

Beslenme Problemleri Açısından Farkındalık, Tarama ve Takip Nasıl Yapılmalıdır?

Dr.Tülay Kuş

Prevalence of cancer-related MN Cancer out- and inpatients

Cancer outpatients (1'000 pts, NRS-2002) 33.8 %

Bozzetti et al. Support Care Cancer 2009;17:279

Cancer inpatients (71 pts, PG-SGA)76 %

Bauer et al. Eur J Clin Nutr 2002;56:779

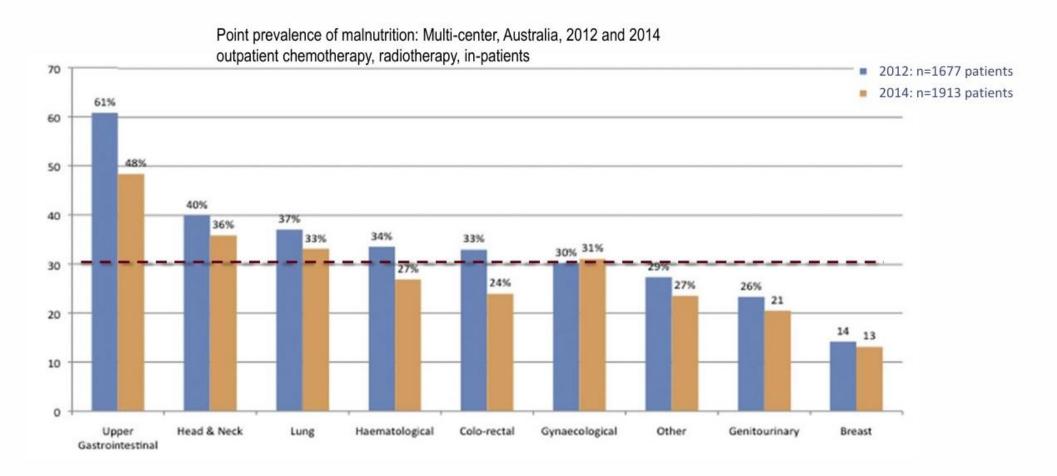
Colorectal cancer (inpatients, 234 pts, PG-SGA)
 41 %

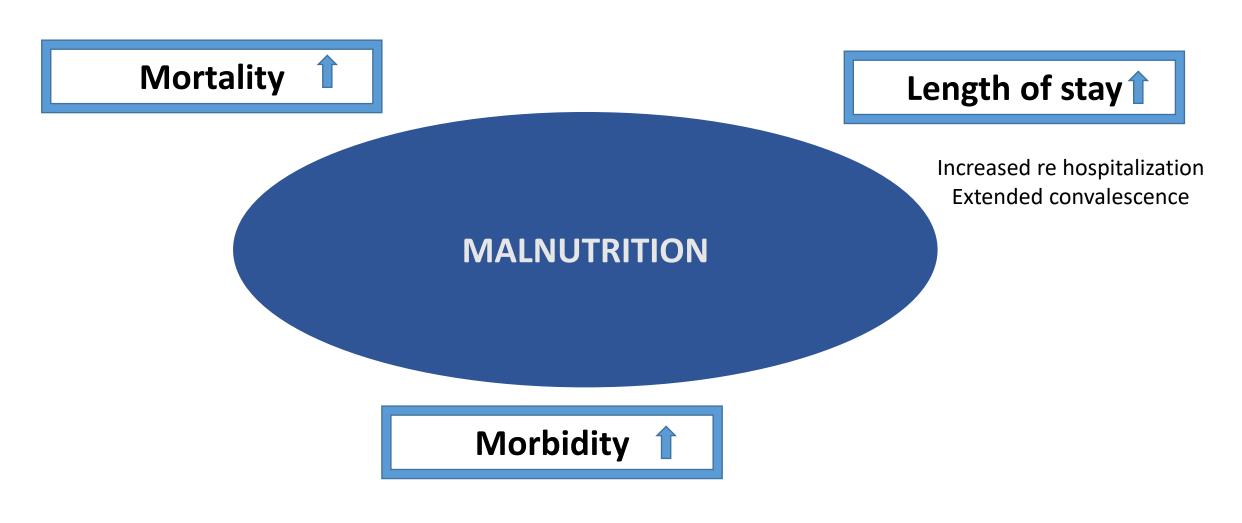
Gubta et al. Eur J Clin Nutr 2004;59:35

Ovarian cancer (inpatients, 132 pts, SGA) 50 %

Gubta et al. J Ovar Res 2008;1:5

MALNUTRITION: INCIDENCE AND DIAGNOSIS





Increased infection; deteriorated wound healing; diminished tolerance to anti-cancer therapy; organ dysfunction; increased complications

Cancer: disease and nutrition are key determinants of patients' quality of life

Paula Ravasco ¹, Isabel Monteiro-Grillo, Pedro Marques Vidal, Maria E Camilo

Party of

Variable	Global	function scores		Global symptom scores ^a			Global single item scores ^a		
	F-test	Estimates of effect size (%) ^b	P value	F-test	Estimates of effect size (%) ^b	P value	F-test	Estimates of effect size (%) ^b	P value
Stage	1.6	1	0.18	56.5	22	0.001	103.7	30	0.0001
Location	111.2	30	0.0001	77.2	41	0.0001	49.2	20	0.001
Energy intake	27.2	10	0.01	1.0	3	0.35	3.9	4	0.07
Protein intake	27.2	10	0.01	1.0	4	0.25	4.2	5	0.07
Weight loss	133.7	30	0.0001	0.05	1	0.82	1.2	3	0.10
Duration of disease	1.5	3	0.14	10.0	7	0.06	1.2	3	0.30
Chemotherapy	35.3	10	0.001	2.1	4	0.22	1.3	1	0.25
Surgery	6.1	6	0.01	1.4	1	0.86	3.0	4	0.09

^a Due to the potential association between symptoms and diagnoses, associations were adjusted for cancer location

^b The sum of percentages may not equal 100% due to the corrected error size

Causes of Cancer-related Malnutrition

 Deterioration in taste, smell and appetite, as a consequence of the disease and/or therapy Altered food preferences/avoidance/aversion Anorexia Dysphagia, odynophagia Partial/total gastrointestinal obstruction or dysfunction Early satiety, nausea and vomiting Soreness, xerostomia, sticky saliva, painful throat, trismus Oral lesions and oesophagitis Radiotherapy-/chemotherapy-induced mucositis Acute or chronic radiation enteritis during and after radiotherapy Depression, anxiety Pain

High Risk Populationf for Cancer-related Malnutrition

*The topmost cancers associated with weight loss and malnutrition are pancreatic, hepatic, gastric, oesophageal, head and neck and lung cancer; in the case of incurable cancers, all patients are at elevated risk of malnutrition

**Future risks of weight loss; Aggressive treatment, for example, radiotherapy with concurrent chemotherapy, is often associated with acute weight loss of >10%. For radiotherapy, the site of treatment may have important nutritional consequences, for example radiation to the oral cavity, laryngeal, pharyngeal and oesophageal regions: the resulting pain and mucositis impair dietary intake

The early

assessment of nutritional status, including body composition when feasible, is now recommended by international guidelines (European Society for Clinical Nutrition and Metabolism [ESPEN] and European Society for Medical Oncology [ESMO]) on the management of patients with cancer.

Nutritional screening needs to be *simple *rapid *easily performed

on hospital admission or at each oncological visit.



OPEN ACCESS | GUIDELINE SUMMARY | © (†) | January 14, 2021



Nutrition in Cancer Care: A Brief, Practical Guide With a Focus on Clinical Practice

Although there is no evidence from randomized clinical trials showing the benefit of *regular nutritional screening* in heterogenous cancer populations, there are some high-risk cancer sites (head and neck, upper GI) where close monitoring (eg, weekly) of the patient's nutritional status is essential.

Other cancer sites where the risk of developing weight loss is lower should be screened on a more individual basis according to the clinical situation, for example, with clinical deterioration because of disease progression and/or toxicities.

From a more practical perspective, every patient with cancer should be screened, at least, at diagnosis of cancer, on hospital admission, on clinical deterioration, and when reporting weight loss while receiving systemic treatment, radiotherapy, or surgery.



Contents lists available at ScienceDirect

Clinical Nutrition

journal homepage: http://www.elsevier.com/locate/clnu

ESPEN Guideline

ESPEN practical guideline: Clinical Nutrition in cancer

Maurizio Muscaritoli a, *, Jann Arends b, Patrick Bachmann c, Vickie Baracos d,

To detect nutritional disturbances at an early stage, we recommend to regularly evaluate <u>nutritional intake</u>, <u>weight</u> <u>change</u>, <u>and body mass index (BMI)</u>, beginning with cancer diagnosis and repeated depending on the stability of the clinical situation.

(Recommendation B1-1; strength of recommendation strong e level of evidence very low e strong consensus)



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Clinical Nutrition

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ESPEN Guideline

ESPEN practical guideline: Clinical Nutrition in cancer

Maurizio Muscaritoli ^{a, *}, Jann Arends ^b, Patrick Bachmann ^c, Vickie Baracos ^d,

Nutritional Assessment

In patients with abnormal screening, we recommend objective and quantitative assessment of nutritional intake, nutrition impact symptoms, muscle mass, physical performance and the degree of systemic inflammation. (Recommendation B1-2; strength of recommendation strong e level of evidence very low e consensus)

Malnutrition screening tools

The severity of food intake impairment may be assessed with validated clinical tools:

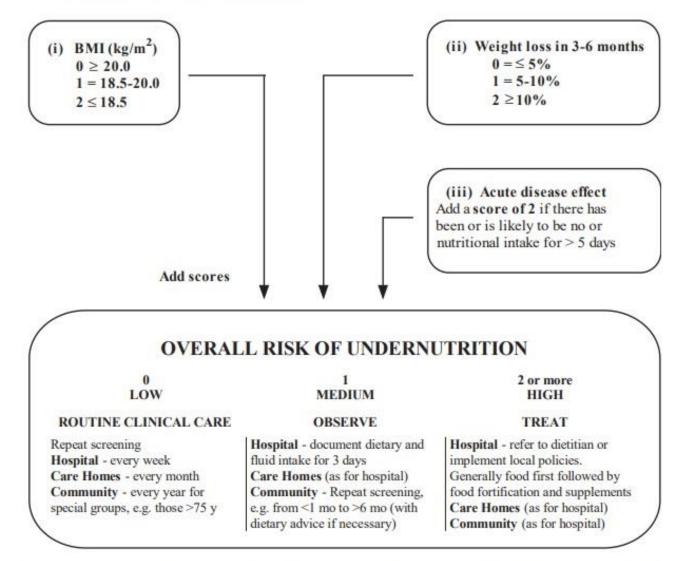
1.MUST

2. NRS 2002

3. Mini Nutritional Assessment (MNA)

4. The Patient-Generated Subjective Global Assessment (PG-SGA)

Malnutrition Universal Screening Tool (MUST) for adults



Can be adapted for special circumstances (e.g. when weight and height cannot be measured or when there are fluid disturbances) using specified alternative measurements including subjective criteria. It also identifies obesity $(BMI > 30 \text{ kg/m}^2)$.

Nutritional Risk Screening (NRS 2002)

Table 1 Initial screening						
1	Is BMI < 20.5?	Yes	No			
2	Has the patient lost weight within the last 3 months?					
3	Has the patient had a reduced dietary intake in the last week?					
4	Is the patient severely ill ? (e.g. in intensive therapy)					

Yes: If the answer is 'Yes' to any question, the screening in Table 2 is performed.

No: If the answer is 'No' to all questions, the patient is re-screened at weekly intervals. If the patient e.g. is scheduled for a major operation, a preventive nutritional care plan is considered to avoid the associated risk status.



	Impaired nutritional status	Severity of d	isease (≈ increase in requirements)
Absent Score 0	Normal nutritional status	Absent Score 0	Normal nutritional requirements
Mild Score 1	Wt loss >5% in 3 mths or Food intake below 50-75% of normal requirement in preceding week	Mild Score 1	Hip fracture* Chronic patients, in particular with acute complications: cirrhosis*, COPD*. Chronic hemodialysis, diabetes, oncology
Moderate Score 2	Wt loss > 5% in 2 mths or BMI 18.5 – 20.5 + impaired general condition or Food intake 25–60% of normal requirement in preceding week	Moderate Score 2	Major abdominal surgery* Stroke* Severe pneumonia, hematologic malignancy
Severe Score 3	Wt loss >5% in 1 mth (>15% in 3 mths) or BMI <18.5 + impaired general condition or Food intake 0-25% of normal requirement in preceding week in preceding week.	Severe Score 3	Head injury* Bone marrow transplantation* <i>Intensive care patients (APACHE>10)</i> .
Score:	+	Score:	= Total score

Score ≥3: the patient is nutritionally at-risk and a nutritional care plan is initiated

Score <3: weekly rescreening of the patient. If the patient e.g. is scheduled for a major operation, a preventive nutritional care plan is considered to avoid the associated risk status.

The Journal of Nutrition, Health & Aging© Volume 13, Number 9, 2009

VALIDATION OF THE MINI NUTRITIONAL ASSESSMENT® SHORT-FORM (MNA-SF)

VALIDATION OF THE MINI NUTRITIONAL ASSESSMENT SHORT-FORM (MNA®-SF): A PRACTICAL TOOL FOR IDENTIFICATION OF NUTRITIONAL STATUS

M.J. KAISER¹, J.M. BAUER¹, C. RAMSCH², W. UTER², Y. GUIGOZ³, T. CEDERHOLM⁴, D.R. THOMAS⁵,

S	creening		
A	Has food intake swallowing diffi	declined over the past 3 months due to loss of appetite, digestive problems, chewing or iculties?	
	1 = moderate	ecrease in food intake e decrease in food intake ase in food intake	
В	Weight loss dur	ing the last 3 months	
	1 = does not	oss between 1 and 3 kg (2.2 and 6.6 lbs)	
c	Mobility		
	0 = bed or ch	pair bound	
		et out of bed / chair but does not go out	
D	Has suffered psy	ychological stress or acute disease in the past 3 months?	
	0 = yes	2 = no	
E	Neuropsycholog	gical problems	
	0 = severe de	ementia or depression	
	1 = mild den		
	2 = no psych	ological problems	
F	Body Mass Inde	x (BMI) (weight in kg) / (height in m²)	
	0 = BMI less t	than 19	
		o less than 21	
	2 = BMI 21 to	o less than 23	
	3 = BMI 23 o	r greater	

F2 Calf circumfere	DO NOT ANSWER QUESTION F2 IF QUESTION F1 IS ALREAD	
0 = CC less to 3 = CC 31 or	nan 31	
Screening score (max. 14 points)	2	00
12-14 points:	Normal nutritional status	
8-11 points:	At risk of malnutrition	
0-7 points:	Malnourished	



Scored Patient-Generated Subjective Global

Assessment (PG-SGA)	
History: Boxes 1 - 4 are designed to be completed by the patient. [Boxes 1-4 are referred to as the PG-SGA Short Form (SF)]	
In summary of my current and recent weight: I currently weigh aboutkg I am aboutkg One month ago I weighed aboutkg Six months ago I weighed aboutkg During the past two weeks my weight has: decreased (1) not changed (0) increased (0) Box 1	2. Food intake: As compared to my normal intake, I would rate my food intake during the past month as unchanged (0) more than usual (0) less than usual (1) I am now taking normal food but less than normal amount (1) little solid food (2) only liquids (3) only nutritional supplements (3) very little of anything (4) only tube feedings or only nutrition by vein (0) Box 2
3. Symptoms: I have had the following problems that have kept me from eating enough during the past two weeks (check all that apply) no problems eating (0) no appetite, just did not feel like eating (3) vomiting (3) nausea (1) diarrhea (3) constipation (1) dry mouth (1) mouth sores (2) smells bother me (1) things taste funny or have no taste (1) feel full quickly (1) problems swallowing (2) fatigue (1) pain; where? (3) other (1)** **Examples: depression, money, or dental problems Box 3	4. Activities and Function: Over the past month, I would generally rate my activity as: normal with no limitations (0) not my normal self, but able to be up and about with fairly normal activities (1) not feeling up to most things, but in bed or chair less than half the day (2) able to do little activity and spend most of the day in bed or chair (3) pretty much bed ridden, rarely out of bed (3) Box 4

Patient Identification Information

The remainder of this form is to be completed by your doctor, nurse, dietitian, or therapist. Thank you.

Additive Score of Boxes 1-4

Scored Patient-Generated Subjective Global Assessment (PG-SGA)

Worksheet 1 – Scorin	Additive Score of Boxes 1-4 (See Side 1)								
To determine score, use 1-mon 1-month weight data. Use point has lost weight during the past 2	s below to score weigh	t change and add	d one extra point if patient	5. Worksheet 2 – Disease and its relation to nutritional requirements: Score is derived by adding 1 point for each of the following conditions: Cancer Presence of decubitus, open wound or fistula					
Weight loss in 1 month	Points		s in 6 months				<u> </u>	•	
10% or greater	4		or greater	☐ AIDS			Presen	ce of trauma	
5-9.9%	3		0- 19.9%	☐ Pulmonary	or cardia	c cachexia	Age gr	eater than 65	
3-4.9% 2-2.9%	2		6- 9.9% 2- 5.9%	20-20-20-20-20-20-20-20-20-20-20-20-20-2					
0-1.9%	0		0- 1.9%	Chronic re		•			
0 1.270	· ·	2	1.770	Other relevant	_				
	Numerical sco	ore from V	Vorksheet 1	Primary diseas	e staging	(circle if kr	own or appro Nu	priate) I II III IV Oth Imerical score from	m Worksheet 2 B
6. Worksheet 3 – Met Score for metabolic stress is patient who has a fever of 3 Stress none (s determined by a nu 8.8 °C (3 points) for	umber of varia	oint) and who is on 10 m		cally (2 po				The score is additive so that a 5 points.
Fever no fever	•	> 37.2 and		and < 38.8		> 38.8 °C			
Fever duration no fever		< 72 hours	72 hou			> 72 hours			
	costeroids	low dose		ate dose		high dose			
		(< 10 mg pr		and < 30 mg		(≥ 30 mg pr		umerical score from	m Worksheet 3 C
		equivalents	/day) predni	sone equivalents/day)		equivalents/	day)	umericai score iroi	iii worksneet 3 C
7. Worksheet 4 – Physe Exam includes a subjective ev Definition of categories: 0 = n Muscle Status temples (temporalis muscle) clavicles (pectoralis & deltoids shoulders (deltoids) interosseous muscles scapula (latissimus dorsi, trape	aluation of 3 aspects of a abnormality, 1+ = m 0	iild, 2+ = moders + 2+ 3+ + 2+ 3+ + 2+ 3+ + 2+ 3+	ate, 3+ = severe. Rating in the Fat St orbital triceps fat over Glo	nese categories is <i>not</i> addit tores fat pads 0 skin fold 0 rlying lower ribs 0		3+ Poi tota 3+ 3+	Ily assess the deg nt score for the phys I body deficit. No M	ree of deficit (or presence of e sical exam is determined by the o	
thigh (quadriceps) calf (gastrocnemius) Global muscle status rati	0 1+	+ 2+ 3+ + 2+ 3+ + 2+ 3+	ankle e sacral e ascites	edema 0	1+ 2+ 1+ 2+ 1+ 2+	3+		Numerical Score f	or Worksheet 4 D
Giovai muscie status fati	s U 17	21 31		bal fluid status rating 0		3+ To:	tal PG-SG	A Score (Total numerica	al score of A+B+C+D)
Clinician Signature			_RD RN PA MD DO Otl					egory Rating (Stage A, Stag	
Worksheet 5 – PG-SGA Stage A Well-nourished No weight loss OR recent non-fluid v Nutrient intake No deficit OR Significar recent improvement allowing adequate intake No deficit OR Significar recent improvement Physical Exam No deficit OR chronic deficit to With recent	Stage B Moderate/suspected t 5% loss in 1 month (t gain OR Progressive weig ant Definite decrease in in Presence of NIS (Box	malnutrition ≤10% in 6 months) tht loss take 3 of PG-SGA) efficit tion oderate loss	ies Stage C Severety malnourished > 5% loss in 1 month (>10% in 6 m OR Progressive weight loss Severe deficit in intake Presence of NIS (Box 3 of PG-SGA Severe functional deficit OR Recent significant deterioratic Obvious signs of malnutrition (e.g., severe loss muscle, fat,	patient & family nutritional supp First line nutrit Triage based on 0-1 No inter 2-3 Patient & indicate: 4-8 Require:	education, sy ements, enter ion interventa in PG-SGA po- vention requi- ic family educa- l by symptom intervention	ymptom manage ral, or parenteral ion includes opto bint score red at this time. ration by dietitian a survey (Box 3) by dietitian, in c	ment including phar triage). imal symptom mand Re-assessment on ro, nurse, or other clianal lab values as aponjunction with nur- symptom management	macologic intervention, and appro- ingement. Dutine and regular basis during tre- nician with pharmacologic interve- propriate. See or physician as indicated by sy- ent and/or nutrient intervention op ©FD Ottery 2005, 2	mptoms (Box 3). stions. 2006, 2015, 2020 v.4.3.20
clinical improvement	palpation &/or loss of		possible edema)				email: faithe	otterymdphd@gmail.c	com or info@pt-global.org

possible edema)

According to the results of this meta-analysis, PG-SGA showed the best diagnostic performance among the three modalities with the sensitivity of 0.964, specificity of 0.905.

X. Ruan, R. Nakyeyune, Y. Shao et al.

Clinical Nutrition 40 (2021) 1733-1743

Table 2Quality of the body of evidence for each outcome of interest reflecting the GRADE) approach.

Outcome	Nº of studies (Nº of patients)	Study design	Risk of bias	Indirectness	Inconsistency	Imprecision	Publication bias	Quality of evidence (GRADE
Construct validity of the MNA	8 studies 1233 patients	cross- sectional studies	serious ^a	serious ^b	serious ^c	not serious	strongly suspected ^d	⊕○○○ VERY LOW
Construct validity of the NRS- 2002	28 studies 5160 patients	cross- sectional studies	serious ^a	serious ^b	serious ^c	not serious	strongly suspected ^d	⊕○○○ VERY LOW
Construct validity of the PG-SGA	8 studies 1265 patients	cross- sectional studies	serious ^a	serious ^b	not serious	not serious	strongly suspected ^d	⊕⊕⊕○ MODERATE

^a Found to have a risk of bias when evaluated using the QUADAS-2.

Table 3Result of meta-analysis and Bayes analysis.

9	MNA	NRS-2002	PG-SGA
Sensitivity (95% CI)	0.910 (0.763, 0.970)	0.747 (0.680, 0.804)	0.964 (0.913, 0.986)
Specificity (95% CI)	0.720 (0.623, 0.800)	0.854 (0.808, 0.891)	0.905 (0.807, 0.956)
DOR (95% CI)	26.039 (9.732, 69.671)	17.361 (12.922, 23.325)	257.204 (62.758, 1054.108)
LR+ (95% CI)	3.245 (2.464, 4.275)	5.133 (4.029, 6.539)	10.173 (4.775, 21.676)
LR ⁻ (95% CI)	0.124 (0.047, 0.333)	0.296 (0.237, 0.369)	0.039 (0.015, 0.101)
1/LR- (95% CI)	8.023 (3.000, 21.461)	3.382 (2.713, 4.218)	25.283 (9.874, 64.739)

Abbreviations: MNA mini nutritional assessment, NRS-2002 nutritional risk screening 2002, PG-SGA patient generated subjective global assessment, SGA subjective global assessment; DOR diagnostic odds ratio, LR likelihood ratio, CI confidence interval.

^b The indirectness arises from the differences in populations, reference tests, cut-off values, how and when measurements were taken, as well as training and expertise of the individuals performing and interpreting the tests.

^c The results of the included studies varied widely.

d Publication bias was assessed by Stata.

Scored Patient-Generated Subjective Global Assessment (PG-SGA)

Worksheet 1 – Scoring Weight Loss To determine score, use 1-month weight data if available. Use 6-month data only if there is no		S 60 14 17500 51	Additive Score of Boxes 1-4 (See Side 1)				
1-month weight data. Use pthas lost weight during the pa Weight loss in 1 month 10% or greater 5-9.9% 3-4.9% 2-2.9% 0-1.9% 6. Worksheet 3 – M	Points 4 3 2 1 0 Numerical s	Weight loss 20% score from V	d one extra point if patient 1 of PG-SGA. s in 6 months or greater 0-19.9% 6-9.9% 2-5.9% 0-1.9% Vorksheet 1	5. Worksheet 2 – Disease and its relation to nutritional requirements: Score is derived by adding 1 point for each of the following conditions: Cancer Presence of decubitus, open wound or fistula AIDS Presence of trauma Pulmonary or cardiac cachexia Age greater than 65 Chronic renal insufficiency Other relevant diagnoses (specify) Primary disease staging (circle if known or appropriate) I II III IV Other Numerical score from Worksheet 2 B			
	of 38.8 °C (3 points)			of prednisone chronically (2 points) would have an additive score for this section ts.			
Fever no fer Fever duration no fer Fever duration no fer no fer no control of the fever duration no control of the fever duration of categories: 0 Muscle Status temples (temporalis muscle clavicles (pectoralis & delta shoulders (deltoids) interosseous muscles scapula (latissimus dorsi, transplanticular (gastrocnemius) Global muscle status response del fector duration of the fever duration of t	hy c ev = r SO	creening creening staff a	g tool for or tool is limind the lengt	ecific nutritional assessment and incological patients, but its use as a more than fat deficit/loss. Thuid). The property of the need for specially trained the of time needed to carry out the pated in approximately 15 minutes Torksheet 3 C Torksheet 4 C Torksheet 4 D Torksheet 4 D Torksheet 4 D Torksheet 4 D			
Clinician Signature			RD RN PA MD DO Othe	Date Date Global PG-SGA Category Rating (Stage A, Stage B or Stage C)			
Worksheet 5 – PG-SC Stage A Category Well-nourished No weight loss OR recent non-flt Nutrient intake No deficit OR Sigr recent improvement allov adequate intake Functioning Physical Exam No deficit OR Sigr recent improvement No deficit OR Sigr recent improvement Physical improvement No deficit OR chre deficit but with rec clinical improvemen	Stage B Moderate/suspec Sy loss in 1 m of Sy loss in 1 m of Sy loss in 1 m of Progressive inficant at Presence of NIS (cecent wing inficant of Recent deter onic Evidence of miss (of muscle mass (of muscle	ted malnutrition onth (<10% in 6 months) weight loss in intake Box 3 of PG-SGA) and deficit ioration to moderate loss k/or muscle tone on	Stage C Severely malnourished > 5% loss in 1 month (>10% in 6 mon OR Progressive weight loss Severe deficit in intake Presence of NIS (Box 3 of PG-SGA) Severe functional deficit OR Recent significant deterioration Obvious signs of malnutrition (e.g., severe loss muscle, fat, possible edema)	Nutritional Triage Recommendations: Additive score is used to define specific nutritional interventions including patient & family education, symptom management including pharmacologic intervention, and appropriate nutrient intervention (food, nutritional supplements, enteral, or parenteral triage). First line nutrition intervention includes optimal symptom management. Triage based on PG-SGA point score 0-1 No intervention required at this time. Re-assessment on routine and regular basis during treatment. 2-3 Patient & family education by dietitian, nurse, or other clinician with pharmacologic intervention as indicated by symptom survey (Box 3) and lab values as appropriate. 4-8 Requires intervention by dietitian, in conjunction with nurse or physician as indicated by symptoms (Box 3). 2 9 Indicates a critical need for improved symptom management and/or nutrient intervention options. ©FD Ottery 2005, 2006, 2015, 2020 v.4.3.20 email: faithotterymdphd@gmail.com or info@pt-global.org			

NUTRISCORE

A. Have you lost weight involuntarily in the last 3 months?

NoI am not sure

If yes, how much weight (in kilograms) have you lost?

•	1-5	1
•	6-10	2
•	11-15	3
•	>15	4
•	Unsure	2

B. Have you been eating poorly in the last week because of a decreased appetite?

NoYes

Location / Neoplasm	Nutritional risk	Score
Head and neck	High*	+ 2
Upper GI tract: oesophagus, gastric, pancreas, intestines		
Lymphoma that compromised GI tract		

Lung Abdominal and pelvis: liver, biliary tract, renal, ovaries, endometrial	Medium	+ 1
Breast Central Nervous System Bladder, prostate Colorectal Leukaemia, other lymphomas Others	Low	+ 0
Treatment	YES (+2)	NO (+0)
The patient is receiving concomitant chemo radiotherapy		
The patient is receiving hyper fractionated radiation therapy		
Haematopoietic stem cell transplantation		
	YES (+1)	NO (+0)
The patient is receiving chemotherapy		
The patient is only receiving radiotherapy		
	YES (+0)	NO (+0)
Other treatments or only symptomatic treatment		

^{*}Please repeat the screening every week for those patients at high risk

Total Score

Score ≥ 5: the patient is at nutritional risk. Please refer to a dietician.

A new nutritional screening tool for oncological outpatients to detect nutritional risk.

Comparative Study > Nutrition. 2017 Jan:33:297-303. doi: 10.1016/j.nut.2016.07.015.

Epub 2016 Aug 13.

NUTRISCORE: A new nutritional screening tool for oncological outpatients

Lorena Arribas ¹, Laura Hurtós ², Maria José Sendrós ³, Inmaculada Peiró ², Neus Salleras ⁴, Eduard Fort ², Jose Manuel Sánchez-Migallón ³

Affiliations + expand

PMID: 27751743 DOI: 10.1016/j.nut.2016.07.015

Abstract

Objectives: The aim of this study was to design a new nutritional screening tool (NUTRISCORE) to detect nutritional risk in outpatients with cancer.

Methods: A multicenter, cross-sectional study was conducted. We randomly selected outpatients receiving onco-specific, palliative, or symptomatic treatment for malignant neoplasms (including solid tumors and hematologic malignancies). These patients were assessed using the NUTRISCORE tool, the Malnutrition Screening Tool (MST), and the Patient-Generated Subjective Global Assessment (PG-SGA) to detect risk for malnutrition. The new tool included questions regarding the cancer site and active

Using the PG-SGA as a reference method, NUTRISCORE had 97.3% sensitivity and 95.9% specificity.

The so-called Global Leadership Initiative on Malnutrition (GLIM) criteria recommend that patients at nutritional risk, based on a validated screening tool, are assessed for the presence of aetiological and phenotypic criteria (Table 2).

Table 2 GLIM Criteria.

Adapted from: Cederholm T, Jensen GL, Correia MITD, et al; GLIM Core Leadership Committee; GLIM Working Group. GLIM criteria for the diagnosis of malnutrition – a consensus report from the global clinical nutrition community. Clin Nutr 2019; 38:1–9.

Phenotypic criteria	Non-volitional weight loss ¹	
	Low BMI ²	
	Reduced muscle mass ³	
Aetiological criteria	Reduced food intake or assimilation⁴	
	Disease burden/inflammatory condition ⁵	

>5% within the past 6 months, or >10% beyond 6 months.

5 Acute disease/injury or chronic disease-related (C-reactive protein may be used as a supportive laboratory measure).
Abbreviations: BIA, bioimpedance analysis; BMI, body mass index; CT, computed tomography; DEXA, dual-energy X-ray absorptiometry; ER, energy requirement; GI, gastrointestinal; GLIM, Global Leadership Initiative on Malnutrition; MRI, magnetic resonance imaging.

^{2 &}lt; 20 if < 70 years, or < 22 if > 70 years; Asia: < 18.5 if < 70 years, or < 20 if > 70 years.

³ Reduced by validated body composition measuring techniques (i.e. DEXA, BIA, CT, MRI; when not available, physical examination or standard anthropometric measures such as mid-arm muscle or calf circumferences may be used).

<sup>⁴ ≤50% of ERs >1 week, or any reduction for >2 weeks, or any chronic GI condition that adversely impacts food assimilation
or absorption.</sup>

Table 2 GLIM Criteria.

Adapted from: Cederholm T, Jensen GL, Correia MITD, et al; GLIM Core Leadership Committee; GLIM Working Group. GLIM criteria for the diagnosis of malnutrition – a consensus report from the global clinical nutrition community. Clin Nutr 2019; 38:1–9.

Phenotypic criteria	Non-volitional weight loss1	
	Low BMI ²	
	Reduced muscle mass ³	
Aetiological criteria	Reduced food intake or assimilation⁴	
	Disease burden/inflammatory condition ⁵	

Patients with at least one aetiological and one phenotypic criterion can be diagnosed with malnutrition

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Applied nutritional investigation

Nutritional risk and malnutrition rates at diagnosis of cancer in patients treated in outpatient settings: Early intervention protocol



Elena Álvaro Sanz, Ph.D. a, Marga Garrido Siles, Pharm.D., Ph.D. a*, Laura Rey Fernández, Dietitian a,

Comparison for nutritional risk

	NO nutritional risk, % (n)	Nutritional risk, % (n)	P-value
Age (y)			
<70	78.4 (174)	21.6 (48)	0.976
≥70	79.5 (58)	20.5(15)	
Sex			
Male	73 (100)	27 (37)	0.039
Female	83.5 (132)	16.5 (26)	
Grouped tumor location			
Upper	37.8 (17)	62.2 (28)	< 0.001
gastrointestinal/Head and			
Neck	86 (215)	14 (35)	
All others			
Treatment intention			
Curative/Radical	86.6 (149)	13.4(23)	< 0.001
Palliative	67.5 (83)	32.5 (40)	
Weight loss at diagnosis, % (median)	0 ± 5.4	13.5 ± 7.1	< 0.001
GPS			
0	87.8 (115)	12.2 (16)	
1	74.2 (89)	25.8 (31)	< 0.001
2	26.7 (4)	73.3 (11)	
INI risk	38.9 (208)	65.5 (58)	< 0.001
Cachexia			
Presence of cachexia	50.8 (60)	49.2 (58)	< 0.001
Absence of cachexia	97.2 (172)	2.8 (5)	

GPS, Glasgow Prognostic Score; INI, Inflammatory-Nutritional Index

Glasgow Prognostic Score (GPS) for Cancer Outcomes

Provides cancer prognosis based on serum biomarkers.

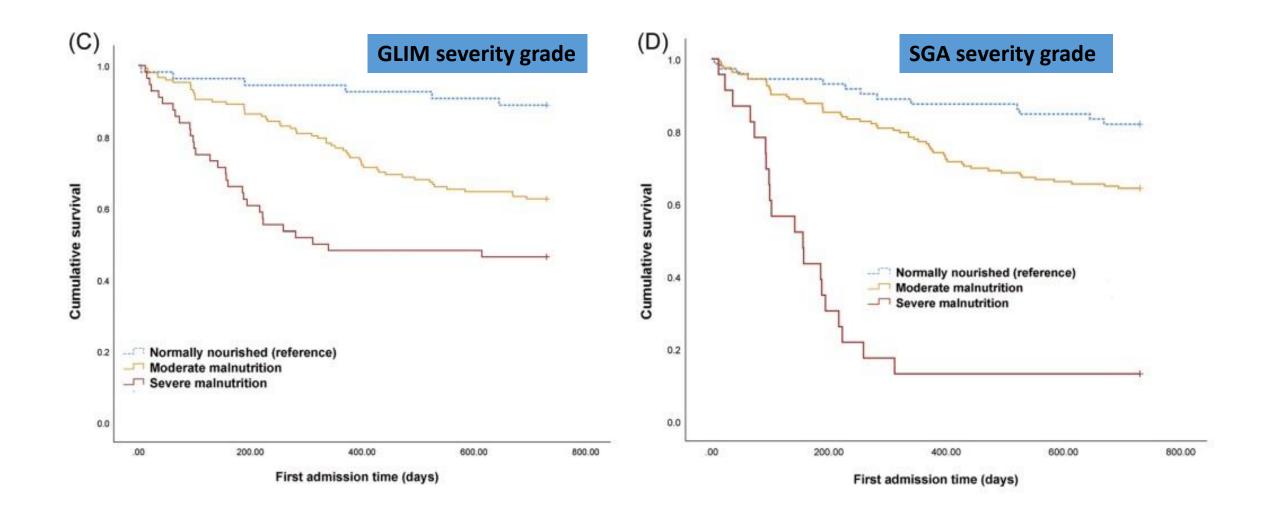
When to Use 🗸	Pearls/Pitfalls 🗸	Why Use 🗸	
	CRP ≤10 mg/L	CRP ≤10 mg/L	
	CRP >10 mg/L		
nin	Albumin <3.5 g/dL	(35 g/L)	
	Albumin ≥3.5 g/dL	(35 a/L)	

Patients were considered at risk when the Nutriscore was ≥5 points. The PG-SGA was used to evaluate patients' nutritional status

TABLE 3 The univariable and multivariable Cox regression analysis of malnutrition defined by the GLIM criteria and the SGA.

	2-year incidence of unplanned hospital admission				
	Univariable analysis	Univariable analysis		Multivariable analysis	
Nutrition assessment	HR (95% CI)	P value	HR (95% CI)	P value	
Malnutrition according to the GLIM criter	ia <u> </u>				
No (normally nourished)	Reference	877	Reference	-	
Yes (malnutrition)	4.61 (2.01-10.55)	<0.001	2.85 (1.22-6.68)	0.016	
Stage 1 (moderate malnutrition)	3.85 (1.66-8.95)	0.002	2.43 (1.02-5.77)	0.045	
Stage 2 (severe malnutrition)	7.26 (3.02-17.47)	<0.001	4.32 (1.75-10.66)	0.002	
Malnutrition according to the SGA					
No (normally nourished)	Reference	=	Reference	i o	
Yes (malnutrition)	2.76 (1.53-4.96)	0.001	2.07 (1.13-3.79)	0.019	
SGA-B	2.20 (1.21-4.02)	0.010	1.63 (0.88-3.04)	0.122	
SGA-C	12.48 (6.11-25.51)	<0.001	8.39 (3.98-17.71)	<0.001	

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Kaplan-Meier curves for 2-year unplanned hospital admission stratified by the GLIM severity grade and SGA severity grade.

GLIM, Global Leadership Initiative on Malnutrition; SGA, Subjective Global Assessment.

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Body Composition Assessment

• <u>Bioimpedance analysis (BIA)</u>, which derives fat mass and fat-free mass from hydration status using validated formulae, is used routinely.

 Also, the use of other imaging techniques, including dual-energy X-ray absorptiometry (DEXA), MRI and USG of quadriceps muscle, have been proposed <u>but their feasibility and reliability remain questionable at the time of publication</u>.

 The gold standard for the measurement of body composition changes in patients with cancer is the analysis of tissue density using a <u>computed</u> <u>tomography scan</u> at the level of the 3. lumbar vertebra. OS, defined as the time between PD and death. Patients were observed until death or 1 September 2015, at which time they were censored. Survival data was obtained from

To determine skeletal muscle mass at the L3 level, the cross-sectional skeletal muscle surface (cm²) was identified and quantified by HU thresholds of -29 to +150.^{1,3,7}

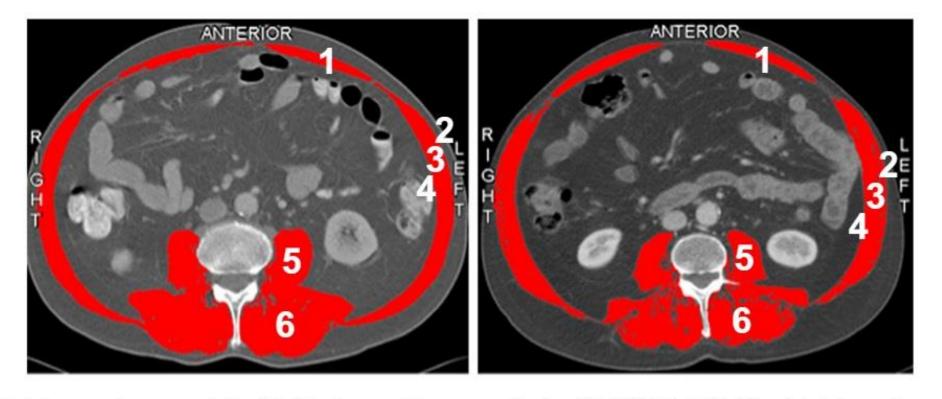


FIG. 1 Computed tomography scans at the third lumbar vertebrae level of two male patients. *Right* Patient with a low skeletal muscle index (SMI, 56.8) (muscle mass) but a normal muscle attenuation index (MAI, 49.8) (muscle quality). *Right* Patient with a normal SMI

and a low MAI (MAI, 24.0). The skeletal muscle area is highlighted in *red*. 1 rectus abdominis, 2 external oblique, 3 internal oblique, 4 transverse abdominal, 5 psoas, 6 paraspinal



Osteoporosis and Sarcopenia

Osteoporesis Sarcopenia

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Original article

Sarcopenia and mortality in cancer: A meta-analysis

Philip Chun-Ming Au ^{a, 1}. Hang-Long Li ^{d, 1}. Grace Koon-Yee Lee ^{a, 1}. Gloria Hoi-Yee Li ^a.



Pooled hazard ratios of low lean mass on mortality according to cancer type.

Cancer type	Number of studies	Overall (HR [95% CI]), I ²
Bile duct (excludes intrahepatic)	2	2.58 [1.82, 3.64], 0%
Breast	3	1.69 [0.79, 3.58]; 61%
Gastrointestinal	18	1.56 [1.36, 1.78]; 48%
Head and neck	1	1.92 [1.19, 3.11]; NA
Hematopoietic	3	1.34 [0.51, 3.53]; 73%
Liver and intrahepatic bile duct	17	2.22 [1.86, 2.65]; 24% ^a
Lung	5	2.19 [1.28, 3.75]; 60%
Ovarian and endometrium	6	1.24 [0.91, 1.70]; 49% ^b
Pancreatic	8	1.63 [1.44, 1.84]; 0%
Prostate	1	0.90 [0.54, 1.50]; NA
Urinary tract	12	1.88 [1.52, 2.34]; 41%
Mixed	5	1.19 [1.03, 1.38]; 61%
Overall	81	1.68 [1.55, 1.83]; 63%

HR, hazard ratio; CI, confidence interval; NA, not applicable.

Results: Altogether 100 studies evaluated the association between lean mass and cancer mortality. The overall pooled HR on cancer mortality was 1.41 (95% CI, 1.24 to 1.59) for every standard deviation decrease in lean mass and 1.69 (95% CI, 1.56 to 1.83) for patients with sarcopenia (binary cutoffs). Overall mortality was also significantly associated with sarcopenia in across various cancer types, except for hematopoietic, breast, ovarian and endometrial, and prostate cancer.

^a Higashi 2016 performed subgroup analysis, hepatocellular carcinoma subgroup was chosen (appendix p27, ref 58).

^b Rutten 2017 included both lean mass measurements, L3 Skeletal Muscle Index

The key is to regularly measure changes in body compartments



Exact measurement of body compartments is difficult

The key is to regularly measure changes in body compartments



Patients should be empowered

and responsible for monitoring their body weight (BW)
 every 2-3 weeks, and immediately report non-volitional weight loss >5% of their usual BW. It is also advisable that changes in their functional ability be reported even in the absence of significant weight loss

The metabolic-nutritional pathway

Regular nutritional screening at diagnosis, on hospital admission and each visit

*Early identification of nutritional deficits and patients who had high risk for malnutrition, then close follow up *Patients education *Solving all causes of malnutrition

Follow-up periodical reevaluation

Nutritional assessment

- Professional counseling:
 dietitian
 - Nutrition care plan





(enteral tubes, parenteral infusions), if needed (eg, short bowel syndrome, oral mucositis, etc)



Oral nutritional supplements Physical activity