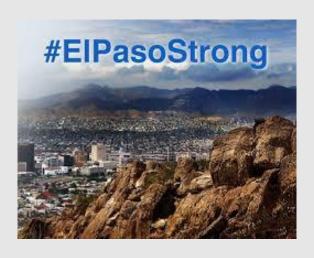
NUTRITIONAL SUPPORT OF THE TRAUMA PATIENT



Alan H. Tyroch, FACS, FCCM Professor and Chair of Surgery Trauma Medical Director

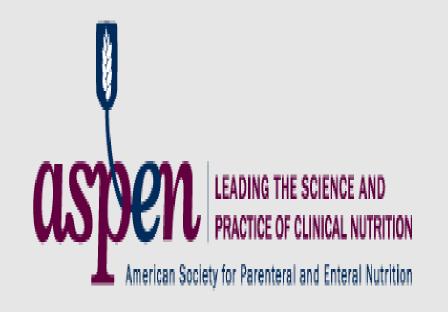




KEY POINTS

- Nutritional assessment on ICU admission
 - Energy requirements
 - Protein requirements
- Initiate enteral nutrition (EN) 24-48 hours after admission.
 - Reach goal < a week from admission (preferably much sooner)
- Take steps to reduce aspiration risk and improve tolerance to feeding.
- Do NOT us gastric residual volumes to monitor patients on EN.

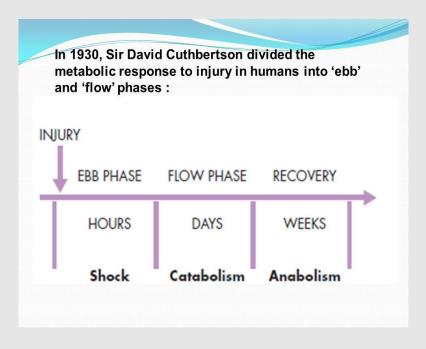
FEED the GUT!
FEED the GUT!
FEED the GUT!
(I hate TPN!)

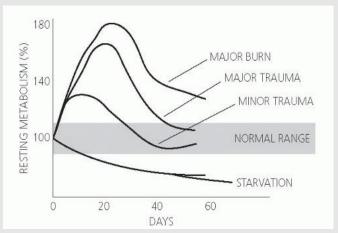


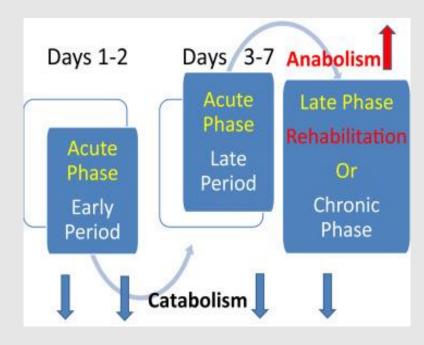


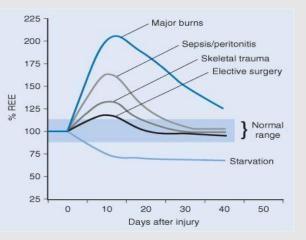


METABOLIC RESPONSE









NUTRITIONAL ASSESSMENT

Comorbid **Conditions Aspiration Function** Risk of GI Tract

DO NOT USE

Traditional Serum Markers 24-hour Urine Urea Nitrogen

BEST TEST
Indirect Calorimetry

CT Scan
Ultrasound

PATIENT ASSESSMENT

Patient history

Physical exam

- Anthropometrics
 - Height
 - Weight (ideal & dry)
 - BMI

Usual labs



NUTRIC Score

The NUTRIC Score is designed to quantify the risk of critically ill patients developing adverse events that may be modified by aggressive nutrition therapy. The score, of 1-10, is based on 6 variables that are explained below in Table 1. The scoring system is shown in Tables 2 and 3.

Table 1: NUTRIC Score variables

Variable	Range	Points
Age	<50	0
	50 - <75	1
	>75	2
APACHE II	<15	0
	15 - <20	1
	20-28	2
	≥28	3
SOFA	<6	0
	6 - <10	1
	≥10	2
Number of Co-morbidities	0-1	0
	≥2	1
Days from hospital to ICU admission	0-<1	0
	≥1	1
IL-6	0 - <400	0
	≥ 400	1

Table 2: NUTRIC Score scoring system: if IL-6 available

Sum of points	Category	Explanation
6-10	High Score	 Associated with worse clinical outcomes (mortality, ventilation). These patients are the most likely to benefit from aggressive nutrition therapy.
0-5	Low Score	These patients have a low malnutrition risk.

Table 3. NUTRIC Score scoring system: If no IL-6 available*

Sum of points	Category	Explanation
5-9	High Score	Associated with worse clinical outcomes (mortality, ventilation). These patients are the most likely to benefit from aggressive
		nutrition therapy.
0-4	Low Score	These patients have a low malnutrition risk.

^{*}It is acceptable to not include IL-6 data when it is not routinely available; it was shown to contribute very little to the overall prediction of the NUTRIC score. 2

December 16th 2015

	Nutritional status	Disease/surgery severity	Age
0	Normal	Normal	<70
1	Weight loss >5%/3 months or Food intake <75%	Includes chronic disease, hip fracture, cancer, minor surgery	≥70
2	Weight loss >5%/2 months or Food intake <50% or BMI 18.5-20.5	Includes major surgery, myocardial infarction, pneumonia, lymphoma, leukemia	
3	Weight loss >5%/1 month (or >15%/3 months) or Food intake <25% or BMI <18.5	Includes head trauma, transplantation, intensive care patients	

BMI: body mass index. The Nutritional Risk Score (NRS) is calculated by adding 3 different components: nutritional status + disease/surgery severity + age. Only the more severe contribution to the overall score of each of these 3 elements is considered in the overall score.

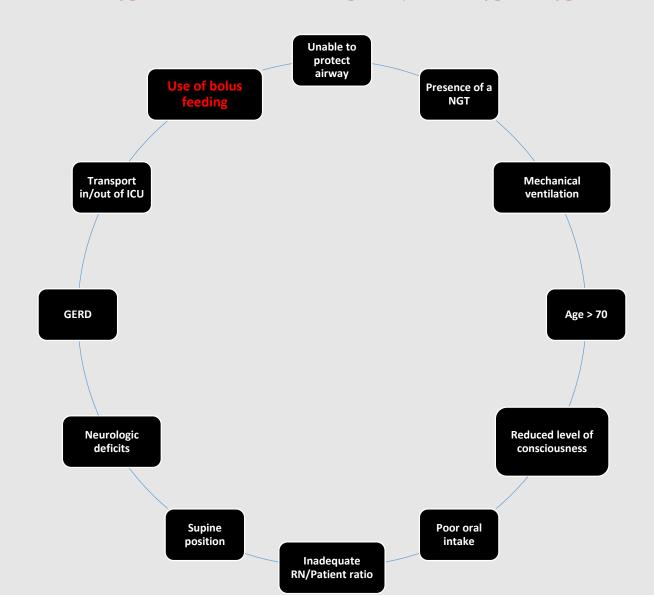
¹ Heyland DK, Dhaliwal R, Jiang X, Day AG. Identifying critically ill patients who benefit the most from nutrition therapy: the development and initial validation of a novel risk assessment tool. Critical Care. 2011;15(6):R268.
²Rahman A, Hasan RM, Agarwala R, Martin C, Day AG, Heyland DK. Identifying critically-ill patients who will benefit most from nutritional therapy: Further validation of the "modified NUTRIC" nutritional risk assessment tool. Clin Nutr. 2015. [Epub ahead of print]

ESPEN RECOMMENDATIONS

- Predicted ICU stay > 2 days.
- Mechanical ventilation
- Active infection
- Underfed > 5 days
- Presenting with a severe chronic disease

"Medical nutrition therapy shall be considered for all patients staying in the ICU, mainly for more than 48 hours."

ASPIRATION RISKS



NUTRITIONAL SUPPORT Inadequacy

< 50% of patients reach their target goal of energy intake.</p>

We provide only 60%-80% of energy requirements.

Patients receive ~80% of what is prescribed.

Feeding is held for too long and for inappropriate reasons

INITIATION OF NUTRITIONAL SUPPORT

Calculate energy requirements: 25-30 kcal/kg/day

- Protein Provision: 1.2-2.0 g/kg/day (more for major trauma & burns)
 - ESPEN: 1.3 g/kg/day
- Start tube feeds at 20cc/hr and increase by 10cc/hr every 4-6 hours until goal (conservative approach)

- Provide free water at 30cc/kg (NS if TBI)
 - Enteral nutrition is ~85% water

SPECIAL POPULATIONS Renal Failure

Standard enteral formula

Use dry weight for calculation

Energy provision: 25-30 kcal/kg/day

Protein provision: 1.2-2 g/kg actual body weight

- Hemodialysis or CRRT:
 - Increase protein to 2.5 g/kg/day

SPECIAL POPULATIONS Hepatic Failure

 Use dry weight to determine energy and protein requirements in those with cirrhosis and hepatic failure.

- Avoid restricting protein.
- Standard enteral formula.
 - Branched-chain amino acid formulas had no effect on coma grade

RATIONAL OF ENTERAL VERSUS PARENTERAL NUTRITION

- Reduction of infectious morbidity:
 - Pneumonia
 - CAUTI
 - Abdominal abscess (trauma)
- Reduced ICU LOS.

Minimal impact on mortality.

ENTERAL VERSUS PARENTERAL NUTRITION Infectious Complications

ASPEN

Risk Ratio Risk Ratio Events Total Events Total Weight M-H, Random, 95% Cl Year M-H. Random, 95% CI Study or Subgroup Adams 1986 0.88 [0.60, 1.30] 1986 Young 1987 1.03 [0.31, 3.39] 1987 Peterson 1988 0.30 [0.07, 1.25] 1988 Moore 1989 0.47 [0.19, 1.19] 1989 Kudsk 1992 0.39 [0.20, 0.78] 1992 Kalfarentzos 1997 20 11.2% 0.56 [0.23, 1.32] 1997 21 13.5% Woodcock 2001 0.72 [0.34, 1.52] 2001 Casas 2007 0.33 [0.04, 2.73] 2007 Chen 2011 49 10.5% 0.28 [0.11, 0.69] 2011 Total (95% CI) 249 247 100.0% 0.56 [0.39, 0.79] 101 Total events Heterogeneity: $Tau^2 = 0.09$; $Chi^2 = 12.10$, df = 8 (P = 0.15); $I^2 = 34\%$ Test for overall effect: Z = 3.26 (P = 0.001) Favors EN Favors PN

ESPEN

	EEN	4	EP			Risk Ratio		Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	Year	M-H, Random, 95% CI
2.1.1 ICU studies				200100				11 110000011111 Dental Control
Kompan 2004	9	27	16	25	9.5%	0.52 [0.28, 0.96]	2004	
Lam 2008	10	41	25	41	9.8%	0.40 [0.22, 0.72]	2008	
Altintas 2011	7	30	13	41	7.0%	0.74 [0.33, 1.62]	2011	
Justo Meirelles 2011	2	12	4	10	2.7%	0.42 [0.10, 1.82]	2011	
Harvey 2014	194	1197	194	1191	18.3%	0.99 [0.83, 1.19]	2014	+
Reignier 2017	173	1202	194	1208	18.2%	0.90 [0.74, 1.08]	2017	•
Subtotal (95% CI)		2509		2516	65.6%	0.75 [0.57, 0.98]		•
Total events	395		446					
Heterogeneity: Tau* = 1	0.05; Chi ²	= 12.66	6, df = 50	P = 0.03	3); $l^2 = 61$	%		
Test for overall effect: 2	Z = 2.14 (P	= 0.03	0					
2.1.2 Studies with unc	lear prop	ortion	of ICU pa	tients				
Aiko 2001	0	13	1	11	0.7%	0.29 [0.01, 6.38]	2001	
Bozzetti 2001	25	159	42	158	12.7%	0.59 [0.38, 0.92]	2001	(
Gupta 2003	1	8	2	9	1.3%	0.56 [0.06, 5.09]	2003	1
Eckerwall 2006	3	23	0	25	0.8%	7.58 [0.41, 139.32]	2006	,
Petrov 2006	11	35	27	34	11.1%	0.40 [0.24, 0.66]	2006	
Sun 2013	3	30	10	30	3.8%	0.30 [0.09, 0.98]	2013	·
Boelens 2014	4	61	8	62	4.1%	0.51 [0.16, 1.60]	2014	
Subtotal (95% CI)		329		329	34.4%	0.50 [0.37, 0.67]		•
Total events	47		90					1988
Heterogeneity: Tau* = 1	0.00; Chi*	= 5.66,	df = 6 (P	= 0.46)	$ ^{2} = 0\%$			
Test for overall effect: 2	Z = 4.49 (P	< 0.00	001)					
Total (95% CI)		2838		2845	100.0%	0.63 [0.49, 0.82]		•
Total events	442		536					0.000
Heterogeneity: Tau* = :	0.09; Chi ²	= 29.81	1, df = 12	(P = 0.1)	003); (*=	60%		0.01 0.1 1 10 100
Test for overall effect: 2								0.01 0.1 1 10 100 Favours EEN Favours EPN
Test for subgroup diffe	rences: C	hi*= 3.	92, df = 1	(P = 0.	05), P = 7	4.5%		rayouis EEN Favouis EFN

INITIATION OF ENTERAL NUTRITION (24 - 48 Hours)

RATIONALE

- Supports intestinal integrity.
- Stimulates intestinal blood flow.

- Induces release of trophic agents.
- Supports immunocytes.

KEY POINTS

 Presence of bowel sounds is NOT required.

 Do NOT wait for flatus or bowel movement.

EARLY VERSUS DELAYED ENTERAL NUTRITIONAL SUPPORT Mortality

	Early	EN	Delayed	None		Risk Ratio	Risk Ratio			
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	Year	M-H, Random, 95% C		
Sagar 1979	0	15	0	15	100000000000000000000000000000000000000	Not estimable	1979			
Moore 1986	1	32	2	31	2.3%	0.48 [0.05, 5.07]	1986	+ -		
Chiarelli 1990	0	10	0	10		Not estimable	1990			
Schroeder 1991	0	16	0	16		Not estimable	1991			
Eyer 1993	2	19	2	19	3.7%	1.00 [0.16, 6.38]	1993	-		
Beier-Holgersen 1996	2	30	4	30	4.9%	0.50 [0.10, 2.53]	1996	·		
Carr 1996	0	14	1	14	1.3%	0.33 [0.01, 7.55]	1996	• •		
Chuntrasakul 1996	1	21	3	17	2.7%	0.27 [0.03, 2.37]	1996	• • • • • • • • • • • • • • • • • • • 		
Watters 1997	0	14	0	14		Not estimable	1997			
Singh 1998	4	21	4	22	8.2%	1.05 [0.30, 3.66]	1998	-		
Kompan 1999	0	14	1	14	1.3%	0.33 [0.01, 7.55]	1999			
Minard 2000	1	12	4	15	3.0%	0.31 [0.04, 2.44]	2000			
Pupelis 2000	1	11	5	18	3.2%	0.33 [0.04, 2.45]	2000			
Pupelis 2001	1	30	7	30	3.1%	0.14 [0.02, 1.09]	2001	•		
Dvorak 2004	0	7	0	10		Not estimable	2004			
Kompan 2004	0	27	1	25	1.3%	0.31 [0.01, 7.26]	2004	• •		
Peck 2004	4	14	5	13	11.0%	0.74 [0.25, 2.18]	2004	-		
Malhotra 2004	12	100	16	100	26.5%	0.75 [0.37, 1.50]	2004			
Nguyen 2008	6	14	6	14	17.5%	1.00 [0.43, 2.35]	2008	-+-		
Moses 2009	3	29	3	30	5.6%	1.03 [0.23, 4.71]	2009	9		
Chourdakis 2012	3	34	2	25	4.4%	1.10 [0.20, 6.12]	2012			
Total (95% CI)		469		467	100.0%	0.70 [0.49, 1.00]		•		
Total events	41		66							
Heterogeneity: Tau ² = 0	.00; Chi2=	7.23,	f = 15 (P	= 0.95);	$l^2 = 0\%$					
Test for overall effect: Z								0.1 0.2 0.5 1 2 Favors Early EN Favors De		

EARLY VERSUS DELAYED ENTERAL NUTRITIONAL SUPPORT Infectious Complications

	Early	EN	Delayed/	None		Risk Ratio		Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	Year	M-H, Random, 95% CI
Sagar 1979	3	15	5	15	3.1%	0.60 [0.17, 2.07]	1979	
Moore 1986	3	32	9	31	3.3%	0.32 [0.10, 1.08]	1986	· · · · · · · · · · · · · · · · · · ·
Schroeder 1991	1	16	0	16	0.5%	3.00 [0.13, 68.57]	1991	
Carr 1996	0	14	3	14	0.6%	0.14 [0.01, 2.53]	1996	· · · · · · · · · · · · · · · · · · ·
Beier-Holgersen 1996	2	30	14	30	2.5%	0.14 [0.04, 0.57]	1996	
Singh 1998	7	21	12	22	7.6%	0.61 [0.30, 1.25]	1998	3 3 3 4
Minard 2000	6	12	7	15	6.6%	1.07 [0.49, 2.34]	2000	
Malhotra 2004	54	100	67	100	20.9%	0.81 [0.64, 1.01]	2004	· ·
Kompan 2004	9	27	16	25	9.4%	0.52 [0.28, 0.96]	2004	
Peck 2004	12	14	11	13	17.7%	1.01 [0.74, 1.39]	2004	· ·
Nguyen 2008	3	14	6	14	3.5%	0.50 [0.15, 1.61]	2008	· ·
Moses 2009	17	29	19	30	14.5%	0.93 [0.61, 1.39]	2009	i —
Chourdakis 2012	13	34	12	25	9.8%	0.80 [0.44, 1.44]	2012	(
Total (95% CI)		358		350	100.0%	0.74 [0.58, 0.93]		•
Total events	130		181					
Heterogeneity: Tau ² = 0	.05; Chi2=	19.58,	df = 12 (P	= 0.08)	F = 39%			14 13 15 14 15 15
Test for overall effect: Z	= 2.54 (P	= 0.01)			Ş.			0.1 0.2 0.5 1 2 5 10 Favors Early EN Favors Delayed/No

SMALL BOWEL VERSUS GASTRIC FEEDINGS

Acceptable to initiate enteral nutrition in the stomach.

ASPEN: Yes

ESPEN: Yes

- No difference, to include:
 - LOS
 - Mortality
 - Nutrient delivery
 - Pneumonia

SMALL BOWEL VERSUS GASTRIC FEEDINGS **Nutritional Efficiency**

ASPEN

Total Events Total Weight M-H, Random, 95% CI

3.9%

2.5%

7.2%

0.6%

0.9%

5.1%

4.7%

54 17.5%

89 12.4%

494 100.0%

43 10.4%

41 20.6%

19

23

15

11

35

51 14.2%

51

153

Risk Ratio

0.67 [0.22, 1.99]

0.65 [0.34, 1.22]

0.69 [0.46, 1.05]

1.46 [0.37, 5.78]

1.07 [0.49, 2.34]

0.16 [0.01, 3.03]

2.26 [0.22, 23.71]

0.82 [0.48, 1.39]

0.35 [0.14, 0.90]

1.97 [0.73, 5.28] 0.56 [0.35, 0.89]

0.93 [0.52, 1.65]

0.75 [0.60, 0.93]

Small Bowel

10

110

Heterogeneity: $Tau^2 = 0.02$; $Chi^2 = 12.33$, df = 11 (P = 0.34); $I^2 = 11\%$

37

Study or Subgroup

Montecalvo, 1992

Kortbeek, 1999

Taylor, 1999

Kearns, 2000

Minard, 2000

Davies, 2002

Montejo, 2002

Hsu, 2009

White, 2010

Davies, 2012

Total (95% CI)

Total events

Acosta-Escribano, 2010

Test for overall effect: Z = 2.56 (P = 0.01)

Day, 2001

Gastric

M-H, Random, 95% CI

Risk Ratio

0.5

Favors Small bowel Favors Gastric

ESPEN

	Post Pyloric F	eeding	Gastric Fe	eding		Risk Ratio		Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	Year	M-H, Random, 95% CI
Davies 2002	- (31	11	35	29.8%	0.41 [0.15, 1.16]	2002	+
Montejo 2002	1	50	25	51	16.8%	0.04 [0.01, 0.29]	2002	
Acosta-Escribano 2010	3	50	10	54	26.6%	0.32 [0.09, 1.11]	2010	-
Davies 2012	0	91	8	89	10.1%	0.06 [0.00, 0.98]	2012	
Wan 2015	1	35	14	35	16.7%	0.07 (0.01, 0.51)	2015	-
Total (95% CI)		257		264	100.0%	0.16 [0.06, 0.45]		•
Total events	1		68					n 1978 n 2
Heterogeneity: Tauf = 0.	65; Chi ² = 7.94,	d + 4	P = 0.09); f	. 50%				kan ah la tari
Test for overall effect: 2		1900 1011	MILLION (17)					0.001 0'.1 1 10 1000' Favours Post Pyloric Feed Favours Castric Feeding

SMALL BOWEL VERSUS GASTRIC FEEDINGS Pneumonia

	Small Bo	owel	Gastr	ic		Risk Ratio	Risk Ratio
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	M-H, Random, 95% CI
Montecalvo, 1992	4	19	6	19	3.9%	0.67 [0.22, 1.99]	
Kortbeek, 1999	10	37	18	43	10.4%	0.65 [0.34, 1.22]	
Taylor, 1999	18	41	26	41	20.6%	0.69 [0.46, 1.05]	
Kearns, 2000	4	21	3	23	2.5%	1.46 [0.37, 5.78]	
Minard, 2000	6	12	7	15	7.2%	1.07 [0.49, 2.34]	
Day, 2001	0	14	2	11	0.6%	0.16 [0.01, 3.03]	-
Davies, 2002	2	31	1	35	0.9%	2.26 [0.22, 23.71]	
Montejo, 2002	16	50	20	51	14.2%	0.82 [0.48, 1.39]	
Hsu, 2009	5	59	15	62	5.1%	0.35 [0.14, 0.90]	
White, 2010	11	57	5	51	4.7%	1.97 [0.73, 5.28]	
Acosta-Escribano, 2010	16	50	31	54	17.5%	0.56 [0.35, 0.89]	
Davies, 2012	18	91	19	89	12.4%	0.93 [0.52, 1.65]	
Total (95% CI)		482		494	100.0%	0.75 [0.60, 0.93]	•
Total events	110		153				
Heterogeneity: Tau ² = 0.02	2; Chi ² = 12	.33, df=	= 11 (P =	0.34); (²=11%		0.1 0.2 0.5 1 2 5 10
Test for overall effect: $Z = 2$	2.56 (P = 0.	.01)	(7.)				
Test for overall effect: Z = 2	2.56 (P = U.	.01)					Favors Small bowel Favors Gastric

IS ENTERAL NUTRITION SAFE WITH HEMODYNAMIC INSTABILITY? ASPEN

Do not start in the setting of hemodynamic compromise or instability.

- Hypotensive (MAP < 50mm Hg)
- Vasopresors are being initiated
- Escalating doses of vasopressors are required for hemodynamic instability

EN may be started if vasopressor support is being withdrawn, but watch for abdominal distention, increased residuals or worsening metabolic acidosis or rising base deficit.

IS ENTERAL NUTRITION SAFE WITH HEMODYNAMIC INSTABILITY? ESPEN

Uncontrolled shock

Intestinal ischemia

Uncontrolled hypoxemia and acidosis

Intestinal obstruction

Uncontrolled UGI hemorrhage

Abdominal compartment syndrome

Gastric aspirate > 500cc/6hr

High-output fistula without distal feeding access

GASTRIC RESIDUALS

- Gastric residuals should NOT be used as part of routine care.
- Residuals do NOT correlate with:
 - Pneumonia
 - Regurgitation
 - Aspiration
- If you must check, hold EN if gastric residual is > 500cc in the absence of other signs of intolerance.

ASPEN: Do not check residuals.

ESPEN: Suggest hold EN if > 500cc/6hrs.

DAILY VOLUME GOAL Nurse-Driven

 Use volume-based feeding protocols in which 24-hour daily volumes are targeted instead of simply hourly rates.

This allows the RN to increase feeding rates to make up for lost volume.

EMPOWER THE NURSES!

UMC PREOPERATIVE ENTERAL NUTRITION GUIDELINE

Continue enteral nutrition until one hour prior to surgery on <u>ventilated patients</u> except for the following procedures (hold for 8 hours):

- Tracheostomy
- Laparotomy
- Spine surgery
- Oral maxillofacial procedures
- If patient will be placed in prone position
- Thoracotomy, especially if patient is to be placed in lateral position or if there is need to change to a double-lumen tube.

TUBE PLACEMENT

- CXR is the gold standard
- Capnometry is an adjunct
- Auscultation is of <u>minimal</u> value (dangerous)



DIARRHEA

- Do NOT stop enteral feeds.
 - If necessary, reduce rate.
- Review medication list (most likely source), including IV antibiotics.
- Rule-out infectious etiology (C.diff).
- Significant reduction in diarrhea with continuous vs. bolus feeding.

	Continuo	us EN	Intermittent EN			Risk Ratio		Risk	Risk Ratio		
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Random, 95% CI	Year	M-H, Rande	om, 95% CI		
Steevens 2002	2	9	5	9	33.3%	0.40 [0.10, 1.55]	2002				
Serpa 2006	Ö	14	3	14	7.4%	0.14 [0.01, 2.53]	2006 -				
Kadami 2014	2	15	5	15	28.1%	0.40 [0.09, 1.75]	2014	_	_		
MacLeod 2017	3	81	5	79	31.3%	0.59 [0.14, 2.37]	2017	-			
Total (95% CI)		119		117	100.0%	0.42 [0.19, 0.91]		-			
Total events	7		18						50		
Heterogeneity: Tau ² =	0.00; Chi ²	= 0.77.	df = 3 (P)	0.86);	$t^2 = 0\%$		24		- 1	110	
Test for overall effect:	Z = 2.19 G	P = 0.03	3)				0.0	하는 그 없었다면 되고 하면 하는데 하다면 되고 있다.	Favours intermittent EN	100	

Help! My tube is clogged!

- Prevention is the best key
- Irrigate with ~25cc of water every 4-6 hrs and before & after Rxs.
- If clogged, do this:
 - Push back and forth with a 60cc syringe containing warm water.
 - If that does not work, let the warm water sit for ~20 minutes
- Sorry! Coke, Pepsi, Gingerale or meat tenderizer really do not work even though many think otherwise. Carbonation may make matters worse.
- Pancrease (Not enteric coated)
 - Viokase

TARGETED GLUCOSE RANGE

SCCM: 150-180 mg/dl

ASPEN: 140-180 mg/dl

ESPEN: 150-180 mg/dl

Glucose > 200 is NOT OK! Consider insulin drip

PERCUTANEOUS GASTROSTOMY PEARLS

Literature supports use within 4 hours. No joke!

ASPEN: Yes

ESPEN: Yes

Restart EN at same rate prior to stopping for procedure.

If concern for patient pulling PEG out, place an abdominal binder.

Don't place bumper too tight to the skin.

OPEN GASTROSTOMY, G-J & J TUBES

- Moss tube
- MIC tube
- Foley
- Feeding jejunostomy tube

Pearls

- Always use water or saline for the balloon;
 NEVER use air.
- Can't check residuals via a J-tube.
- If the tube fall out, immediately replace with a Foley catheter before tract closes.

