indirect outcomes Small sample | Obese patients referred for PN BMI = 33 ± 5.5 kg/m ² Weight 77-114 kg Hypocaloric high protein PN, n = 9 vs Eucaloric high protein PN, n = 7 Total N = 16 | Evaluate impact of hypocaloric PN on nitrogen balance Daily feeding plan: • Eucaloric goal with kcal at 100% REE, kcal/nitrogen 150:1, actual intake, actual intake 2492 ± 298 kcal (25 kcal/kg actual weight) and 130 ± 15 g protein (1.2 g/kg or 2 g/kg ideal weight) Hypocaloric goal with 50% REE and kcal/nitrogen 75:1, actual intake 1285 ± 374 kcal (14 kcal/kg actual weight) and 111 ± 32 g protein (1.3 g/kg actual weight, 2 g/kg ideal weight) | Daily Nutrient Delivery: • Hypocaloric 585 ± 170 nonprotein kcal, 110.9 ± 32 g protein • Eucaloric 1972 ± 235 nonprotein kcal, 130 ± 15.5 g protein Change in body weight • Hypocaloric vs Eucaloric: -4.1 ± 6. kg vs -7.4 ± 8.4kg (-4.5% vs7.3%) Nitrogen Balance: • Hypocaloric vs Eucaloric, 1.3 ± 3.62 vs2.83 ± 6.9 g | |
| Dickerson et al, 1986 ⁶⁴ | Prospective cohort Uncontrolled Balanced prognosis Small sample | Obese, stressed surgical patients requiring PN Baseline weight 127 ± 60 kg (range 90-302 kg) N = 13 | Evaluate efficacy of hypocaloric, high-protein feeding Daily Nutrient Delivery: • Nonprotein kcal 881 ± 393 (51% REE) • Protein 129 g or 2.1 ± 0.6 g/kg ideal body weight or 1.2 ± 0.5 g/kg actual weight, 2.1 g/kg ideal weight | Nitrogen Balance: • +2.4 g/day Weight Loss: • 2.3 ± 2.7 kg/week Wound Healing: • All fistulas or dehiscence healed by 35.8 ± 18.1 days Adverse Events in Single Patients: • Ketonuria • Mild skin rash that responded to zinc and lipid intake • Acute renal failure due to antibiotic therapy • Readmission for recurrent anastomotic leak | |

APACHE, Acute Physiology and Chronic Health Evaluation; BMI, body mass index; ICU, intensive care unit; IQR, interquartile range; LOS, length of stay; OR, odds ratio; PN, parenteral nutrition; RCT, randomized controlled trial.

Table 6. GRADE Table Question 3: Are Clinical Outcomes Improved With Hypocaloric, High Protein Diets in Hospitalized Patients?

| Comparison | Outcome | Quantity, Type Evidence | Finding | Final GRADE | Overall Evidence GRADE |
|--|------------------|-------------------------|----------------------------------|-------------|------------------------|
| Hypocaloric/high protein vs eucaloric/high protein | LOS | 1 OBS | 1 decreased ⁶¹ | Low | Low |
| | Nitrogen Balance | 1 RCT, 3 OBS | 4 no difference ⁵⁹⁻⁶² | Low | |
| | Weight Loss | 1 RCT, 1 OBS | 2 no difference ^{59,60} | Low | |

LOS, length of stay; OBS, observational study; RCT, randomized controlled trial.

Data to support this recommendation are in Table 3, where protein intake of 1.2 g/kg actual body weight (2 g/kg ideal body weight) daily was given to patients in 5 observational studies^{59-62,64} with hypocaloric or eucaloric energy intake. An additional study compared protein requirements based on nitrogen balance studies separately for ICU and non-ICU patients. The ICU patients required 2-2.5 g/kg/day and the non-ICU patients 1.8-1.9 g/kg/d to approach nitrogen equilibrium with the higher requirements for those with BMI > 40 kg/m².⁶⁶ These studies included patients up to 302 kg and BMI 50.6 kg/m², however most subjects were considerably below these levels. Data have not been found to establish reasonable nitrogen intake goals for patients beyond these limits. Nitrogen balance was similar at this level of protein intake whether energy intake was hypocaloric or eucaloric. These initial recommendations should be adjusted using nitrogen balance studies, with a goal of nitrogen equilibrium if possible (-4 to +4 g nitrogen/kg/d).⁶¹ While older studies may have suggested increase in albumin or prealbumin concentration as a goal for protein intake, a more recent appreciation of the strong impact of inflammation on these measures makes them unreliable as a marker of nutrition state in most ill, hospitalized patients.

Question 4: In Obese Patients Who Have Had Malabsorptive or Restrictive Surgical Procedures for Weight Loss, What Micronutrients Should Be Evaluated? (Tables 7-8)

Recommendation

Patients who have undergone sleeve gastrectomy, gastric bypass, or biliopancreatic diversion ± duodenal switch have increased risk of nutrient deficiency. In acutely ill hospitalized patients with history of these procedures, evaluation for evidence of depletion of iron, copper, zinc, selenium, thiamine, folate, and vitamins B₁₂, and D is suggested as well as repletion of deficiency states. (weak).

Evidence Grade: Low.

Rationale. Bariatric surgical procedures that change the capacity of the stomach facilitate weight reduction by restriction, that is, increasing satiety and reducing caloric intake.

Procedures that shorten small bowel absorptive capacity result in malabsorption of protein, energy and micronutrients to varying degrees depending on construction of the anatomy. Bilio-pancreatic diversion ± duodenal switch (BPD ± DS) and Roux-en-Y gastric bypass (RYGB) combine these mechanisms. Micronutrient deficiency may well be a comorbidity of severe obesity in that it appears to increase in prevalence as the degree of obesity increases in populations who have had no prior bariatric surgery. This has been documented for alpha & beta carotene, beta cryptoxanthin, lutein/zeaxanthin, lycopene, total carotenoids, iron, selenium, vitamins A, C, D, B₆, B₁₂, and folic acid.⁶⁷⁻⁶⁹

Twenty-one observational studies and 2 RCTs have investigated a variety of micronutrients. These have compared serum levels in cohorts of patients treated with different procedures and have included RYGB, sleeve gastrectomy (SG), BPD ± DS, and adjustable gastric band procedures. The duration of follow-up was generally short, with 16 studies covering 1-3 years,⁶⁹⁻⁸² 3 studies 4-5 years⁸³⁻⁸⁵ and 1 study 7 years.⁸⁶ The study of longest duration documented no deficiency states in patients with restrictive procedures but no malabsorptive component; however, the others have documented an increased risk of deficiency of iron, copper, zinc, selenium, thiamine, folate, and Vitamins B₁₂ and D as compared with preoperative populations.

The proclivity of restrictive or malabsorptive procedures to exacerbate or create micronutrient deficiency states has been acknowledged by recommendations for supplementation published by the American Society for Metabolic and Bariatric Surgery and the Obesity Society.⁸⁷ For all bariatric surgery patients, a daily multiple vitamin/mineral supplement is recommended with 2 daily doses for patients with SG, RYGB, and BPD. For all patients, at least 3000 IU vitamin D daily is recommended to achieve serum 25-hydroxyvitamin D levels > 30 ng/mL; 2 mg copper daily; iron 45-60 mg from diet and supplements; and vitamin B₁₂ should be given as needed to maintain normal serum levels. All patients except those with BPD should take 1200-1500 mg calcium citrate daily. Evaluation of folic acid, iron and 25-hydroxyvitamin D should be done annually. Copper, zinc, selenium, and thiamine should be monitored when patients have specific findings to suggest deficiency. As with other chronic or home medications, these vitamin supplements should be continued or resumed in hospitalized patients.

Table 7. Evidence Summary Question 4: In Obese Patients Who Have Had a Malabsorptive or Restrictive Surgical Procedure, What Micronutrients Should Be Evaluated?

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|------------------------------------|---|--|--|--|--|
| Beckman et al, 2013 ⁷⁹ | Prospective cohort observation Small sample | Women with RYGB, N = 20 | Describe serum 25(OH)D changes and determine if FM loss and vitamin D intake are associated with changes in serum levels at 12 months after RYGB | 25(OH)D increased by 10 ± 2 ng/mL by 12 months 3 patients still had 25(OH)D < 20 ng/mL Weight, FM, BMI, and %EWL changes were associated with 25(OH)D change | |
| Aasheim et al, 2012 ⁹⁴ | Prospective nonrandomized trial Small sample | RYGB, n = 29 Lifestyle management, n = 24 | Assess change in vitamin status in patients taking vitamin supplements 1 year after RYGB vs lifestyle management controls | All vitamins similar between RYGB and control patients except vitamin A A lower in RYGB | |
| Dammis-Machado, 2012 ⁶⁹ | Retrospective record review Similar population Small sample | SG, N = 54 | Describe nutrient deficiencies before and 1, 3, 6, and 12 months after SG | At least 51% had a micronutrient deficiency preoperatively: <ul style="list-style-type: none"> • Vitamin D (83%) • Iron (29%) • Vitamin B6 (11%) • Vitamin B12 (9%) • Folate (6%) • Potassium (7%) By 12 months after SG, prevalence of deficiencies of the following nutrients increased: <ul style="list-style-type: none"> • Vitamin B₆ (17%) • Vitamin B₁₂ (17%) • Folate (14%) | Reduction in gastric acidity may be implicated postoperatively with vitamins B6, B12; folate deficiency may be due to food choices of patients |
| Gletsu-Miller, 2012 ⁹⁵ | Retrospective record review with Prospective cohort observation Small sample | RYGB, N = 136 | Describe number of RYGB patients with copper deficiency and associated hematological and neurological complaints over 12 months. | Prevalence of copper deficiency, 9.6% Incidence of copper deficiency, 18.8% Concomitant complications include anemia, leukopenia, and various neuromuscular abnormalities. | |
| Kehagias et al, 2011 ⁷⁶ | RCT of surgical procedure ITT analysis 5% attrition Small sample | Randomized to RYGB, N = 30 or SG, N = 30 | Describe perioperative safety and 3-year results after RYGB or SG | Preoperative nutrient deficiencies: RYGB vs SG, not significantly different 3 years postoperatively: Vitamin B ₁₂ deficiency in 7/29 (24%) in RYGB vs 1/28 (3.5%) in SG | |

(continued)

Table 7. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|------------------------------------|---|---|---|--|---|
| Leivonen et al, 2011 ⁷⁵ | Retrospective record review Small sample | Patients over age 60 years treated with SG, N = 12 vs patients < age 59 years, N = 43 | Evaluate differences in recovery, weight loss, and vitamin status 12 months after SG in younger vs older patients | Vitamin deficiencies: ● Not significantly different | |
| de Luis et al, 2011 ⁸⁵ | Retrospective record review No information on supplement adherence | BPD patients at baseline and 4 years postoperatively N = 65 | Evaluate influence of BPD on copper and zinc levels | Prevalence of copper deficiency: ● Preoperative, 67.8% ● 6 months, 76.9% ● 12 months, 76.9% ● 24 months, 87.7% ● 36 months, 87.7% ● 48 months, 90.7% Prevalence of zinc deficiency: ● Preoperative, 73.8% ● 6 months, 73.8% ● 12 months, 86.1% ● 24 months, 86.1% ● 36 months, 90.7% ● 48 months, 90.7% | Deficiency prevalence increases over time |
| Alasfar et al, 2011 ⁶⁸ | Controlled cohort observation No information on trace element intake or supplement use | Bariatric surgery patients, N = 66, BMI = 45.3 Nonobese controls, N = 44, BMI = 25.9 | Compare serum trace element (copper, zinc, selenium, magnesium) concentrations in preoperative bariatric surgery vs nonobese control subjects | Selenium concentration significantly lower in obese patients, $P < .001$ | |
| Balsa et al, 2011 ⁸³ | Cohort observation No information on trace element supplement use | RYGB, N = 52 BPD, N = 89 | Compare prevalence of copper and zinc deficiency in RYGB vs BPD patients | Prevalence of copper deficiency, RYGB vs BPD: ● Preoperative, 0% vs 0% ● 6 months, 0% vs 17% ● 12 months, 2% vs 13% ● 24 months, 0% vs 24% ● 48 months, 2% vs 22% ● 60 months, 2% vs 13% Prevalence of zinc deficiency, RYGB vs BPD: ● Preoperative, 12% vs 12% ● 6 months, 6% vs 69% ● 12 months, 2% vs 70% ● 24 months, 6% vs 74% ● 48 months, 15% vs 46% ● 60 months, 21% vs 45% | Copper and zinc deficiencies more common with BPD than RYGB, more prevalent over time |

(continued)

Table 7. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|------------------------------------|--|---|--|--|----------|
| Rosa et al, 2011 ⁹⁶ | Prospective bioavailability studies Small sample | RYGB, N = 9 | Describe iron and zinc plasma response to a tolerance test before and 3 months after RYGB. | Lower plasma zinc response ($P < .01$) and delayed response to iron intake after RYGB The total plasma iron concentration area over 4 hours was not different after surgery ($P > .05$) 24-hour urinary iron and zinc excretion did not change | |
| Gehrer et al, 2010 ⁷⁷ | Retrospective record review | 2004-2006 RYGB, N = 86, SG, N = 50 | Assess frequency of pre- and 3-year postoperative vitamin deficiencies and the success rate of their treatment | Preoperative and postoperative deficiencies: <ul style="list-style-type: none"> • Vitamin B₁₂ in RYGB (58%) vs SG (18%), $P < .0001$ • Vitamin D in RYGB (52%) vs SG (32%), $P < .01$ All deficiencies treatable | |
| Schouten et al, 2010 ⁸⁶ | RCT of laparoscopic band vs open VBG, cohort observation Diagnostic similarity Small sample may lack statistical power | Original study N = 100 2 and 7-years postsurgical data obtained from 91 (91%) with a mean follow-up of 84 months laparoscopic AGB N = 48 VBG N = 43 | Describe the long-term results of restrictive bariatric procedures including weight loss, long-term complications, comorbidities, reoperations, and vitamin status | No significant differences in levels of iron, zinc, folic acid or thiamine, vitamin B ₆ or B ₁₂ between laparoscopic AGB and VGB groups No vitamin deficiencies were present 7 years after restrictive bariatric surgical procedures | |
| Signori et al, 2010 ⁸⁰ | Retrospective record review | RYGB patients, N = 123 Recommended to take 1200-2000 IU vitamin D daily | Compare vitamin D status preoperatively vs 12 months post-RYGB | 25-OH D (ng/mL) 22.7 ± 9.9 vs 29.7 ± 14.1, preop vs 12 months post-RYGB, $P < .001$ | |
| Salle et al, 2010 ⁷⁸ | Retrospective record review | Bariatric surgery patients in Angers, France RYGB, N = 266 SG, N = 33 BPD-DS, N = 25 | Describe zinc and nutrition status before and 6, 12 and 24 months after RYGB, SG, DS | Preoperative: Zinc deficiency (9%) 24 months postoperatively: <ul style="list-style-type: none"> • RYGB (35%) • SG (18%) at 12 months • BPD-DS (92%) Iron deficiency: <ul style="list-style-type: none"> • RYGB (38%) • SG (25%) at 12 months • BPD-DS (58%) | |

(continued)

Table 7. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-----------------------------------|--|--|--|--|--|
| Goldner et al, 2009 ⁸¹ | RCT dose-response trial Small sample | Patients with RYGB and daily vitamin D supplements 800 IU, N = 13 2000 IU, N = 13 5000 IU, N = 15 | Dose-response trial to define dose of vitamin D supplement needed after RYGB | Preoperative serum 25(OH) D: <ul style="list-style-type: none"> 19.1 ± 9.9 vs 15.0 ± 9.3 vs 22.9 ± 10.3 nmol/L in 800 vs 2000 vs 5000 IU groups, <i>P</i> = .01 12 months post-RYGB: <ul style="list-style-type: none"> 27.5 ± 31.0 (n = 9), 800 IU 60.2 ± 37.4 (n = 9), 2000 IU 66.1 ± 42.2 (n = 10), 5000 IU No hypercalcaemia | Recommended to start all patients at 2000 IU/day |
| Coupaye et al, 2009 ⁷² | Prospective cohort Difference in BMI by treatment group Small sample, may lack statistical power No adjustment for inflammation or BMI group difference | Single center 70 consecutive patients who had undergone bariatric surgery AGB: N = 49, BMI 43 RYGB: N = 21, BMI 49 | Compare the vitamin and nutrition status before and 1 year after bariatric surgery in patients receiving systematized nutrition care | Deficiencies of thiamine, vitamin C, and iron in 38%, 47% and 43% of ABG patients preoperatively, not significantly worsened at 1 year In RYGB patients deficiencies of thiamine, iron, vitamin C were in 25%, 57%, and 47% preoperatively, with improvement in thiamine and vitamin C deficiencies at 1 year (12%* <i>P</i> < .05, 37%, 10%* <i>P</i> < .05 respectively) CRP and fibrinogen improved in both groups by 1 year | Vitamin supplements improved postoperative outcomes in RYGB patients |
| Carlin et al, 2009 ⁸² | RCT Small sample | Compare supplementation in female RYGB patients with 50,000 IU vitamin D weekly, N = 30 vs No vitamin D supplementation, N = 30 Both received 800 IU vitamin D and 1500 mg calcium daily | Evaluate the effectiveness of 50,000 IU vitamin D weekly to replenish vitamin D stores 1 year after RYGB | Baseline 25-hydroxyvitamin D: <ul style="list-style-type: none"> 19.7 ± 8.5 vs 18.5 ± 9.4 ng/mL, intervention vs control 12 Month 25-hydroxyvitamin D: <ul style="list-style-type: none"> 37.8 ± 15.6 vs 15.2 ± 7.5 ng/mL, intervention vs control (<i>P</i> < .001) Less decline in bone mineral density in treatment More frequent resolution of hypertension in treatment | |

(continued)

Table 7. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|-------------------------------------|--|---|--|---|--|
| Toh et al, 2009 ⁹⁷ | Retrospective record review Prognostic similarity Small sample No adjustment for supplement adherence rates, interaction of weight loss with vitamin status | Preoperative: n = 232 Postoperative: n = 148; RYGB = 103; SG = 46 | Describe prevalence of nutrient deficiencies in patients who present for bariatric surgery, compare with 12-month postoperative levels | Preoperatively <ul style="list-style-type: none"> Low 25-OH vitamin D in 57% Low iron in 15.7% High CRP in 58.5% Postoperatively, <ul style="list-style-type: none"> Low 25-OH vitamin D reduced to 30% in RYGB, 43% in SG patients Low iron unchanged High CRP improved to 13% and 17% in RYGB and SG patients Vitamin B₁₂ increased from 1% to 11% in RYGB Low RBC folate increased in RYGB from 1% to 12% | Increased B ₁₂ and folate deficiencies with RYGB suggest lack of adherence with supplements |
| Gasteyger et al, 2008 ⁷⁴ | Retrospective record review Small sample Adherence with vitamin supplements not evaluated | Single center Adult patients at 2 year follow-up after RYGB N = 137 (110 women; 27 men) Length of Roux limb: 100cm for BMI ≤ 48, 0 and 150 cm for BMI < 48.0 All patients received a multivitamin supplement 1-6 months after RYGB Supplementation with specific nutrients prescribed for values that fell below the reference range | Assess type, frequency, and pattern of the development of nutrition deficiencies over the first 24 months after RYGB, to determine the amount of supplements prescribed and to evaluate the cost of treatment. | <p>Patients requiring supplementation:</p> <ul style="list-style-type: none"> 3 months, 34% 6 months, 59% 24 months, 98% <p>Most frequent supplements:</p> <ul style="list-style-type: none"> Vitamin B₁₂, iron, calcium/vitamin D in 60% Folate in 40% Vitamin B₆, zinc, magnesium in 15% <p>Mean supplements per patient:</p> <ul style="list-style-type: none"> 24 months, 2.9 ± 1.4 Cost/year US\$417.96 | Nutrition deficiencies are common post RYGB despite multivitamin supplementation |
| Madan et al, 2006 ⁷¹ | Retrospective record review Small sample Incomplete data | All patients undergoing laparoscopic RYGB by 1 surgeon during a 6 month period. N = 100 Only about 30 patients with all vitamin levels at 12 months | Describe preoperative and 1-year post-RYGB vitamin and trace mineral levels | <p>Deficiencies, preoperative vs postoperative:</p> <ul style="list-style-type: none"> Vitamin A, 7% vs 16% Vitamin B 12.5% vs 0% Vitamin D, 40% vs 19% (<i>P</i> < .05) Zinc, 28% vs 36% Iron, 14% & 6% Selenium, 58% & 3% (<i>P</i> < .001) Folate, 2% vs 8% | Did not report thiamine levels |

(continued)

Table 7. (continued)

| Study | Study Design, Quality | Population, Setting, n | Study Objective | Results | Comments |
|------------------------------------|---|---|---|--|----------|
| Clements et al, 2006 ⁷⁰ | Retrospective record review | All patients with laparoscopic RYGB, 2002-2004 (N = 493) with 1- and 2-year follow-up, N = 141 | Evaluate prevalence of vitamin deficiency after RYGB | <p>Vitamin Deficiencies:</p> <ul style="list-style-type: none"> ● A (11%) ● C (34.6%) ● D (7%) ● Thiamine (18.3%) ● Riboflavin (13.6%) ● B₆ (17.6%) ● B₁₂ (3.6%) <p>No difference year 1 vs year 2 postoperatively</p> | |
| Skroubis et al, 2002 ⁸⁴ | Retrospective record review No data on adherence rates No data on baseline comorbid conditions Unclear data on number of subjects at each time point | University medical center in Greece N = 174 RYGB, N = 79 (BMI 45.6 ± 4.9) BPD, N = 95 (BMI 57.2 ± 6.1) | Compare nutrition complications and effectiveness of micronutrient supplementation after RYGB and BPD All patients received a multivitamin and mineral supplement and 2 g of calcium | <p>Iron deficiency:</p> <ul style="list-style-type: none"> ● Low iron and ferritin levels increased with both surgical procedures over time <p>Vitamin B₁₂ deficiency:</p> <ul style="list-style-type: none"> ● Increased with both surgical procedures from preop to 4 years postop with RYGB 33%, BPD 22% <p>Negligible incidence of hypocalbunemia</p> | |

AGB, adjustable gastric banding; BMI, body mass index; BPD, biliopancreatic diversion; CRP, C-reactive protein; DS, duodenal switch; EWL, excess weight loss; FM, fat mass; ITT, intention to treat analysis; IU, international unit; RCT, randomized controlled trial; RYGB, Roux-en-Y gastric bypass; SG, sleeve gastrectomy; VBG, vertical-banded gastroplasty; 25(OH)D = 25-hydroxyvitamin D.

Table 8. GRADE Table Question 4: In Obese Patients Who Have Had a Malabsorptive Surgical Procedure, What Micronutrients Should Be Evaluated?

| Comparison | Outcome/Nutrient Deficiency | Quantity, Type Evidence | Finding | Final GRADE | Overall Evidence GRADE |
|---|-----------------------------|-------------------------|--|-------------|------------------------|
| Preoperative to postoperative RYGB or BPD | Copper | 3 OBS | Increased ^{83,85,95} | Low | Low |
| | Zinc | 3 OBS | Increased ^{83,85} | Low | |
| | Iron | 3 OBS | Increased ^{84,97} | Very low | |
| | Selenium | 1 OBS | | Low | |
| | Thiamine | 1 OBS | Increased ⁷² | Low | |
| | Folic acid | 1 OBS | Increased ⁹⁷ | Low | |
| | Vitamin B ₁₂ | 2 OBS | Increased ^{84,97} | Low | |
| | Vitamin D | 5 OBS, 2 RCT | Increased with supplements decreased ⁹⁷ | Low | |

BPD = biliopancreatic diversion; OBS = observational study; RCT, randomized controlled trial; RYGB = Roux-en-Y gastric bypass.

Compliance with supplement ingestion has been variable, with BPD \pm DS 55%, RYGB 25%.⁸⁸ Patient follow-up with bariatric surgical programs, and hence routine surveillance of nutrition parameters, tends to diminish with time duration after the surgical procedure. The severity and prevalence of deficiency appears to increase with the interval of time after the procedure as well as with the degree of malabsorption induced by the procedure. Data evaluating micronutrient status in patients in the decades following bariatric surgical intervention are not available.

A.S.P.E.N. Board of Directors Providing Final Approval

Deborah A. Andris, MSN, APNP; Phil Ayers, PharmD, BCNSP, FASHP; Albert Baroccas, MD, FACS, FASPEN; Praveen S. Goday, MBBS, CNSC; Carol Ireton-Jones, PhD, RD, LD, CNSD; Tom Jaksic, MD, PhD; Lawrence A. Robinson, BS, MS, PharmD; Gordon Sacks, PharmD, BCNSP, FCCP; Daniel Teitelbaum, MD; Charles W. Van Way III, MD, FASPEN.

A.S.P.E.N. Clinical Guidelines Editorial Board

Charlene Compher, PhD, RD, CNSC, LDN, FADA, FASPEN; Nancy Allen, MS, MLS, RD; Joseph I. Boullata, PharmD, RPh, BCNSP; Carol L. Braunschweig, PhD, RD; Donald E. George, MD; Edwin Simpser, MD; and Patricia A. Worthington, MSN, RN, CNSN.

Acknowledgments

This unfunded project was completed by authors and reviewers using their time as volunteers.

References

- Hoven AD, A.M.A. Obesity as a disease state: it's about patient health. 2013. Available at: <http://www.ama-assn.org/ama/pub/ama-president-blog/road-dr-hoven.page?plckController=Blog&plckBlogPage=BlogViewPost&UID=a0ab4522-eadf-4594-808e-508e4fa2e811&plckPostId=Blog%3Aa0ab4522-eadf-4594-808e-508e4fa2e811Post%3A13a0adac-b70d-431d-a262-999a7a6b427d&plckScript=blogScript&plckElementId=blogDest>.
- Healey KL, Bines JE, Thomas SL, et al. Morphological and functional changes in the colon after massive small bowel resection. *J Pediatr Surg.* 2010;45(8):1581-1590.
- Flegal KM, Carroll MD, Kit BK, Ogden CL. Prevalence of obesity and trends in the distribution of body mass index among US adults, 1999-2010. *JAMA.* 2012;307(5):491-497.
- Surgery ASfMaB. New evidence prompts update to metabolic and bariatric surgery clinical guidelines. 2013. Available at: <http://s3.amazonaws.com/publicASMBS/top5/April2013/CPG.Guidelines.News.April.2013.pdf>
- Guidelines for the use of parenteral and enteral nutrition in adult and pediatric patients. *JPEN J Parenter Enteral Nutr.* Jan-Feb 2002;26(1 suppl):1SA-138SA.
- Guidelines for use of total parenteral nutrition in the hospitalized adult patient. A.S.P.E.N. Board of Directors. *JPEN J Parenter Enteral Nutr.* Sep-Oct 1986;10(5):441-445.
- Arsenault D, Brenn M, Kim S, et al. A.S.P.E.N. Clinical guidelines: hyperglycemia and hypoglycemia in the neonate receiving parenteral nutrition. *JPEN J Parenter Enteral Nutr.* Jan 2012;36(1):81-95.
- August DA, Huhmann MB. A.S.P.E.N. clinical guidelines: nutrition support therapy during adult anticancer treatment and in hematopoietic cell transplantation. *JPEN J Parenter Enteral Nutr.* Sep-Oct 2009;33(5):472-500.
- Brown RO, Compher C. A.S.P.E.N. clinical guidelines: nutrition support in adult acute and chronic renal failure. *JPEN J Parenter Enteral Nutr.* Jul-Aug 2010;34(4):366-377.
- Sabery N, Duggan C. A.S.P.E.N. clinical guidelines: nutrition support of children with human immunodeficiency virus infection. *JPEN J Parenter Enteral Nutr.* Nov-Dec 2009;33(6):588-606.
- Jaksic T, Hull MA, Modi BP, Ching YA, George D, Compher C. A.S.P.E.N. Clinical guidelines: nutrition support of neonates supported with extracorporeal membrane oxygenation. *JPEN J Parenter Enteral Nutr.* May-Jun 2010;34(3):247-253.
- Jesuit C, Dillon C, Compher C, Lenders CM. A.S.P.E.N clinical guidelines: nutrition support of hospitalized pediatric patients with obesity. *JPEN J Parenter Enteral Nutr.* Jan-Feb 2010;34(1):13-20.
- McClave SA, Martindale RG, Vanek VW, et al. Guidelines for the provision and assessment of nutrition support therapy in the adult critically ill patient: Society of Critical Care Medicine (SCCM) and American Society for Parenteral and Enteral Nutrition (A.S.P.E.N.). *JPEN J Parenter Enteral Nutr.* May-Jun 2009;33(3):277-316.

14. Mehta NM, Compher C. A.S.P.E.N. Clinical guidelines: nutrition support of the critically ill child. *JPEN J Parenter Enteral Nutr.* May-Jun 2009;33(3):260-276.
15. Mueller C, Compher C, Ellen DM. A.S.P.E.N. clinical guidelines: nutrition screening, assessment, and intervention in adults. *JPEN J Parenter Enteral Nutr.* Jan 2011;35(1):16-24.
16. Balshem H, Helfand M, Schunemann HJ, et al. GRADE guidelines: 3. Rating the quality of evidence. *J Clin Epidemiol.* Apr 2011;64(4):401-406.
17. Guyatt GH, Oxman AD, Kunz R, et al. GRADE guidelines: 2. Framing the question and deciding on important outcomes. *J Clin Epidemiol.* Apr 2011;64(4):395-400.
18. Guyatt GH, Oxman AD, Vist G, et al. GRADE guidelines: 4. Rating the quality of evidence—study limitations (risk of bias). *J Clin Epidemiol.* Apr 2011;64(4):407-415.
19. Guyatt G, Oxman AD, Akl EA, et al. GRADE guidelines: 1. Introduction- GRADE evidence profiles and summary of findings tables. *J Clin Epidemiol.* Apr 2011;64(4):383-394.
20. Druyan ME, Compher C, Boullata JI, et al. Clinical guidelines for the use of parenteral and enteral nutrition in adult and pediatric patients: applying the GRADE system to development of A.S.P.E.N. clinical guidelines. *JPEN J Parenter Enteral Nutr.* Jan 2012;36(1):77-80.
21. Hoffmann M, Lefering R, Gruber-Rathmann M, Rueger JM, Lehmann W. The impact of BMI on polytrauma outcome. *Injury.* Feb 2012;43(2):184-188.
22. Westerly BD, Dabbagh O. Morbidity and mortality characteristics of morbidly obese patients admitted to hospital and intensive care units. *J Crit Care.* Apr 2011;26(2):180-185.
23. Hutagalung R, Marques J, Kobylka K, et al. The obesity paradox in surgical intensive care unit patients. *Intensive Care Med.* Nov 2011;37(11):1793-1799.
24. Evans DC, Stawicki SP, Davido HT, Eiferman D. Obesity in trauma patients: correlations of body mass index with outcomes, injury patterns, and complications. *Amer Surg.* Aug 2011;77(8):1003-1008.
25. Serrano PE, Khuder SA, Fath JJ. Obesity as a risk factor for nosocomial infections in trauma patients. *J Am Coll Surg.* Jul 2010;211(1):61-67.
26. Wurzinger B, Dunser MW, Wohlmuth C, et al. The association between body-mass index and patient outcome in septic shock: a retrospective cohort study. *Wiener klinische Wochenschrift.* Jan 2010;122(1-2):31-36.
27. Pieracci FM, Hydo L, Pomp A, Eachempati SR, Shou J, Barie PS. The relationship between body mass index and postoperative mortality from critical illness. *Obes Surg.* May 2008;18(5):501-507.
28. Morris AE, Stapleton RD, Rubenfeld GD, Hudson LD, Caldwell E, Steinberg KP. The association between body mass index and clinical outcomes in acute lung injury. *Chest.* Feb 2007;131(2):342-348.
29. Tremblay A, Bandi V. Impact of body mass index on outcomes following critical care. *Chest.* Apr 2003;123(4):1202-1207.
30. Nafiu OO, Ramachandran SK, Wagner DS, Campbell DA Jr, Stanley JC. Contribution of body mass index to postoperative outcome in minority patients. *J Hosp Med.* Feb 2012;7(2):117-123.
31. Park B, Dargon P, Binette C, et al. Obesity is not an independent risk factor for adverse perioperative and long-term clinical outcomes following open AAA repair or EVAR. *Vasc Endovascular Surg.* Oct 2011;45(7):607-613.
32. Ray DE, Matchett SC, Baker K, Wasser T, Young MJ. The effect of body mass index on patient outcomes in a medical ICU. *Chest.* Jun 2005;127(6):2125-2131.
33. Brown CV, Neville AL, Rhee P, Salim A, Velmahos GC, Demetriades D. The impact of obesity on the outcomes of 1,153 critically injured blunt trauma patients. *J Trauma.* Nov 2005;59(5):1048-1051, discussion 1051.
34. O'Brien JM. Obesity-related excess mortality rate in an adult intensive care unit: a risk-adjusted matched cohort study. *Crit Care Med.* Sep 2004;32(9):1980.
35. Aldawood A, Arabi Y, Dabbagh O. Association of obesity with increased mortality in the critically ill patient. *Anaesth Intensive Care.* Oct 2006;34(5):629-633.
36. Frat JP, Gissot V, Ragot S, et al. Impact of obesity in mechanically ventilated patients: a prospective study. *Intensive Care Med.* Nov 2008;34(11):1991-1998.
37. Newell MA, Bard MR, Goettler CE, et al. Body mass index and outcomes in critically injured blunt trauma patients: weighing the impact. *J Am Coll Surg.* May 2007;204(5):1056-1061, discussion 1062-1054.
38. Peake SL, Moran JL, Ghelani DR, Lloyd AJ, Walker MJ. The effect of obesity on 12-month survival following admission to intensive care: a prospective study. *Crit Care Med.* Dec 2006;34(12):2929-2939.
39. Duane TM, Dechert T, Aboutanos MB, Malhotra AK, Ivatury RR. Obesity and outcomes after blunt trauma. *J Trauma.* Nov 2006;61(5):1218-1221.
40. Alban RF, Lyass S, Margulies DR, Shabot MM. Obesity does not affect mortality after trauma. *Amer Surg.* Oct 2006;72(10):966-969.
41. Winkelman C, Maloney B. Obese ICU patients: resource utilization and outcomes. *Clin Nurs Res.* Nov 2005;14(4):303-323, discussion 324-306.
42. O'Brien JM Jr, Phillips GS, Ali NA, Lucarelli M, Marsh CB, Lemeshow S. Body mass index is independently associated with hospital mortality in mechanically ventilated adults with acute lung injury. *Crit Care Med.* Mar 2006;34(3):738-744.
43. Garrouste-Orgeas M, Troche G, Azoulay E, et al. Body mass index. An additional prognostic factor in ICU patients. *Intensive Care Med.* Mar 2004;30(3):437-443.
44. Abhyankar S, Leishear K, Callaghan FM, Demner-Fushman D, McDonald CJ. Lower short- and long-term mortality associated with overweight and obesity in a large cohort study of adult intensive care unit patients. *Crit Care.* Dec 18 2012;16(6):R235.
45. Martino JL, Stapleton RD, Wang M, et al. Extreme obesity and outcomes in critically ill patients. *Chest.* Nov 2011;140(5):1198-1206.
46. Sakr Y, Madl C, Filipescu D, et al. Obesity is associated with increased morbidity but not mortality in critically ill patients. *Intensive Care Med.* Nov 2008;34(11):1999-2009.
47. Dossett LA, Dageforde LA, Swenson BR, et al. Obesity and site-specific nosocomial infection risk in the intensive care unit. *Surg Infect (Larchmt).* Apr 2009;10(2):137-142.
48. Duchesne JC, Schmiege RE Jr, Simmons JD, Islam T, McGinness CL, McSwain NE Jr. Impact of obesity in damage control laparotomy patients. *J Trauma.* Jul 2009;67(1):108-112, discussion 112-104.
49. Das SR, Alexander KP, Chen AY, et al. Impact of body weight and extreme obesity on the presentation, treatment, and in-hospital outcomes of 50,149 patients with ST-segment elevation myocardial infarction results from the NCDR (National Cardiovascular Data Registry). *J Am Coll Cardiol.* Dec 13 2011;58(25):2642-2650.
50. Frankenfield DC, Coleman A, Alam S, Cooney RN. Analysis of estimation methods for resting metabolic rate in critically ill adults. *JPEN J Parenter Enteral Nutr.* Jan-Feb 2009;33(1):27-36.
51. Frankenfield DC, Ashcraft CM, Galvan DA. Prediction of resting metabolic rate in critically ill patients at the extremes of body mass index. *JPEN J Parenter Enteral Nutr.* May-Jun 2013;37(3):361-367.
52. Frankenfield DC, Ashcraft CM, Galvan DA. Longitudinal prediction of metabolic rate in critically ill patients. *JPEN J Parenter Enteral Nutr.* Nov 2012;36(6):700-712.
53. Frankenfield D. Validation of an equation for resting metabolic rate in older obese, critically ill patients. *JPEN J Parenter Enteral Nutr.* Mar 2011;35(2):264-269.
54. Boullata J, Williams J, Cottrell F, Hudson L, Compher C. Accurate determination of energy needs in hospitalized patients. *J Am Diet Assoc.* Mar 2007;107(3):393-401.
55. Anderegg BA, Worrall C, Barbour E, Simpson KN, Delege M. Comparison of resting energy expenditure prediction methods with measured resting energy expenditure in obese, hospitalized adults. *JPEN J Parenter Enteral Nutr.* Mar-Apr 2009;33(2):168-175.
56. Frankenfield DC, Rowe WA, Smith JS, Cooney RN. Validation of several established equations for resting metabolic rate in obese and nonobese people. *J Am Diet Assoc.* Sep 2003;103(9):1152-1159.

57. Dobratz JR, Sibley SD, Beckman TR, et al. Predicting energy expenditure in extremely obese women. *JPEN J Parenter Enteral Nutr.* May-Jun 2007;31(3):217-227.
58. Mifflin MD, St Jeor ST, Hill LA, Scott BJ, Daugherty SA, Koh YO. A new predictive equation for resting energy expenditure in healthy individuals. *Am J Clin Nutr.* Feb 1990;51(2):241-247.
59. Choban PS, Burge JC, Scales D, Flancbaum L. Hypoenergetic nutrition support in hospitalized obese patients: a simplified method for clinical application. *Am J Clin Nutr.* Sep 1997;66(3):546-550.
60. Burge JC, Goon A, Choban PS, Flancbaum L. Efficacy of hypocaloric total parenteral nutrition in hospitalized obese patients: a prospective, double-blind randomized trial. *JPEN J Parenter Enteral Nutr.* May-Jun 1994;18(3):203-207.
61. Dickerson RN, Boschert KJ, Kudsk KA, Brown RO. Hypocaloric enteral tube feeding in critically ill obese patients. *Nutrition.* Mar 2002;18(3):241-246.
62. Dickerson RN, Medling TL, Smith AC, et al. Hypocaloric, high-protein nutrition therapy in older vs younger critically ill patients with obesity. *JPEN J Parenter Enteral Nutr.* May-Jun 2013;37(3):342-351.
63. Hamilton C, Dasari V, Shatnawei A, Lopez R, Steiger E, Seidner D. Hypocaloric home parenteral nutrition and nutrition parameters in patients following bariatric surgery. *Nutr Clin Pract.* Oct 2011;26(5):577-582.
64. Dickerson RN, Rosato EF, Mullen JL. Net protein anabolism with hypocaloric parenteral nutrition in obese stressed patients. *Am J Clin Nutr.* Dec 1986;44(6):747-755.
65. Alberda C, Gramlich L, Jones N, et al. The relationship between nutritional intake and clinical outcomes in critically ill patients: results of an international multicenter observational study. *Intensive Care Med.* Oct 2009;35(10):1728-1737.
66. Choban PS, Dickerson RN. Morbid obesity and nutrition support: is bigger different? *Nutr Clin Pract.* Aug 2005;20(4):480-487.
67. Kimmons JE, Blanck HM, Tohill BC, Zhang J, Khan LK. Associations between body mass index and the prevalence of low micronutrient levels among US adults. *Med Gen Med.* 2006;8(4):59.
68. Alasfar F, Ben-Nakhi M, Khourshed M, Kehinde EO, Alsaleh M. Selenium is significantly depleted among morbidly obese female patients seeking bariatric surgery. *Obes Surg.* Nov 2011;21(11):1710-1713.
69. Damms-Machado A, Friedrich A, Kramer KM, et al. Pre- and postoperative nutritional deficiencies in obese patients undergoing laparoscopic sleeve gastrectomy. *Obes Surg.* Jun 2012;22(6):881-889.
70. Clements RH, Katasani VG, Palepu R, et al. Incidence of vitamin deficiency after laparoscopic Roux-en-Y gastric bypass in a university hospital setting. *Amer Surg.* Dec 2006;72(12):1196-1202, discussion 1203-1204.
71. Madan AK, Orth WS, Tichansky DS, Ternovits CA. Vitamin and trace mineral levels after laparoscopic gastric bypass. *Obes Surg.* May 2006;16(5):603-606.
72. Coupaye M, Puchaux K, Bogard C, et al. Nutritional consequences of adjustable gastric banding and gastric bypass: a 1-year prospective study. *Obes Surg.* Jan 2009;19(1):56-65.
73. Toh SY, Zarshenas N, Jorgensen J. Prevalence of nutrient deficiencies in bariatric patients. *Nutrition.* Nov-Dec 2009;25(11-12):1150-1156.
74. Gasteyger C, Suter M, Gaillard RC, Giusti V. Nutritional deficiencies after Roux-en-Y gastric bypass for morbid obesity often cannot be prevented by standard multivitamin supplementation. *Am J Clin Nutr.* May 2008;87(5):1128-1133.
75. Leivonen MK, Juuti A, Jaser N, Mustonen H. Laparoscopic sleeve gastrectomy in patients over 59 years: early recovery and 12-month follow-up. *Obes Surg.* Aug 2011;21(8):1180-1187.
76. Kehagias I, Karamanakos SN, Argentou M, Kalfarentzos F. Randomized clinical trial of laparoscopic Roux-en-Y gastric bypass versus laparoscopic sleeve gastrectomy for the management of patients with BMI < 50 kg/m². *Obes Surg.* Nov 2011;21(11):1650-1656.
77. Gehrler S, Kern B, Peters T, Christoffel-Courtin C, Peterli R. Fewer nutrient deficiencies after laparoscopic sleeve gastrectomy (LSG) than after laparoscopic Roux-Y-gastric bypass (LRYGB)-a prospective study. *Obes Surg.* Apr 2010;20(4):447-453.
78. Salle A, Demarsy D, Poirier AL, et al. Zinc deficiency: a frequent and underestimated complication after bariatric surgery. *Obes Surg.* Dec 2010;20(12):1660-1670.
79. Beckman LM, Earthman CP, Thomas W, et al. Serum 25(OH) vitamin D concentration changes after Roux-en-Y gastric bypass surgery [published online ahead of print March 21, 2013]. *Obesity (Silver Spring).* doi:10.1002/oby.20464.
80. Signori C, Zalesin KC, Franklin B, Miller WL, McCullough PA. Effect of gastric bypass on vitamin D and secondary hyperparathyroidism. *Obes Surg.* Jul 2010;20(7):949-952.
81. Goldner WS, Stoner JA, Lyden E, et al. Finding the optimal dose of vitamin D following Roux-en-Y gastric bypass: a prospective, randomized pilot clinical trial. *Obes Surg.* Feb 2009;19(2):173-179.
82. Carlin AM, Rao DS, Yager KM, Parikh NJ, Kapke A. Treatment of vitamin D depletion after Roux-en-Y gastric bypass: a randomized prospective clinical trial. *Surg Obes Relat Dis.* Jul-Aug 2009;5(4):444-449.
83. Balsa JA, Botella-Carretero JJ, Gomez-Martin JM, et al. Copper and zinc serum levels after derivative bariatric surgery: differences between Roux-en-Y gastric bypass and biliopancreatic diversion. *Obes Surg.* Jun 2011;21(6):744-750.
84. Skroubis G, Sakellaropoulos G, Pougouras K, Mead N, Nikiforidis G, Kalfarentzos F. Comparison of nutritional deficiencies after Roux-en-Y gastric bypass and after biliopancreatic diversion with Roux-en-Y gastric bypass. *Obes Surg.* Aug 2002;12(4):551-558.
85. de Luis DA, Pacheco D, Izaola O, Terroba MC, Cuellar L, Martin T. Zinc and copper serum levels of morbidly obese patients before and after biliopancreatic diversion: 4 years of follow-up. *J Gastrointest Surg.* Dec 2011;15(12):2178-2181.
86. Schouten R, Wiryasaputra DC, van Dielen FM, van Gemert WG, Greve JW. Long-term results of bariatric restrictive procedures: a prospective study. *Obes Surg.* Dec 2010;20(12):1617-1626.
87. Mechanick JI, Youdim A, Jones DB, et al. Clinical practice guidelines for the perioperative nutritional, metabolic, and nonsurgical support of the bariatric surgery patient—2013 update: cosponsored by American Association of Clinical Endocrinologists, the Obesity Society, and American Society for Metabolic & Bariatric Surgery. *Surg Obes Relat Dis.* Mar-Apr 2013;9(2):159-191.
88. Aasheim ET, Bjorkman S, Sovik TT, et al. Vitamin status after bariatric surgery: a randomized study of gastric bypass and duodenal switch. *Am J Clin Nutr.* Jul 2009;90(1):15-22.
89. Nelson J, Billeter AT, Seifert B, et al. Obese trauma patients are at increased risk of early hypovolemic shock: a retrospective cohort analysis of 1,084 severely injured patients. *Crit Care.* May 8 2012;16(3):R77.
90. Nasraway SA Jr, Albert M, Donnelly AM, Ruthazer R, Shikora SA, Saltzman E. Morbid obesity is an independent determinant of death among surgical critically ill patients. *Crit Care Med.* Apr 2006;34(4):964-970, quiz 971.
91. Nafu OO, Shanks AM, Hayanga AJ, Tremper KK, Campbell DA Jr. The impact of high body mass index on postoperative complications and resource utilization in minority patients. *J Natl Med Assoc.* Jan 2011;103(1):9-15.
92. Kross EK, Sena M, Schmidt K, Stapleton RD. A comparison of predictive equations of energy expenditure and measured energy expenditure in critically ill patients. *J Crit Care.* Jun 2012;27(3):321e5-2.
93. Alves VG, da Rocha EE, Gonzalez MC, da Fonseca RB, Silva MH, Chiesa CA. Assessment of resting energy expenditure of obese patients: comparison of indirect calorimetry with formulae. *Clin Nutr.* Jun 2009;28(3):299-304.

94. Aasheim ET, Johnson LK, Hofso D, Bohmer T, Hjelmsaeth J. Vitamin status after gastric bypass and lifestyle intervention: a comparative prospective study. *Surg Obes Relat Dis*. Mar-Apr 2012;8(2):169-175.
95. Gletsu-Miller N, Broderius M, Frediani JK, et al. Incidence and prevalence of copper deficiency following Roux-en-Y gastric bypass surgery. *Int J Obes (Lond)*. Mar 2012;36(3):328-335.
96. Rosa FT, de Oliveira-Penaforte FR, de Arruda Leme I, Padovan GJ, Ceneviva R, Marchini JS. Altered plasma response to zinc and iron tolerance test after Roux-en-Y gastric bypass. *Surg Obes Relat Dis*. May-Jun 2011;7(3):309-314.
97. Toh SY, Zarshenas N, Jorgensen J. Prevalence of nutrient deficiencies in bariatric patients. *Nutr* 2009;25(11-12):1150-1156.