

The Global leadership initiative on malnutrition (GLIM) A new diagnostic tool for malnutrition in clinical practice

Dr.med. Mohammed Barigou

Endocrinology diabetes and metabolism division_ Clinical nutrition unit CHU Vaudois_ University of Lausanne





Why the GLIM?

- A core leadership committee with representatives of several of the global clinical nutrition societies was created in January 19th 2016:
- ASPEN (www.nutritioncare.org), aspen
- ESPEN (www.espen.org),
- FELANPE (www.felanpeweb.org)
- PENSA (www.pensa-online.org

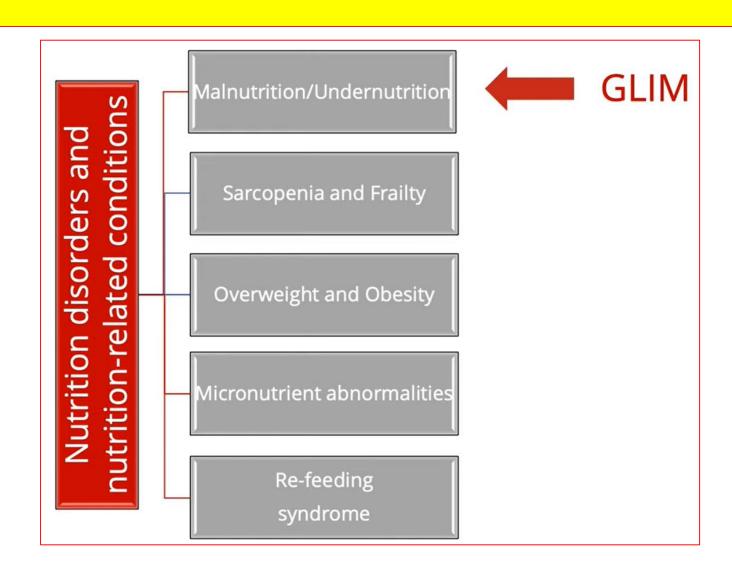


 The core GLIM leadership committee then created a larger supporting working group comprised of invited members that brought additional global diversity and expertise to the consensus effort.

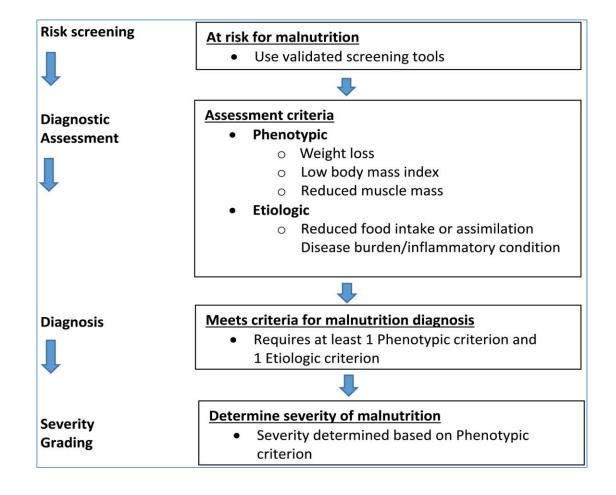
Why the GLIM?

- In order to respond to the needs of the clinical nutrition and medical communities
- Focus on standardizing the clinical practice of malnutrition diagnosis.
- Clarify overlaps with related disease classifications including cachexia.
- The purpose of this specific initiative is to reach global consensus on the identification and endorsement of criteria for the diagnosis of malnutrition in clinical settings.

Spectrum of the GLIM



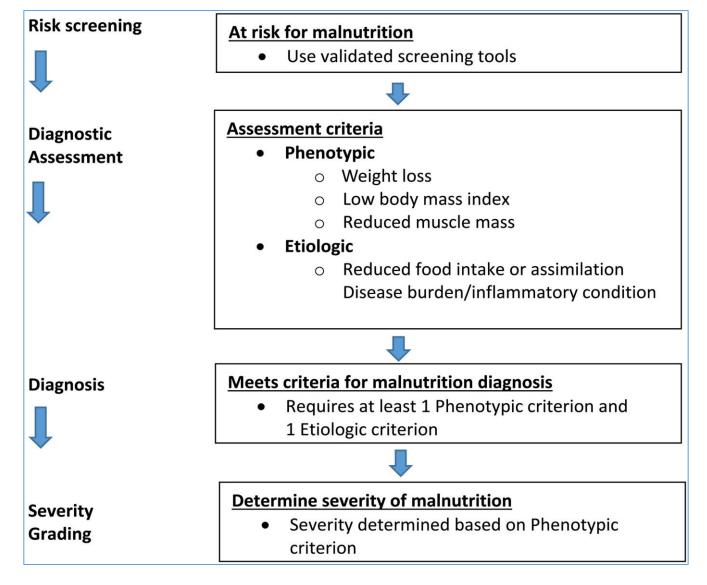
The logic behind the GLIM



Step 1: screening for nutritional risk

	Phenotypic Criteria		Etiologic Criteria			
	Weight loss	Low body mass index (kg/m ²)	Reduced muscle mass/muscle function	Reduced food intake	Severe disease/inflammation	
NRS-2002 ¹²						
Mild	>5% in 3 mo	NS	NA	50-75% of required preceding week	E.g. hip fracture, chronic disease	
Moderate	>5% in 2 mo	18.5–20.5	NA	25-60% of required preceding week	E.g. major abdominal surgery, stroke	
Severe	>5% in 1 mo	<18.5	NA	0–25% of required preceding week	E.g. head injury, bone marrow transplantation, intensive care	
MNA-SF <u>a</u> ²¹						
Mild	1–3 kg in last months	21–23	NS	NS	NS	
Moderate	"Does not know"	19–21	"Does not go out"	Moderate loss of appetite past 3 mo	Mild dementia	
Severe	>3 kg last months	<19	Bed or chair bound	Severe loss of appetite past 3 mo	Acute disease past 3 mo, or severe dementia/depression	
MUST ²²	1	1				
Medium risk	5–10% in 3–6 mo	18.5–20	NA	NS	NA	
High risk	>10% in 3–6 mo	<18.5	NA	Acute illness AND no food intake for >5 d	NA	

GLIM criteria for the diagnosis of malnutrition – A consensus report from the global clinical nutrition community



Step 2: Diagnosis criteria for malnutrition: GLIM

ASPEN/AND7a

SGA4a

Evans 20085c

PEW 200823d

Fearon 20116c

ESPEN 20158a

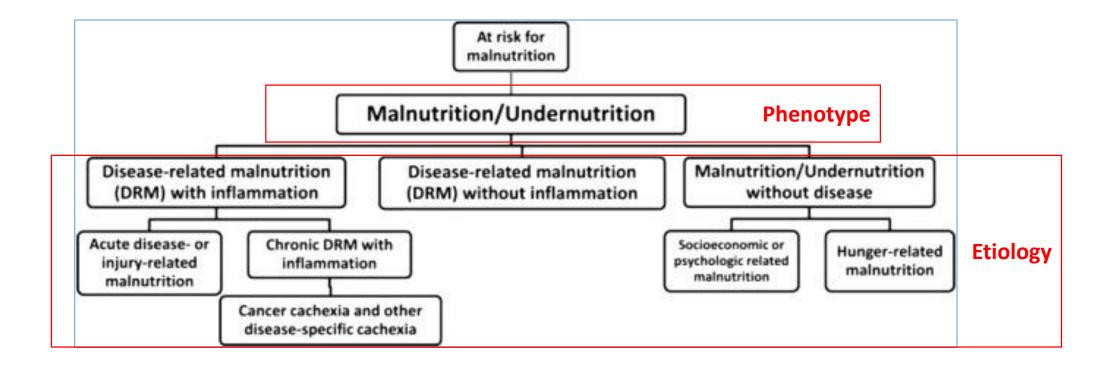
NRS-200212a

MNA-SF21a'b

MUST22a

				_		_	_		
Etiologies	itiologies								
Reduced food intake	Х	Х	Х	Х	Х	Х		X	Х
Disease burden/inflammati on	Х	Х	Х	Х	Х	Х	X	Х	Х
Symptoms									
Anorexia		Х				X	X		Х
Weakness		Х				X	X		
Signs/Phenotype									
Weight loss	X	X	X	X	X	X	X	X	X
Body mass index	Х	Х	Х	Х			Х	Х	X
Lean/fat free/muscle mass		Х		Х	X	х	X	X	Х
Fat mass					X	X		X	
Fluid retention/ascites					X	X			
Muscle function; e.g. grip strength					Х	Х	Х		
Biochemistry							X	X	

Step 2: Diagnosis criteria for malnutrition: GLIM





Contents lists available at ScienceDirect

Clinical Nutrition

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



ESPEN Endorsed Recommendation

GLIM criteria for the diagnosis of malnutrition — A consensus report from the global clinical nutrition community[★]

```
T. Cederholm <sup>a, b, *, 1</sup>, G.L. Jensen <sup>c, 1</sup>, M.I.T.D. Correia <sup>d</sup>, M.C. Gonzalez <sup>e</sup>, R. Fukushima <sup>f</sup>, T. Higashiguchi <sup>g</sup>, G. Baptista <sup>h</sup>, R. Barazzoni <sup>i</sup>, R. Blaauw <sup>j</sup>, A. Coats <sup>k, 1</sup>, A. Crivelli <sup>m</sup>, D.C. Evans <sup>n</sup>, L. Gramlich <sup>o</sup>, V. Fuchs-Tarlovsky <sup>p</sup>, H. Keller <sup>q</sup>, L. Llido <sup>r</sup>, A. Malone <sup>s, t</sup>, K.M. Mogensen <sup>u</sup>, J.E. Morley <sup>v</sup>, M. Muscaritoli <sup>w</sup>, I. Nyulasi <sup>x</sup>, M. Pirlich <sup>y</sup>, V. Pisprasert <sup>z</sup>, M.A.E. de van der Schueren <sup>aa, ab</sup>, S. Siltharm <sup>ac</sup>, P. Singer <sup>ad, ae</sup>, K. Tappenden <sup>af</sup>, N. Velasco <sup>ag</sup>, D. Waitzberg <sup>ah</sup>, P. Yamwong <sup>ai</sup>, J. Yu <sup>aj</sup>, A. Van Gossum <sup>ak, 2</sup>, C. Compher <sup>al, 2</sup>, GLIM Core Leadership Committee, GLIM Working Group<sup>3</sup>
```

Risk screening



Use validated screening tools



Diagnostic Assessment



Assessment criteria

Phenotypic

At risk for malnutrition

- Weight loss
- Low body mass index
- Reduced muscle mass
- Etiologic
 - Reduced food intake or assimilation
 Disease burden/inflammatory condition



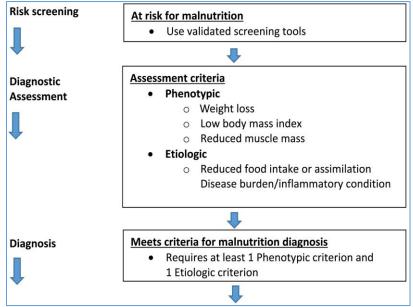
Phenotypic Criteria		Etiologic Criteria		
Weight loss (%)	Low body mass index (kg/m²)	Reduced muscle mass	Reduced food intake or assimilation	Inflammation
>5% within past 6 months, or>10% beyond 6 months	<20 if <70 years, or <22 if >70 years	Reduced by validated body composition measuring	•	Acute disease/injuryd or chronic disease-related
	Asia: <18.5 if <70 years, or <20 if >70 years	techniques	>2 weeks, or any chronic GI condition that adversely impacts food assimilation or absorption	

Step 2: Diagnosis criteria for malnutrition: GLIM

Body composition cut-offs

	Males	Females
Appendicular Skeletal Muscle Index (ASMI, kg/m²)	<7.26	<5.25
ASMI, kg/m ²	<7	<6
ASMI, kg/m ²		
- DXA	<7	<5.4
- BIA	<7	<5.7
Fat free mass index (FFMI, kg/m²)	<17	<15
Appendicular lean mass (ALM, kg)	<21.4	<14.1
Appendicular lean mass adjusted for BMI = ALM/BMI	<0.725	<0.591

Step 3: Grading the severity



	Phenotypic Criteria <u>a</u>					
	Weight loss (%)	Low body mass index (kg/m ²)b	Reduced muscle mass <u>c</u>			
Stage 1/Moderate Malnutrition (Requires 1 phenotypic criterion that meets this grade)	5–10% within the past 6 mo, or 10–20% beyond 6 mo	<20 if <70 yr, <22 if ≥70 yr	Mild to moderate deficit (per validated assessment methods – see below)			
Stage 2/Severe Malnutrition (Requires 1 phenotypic criterion that meets this grade)	>10% within the past 6 mo, or >20% beyond 6 mo	<18.5 if <70 yr, <20 if ≥70 yr	Severe deficit (per validated assessment methods – see below)			

Glim checklist

	Phenotypic criteria	Check if present
Unintentional	> 5% within past 6 months	
Weight loss (%)	> 10% beyond 6 months	
BMI (kg/m²)	< 20 if < 70 years (Asia: < 18.5)	
	< 22 if ≥ 70 years (Asia: < 20)	
Muscle mass	Reduced	
	Etiologic criteria	Check if

	Etiologic criteria	Check if present
Reduced food	Ingestion ≤ 50% of needs from 1 to 2 weeks	
intake	Any reduction for > 2 weeks	
or Assimilation	Any chronic GI condition that adversely impacts food assimilation or absorption	
Disease burden/ Inflammation	Presence of acute disease/injury or chronic disease related	

Malnutrition: if at least one criterion was checked in each section

Determine Malnutrition Severity							
Severity Grade	Phenotypic Criteria						
	Unintentional Weight Loss (%)	Low BMI (kg/m²) a	Reduced Muscle Mass				
Stage 1: Moderate Malnutrition Patient requires 1 phenotypic criterion that meets this grade.	• 5-10% in 6 months; or • 10-20% in more than 6 months	• <20 if <70 years; or • <22 if ≥70 years	Mild-to-moderate deficit (per validated assessment methods on previous page)				
Stage 2: Severe Malnutrition Patient requires 1 phenotypic criterion that meets this grade.	>10% in 6 months; or>20% in more than 6 months	• 18.5 if <70 years; or • <20 if ≥70 years	Severe deficit (per validated assessment methods on previous page)				

What are the advantages of GLIM over other diagnostic criteria?

Combination of criteria to diagnose malnutrition

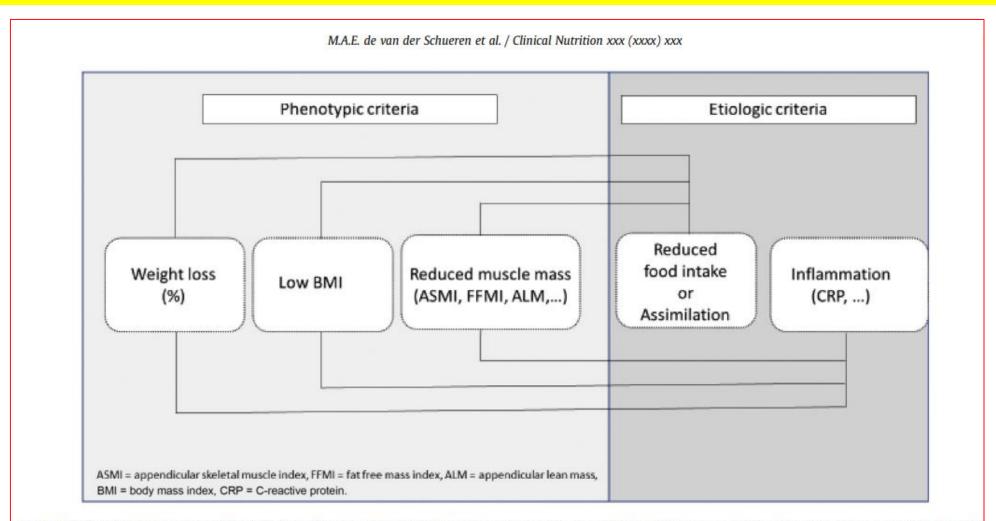
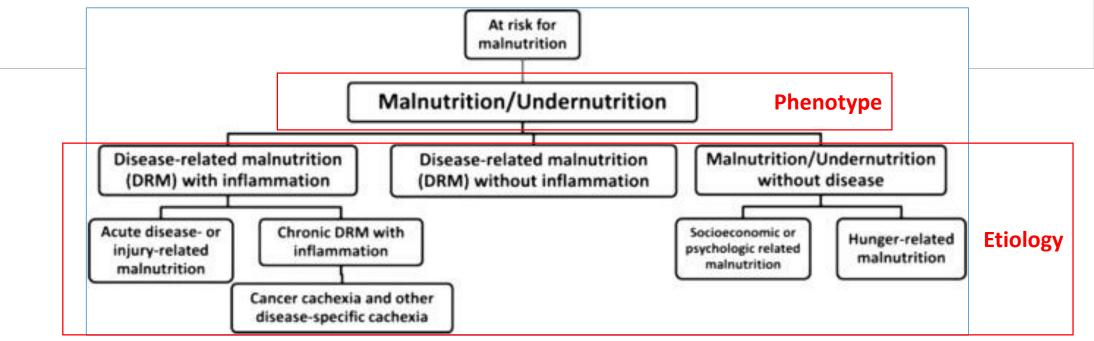


Fig. 1. Potential combinations of GLIM criteria for validation. ALM, appendicular lean mass; ASMI, appendicular skeletal muscle index; BMI, body mass index; CRP, C-reactiv protein; FFMI, fat-free mass index; GLIM, Global Leadership Initiative on Malnutrition. Figure adapted with permission from Reference [32].

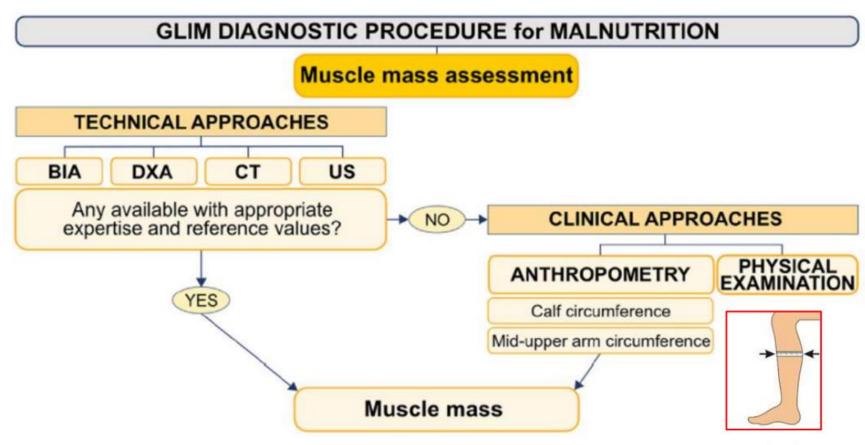
Takes into account the concept of disease related malnutrition

Malnutrition related to

- Chronic disease with inflammation
- Chronic disease with minimal or no perceived inflammation
- Acute disease or injury with severe inflammation
- Starvation including hunger/food shortage associated with socio-economic or environmental factors

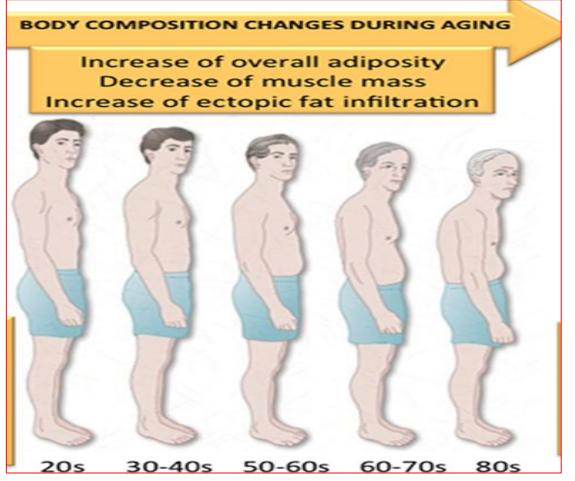


Takes into account the modifications of body composition



Barazzoni R et al. Clin Nutr 2022

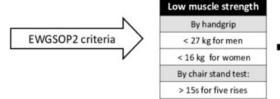
Body composition maters

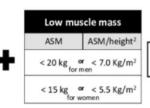


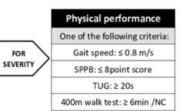


	Males	Females
Appendicular Skeletal Muscle Index (ASMI, kg/m²)	<7.26	<5.25
ASMI, kg/m ²	<7	<6
ASMI, kg/m ²		
- DXA	<7	<5.4
- BIA	<7	<5.7
Fat free mass index (FFMI, kg/m ²)	<17	<15
Appendicular lean mass (ALM, kg)	<21.4	<14.1
Appendicular lean mass adjusted for BMI = ALM/BMI	<0.725	<0.591









Where we are?





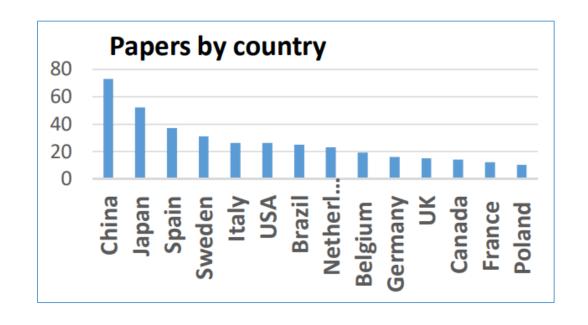
Review 🙃 Full Access

Global Leadership Initiative on Malnutrition (GLIM): Guidance on Validation of the Operational Criteria for the Diagnosis of Protein-Energy Malnutrition in Adults

Heather Keller RD, PhD , Marian A. E. de van der Schueren RD, PhD, for the GLIM Consortium, Gordon L. Jensen MD, PhD, Rocco Barazzoni MD, PhD, Charlene Compher PhD, RD, M. Isabel T. D. Correia MD, PhD, M. Cristina Gonzalez MD, PhD, Harriët Jager-Wittenaar PhD, RD, Matthias Pirlich MD, PhD, Alison Steiber PhD, RDN, Dan Waitzberg MD, PhD, Tommy Cederholm MD, PhD

Where we are?

- Glim bibliometry Jan 2019-2023
- 327 papers in pubmed
- 220 validation studies
- Validation studies:
 - How to define disease: Criterion validity: compared to standard
 - Predictive validity: ability to predict negative outcomes
 - Implementation/ Criterion specification:
 - Muscle mass methodology and cut-off
- burden/inflammation?
- ICD coding for ICD-11 (WHO)

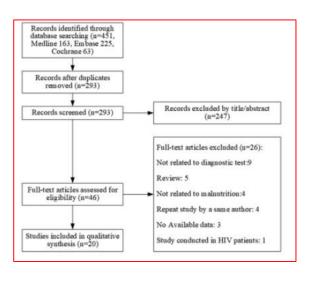


GLIM validation studies

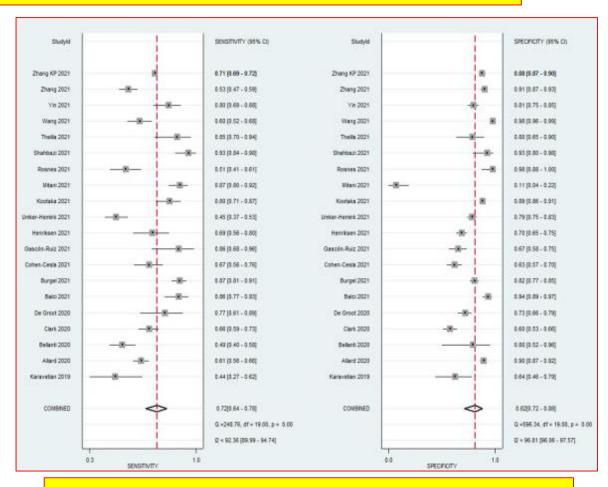
GLIM Criterion validity

GLIM criterion validity:

- 20 studies
- 13 countries
- >10.000 patients
- 7 Cancer
- 8 Inhospital
- 2 CKD
- 2 ICU
- 1 CVD



- 15 studies: SGA or PG-SGA as semi-gold comparator
- Results:
- Amalgamated sensitivity 0.72 (true positives)
- Amalgamated specificity 0.82 (true negatives)



Conclusion: The GLIM criteria "have the potential to be used as the gold standard for diagnosing malnutrition"

GLIM Vs PG-SGA predicitve validity

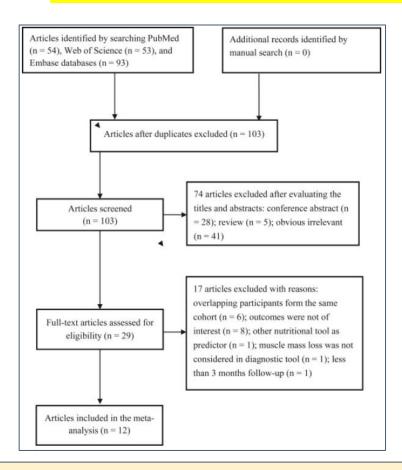
- Exemple of cancer patients
- 468 patients
- 33,7% Vs 39,7% for GLIM
- Results:
 - GLIM was superior to PG-SGA to predict outcomes in cancer patients

	Moderate malnutrition		Severe malnutritio	Severe GLIM vs PG-SGA malnutrition				
	OR (95%CI)	p	OR (95%CI)	p	Changed Log- likelihood	p	AIC	BIC
Severe tox	cicity							
GLIM criteria	2.0(1.1-3.5)	0.023	3.7(1.4– 9.5)	0.008	0.034	0.008	395.653	452.382
PG-SGA	1.5(0.8- 2.7)	0.217	2.9(1.3- 6.3)	0.007	0.026	0.026	397.905	454.634
Discontin	uity of treatr	nent						
GLIM criteria	2.3(0.8– 6.5)	0.121	6.6(1.9– 23.3)	0.003	0.057	0.015	178.821	235.55
PG-SGA	1.7(0.6- 4.9)	0.319	4.7(1.5- 15.4)	0.010	0.042	0.046	181.017	237.746
Artificial nutrition support								
GLIM criteria	1.7(0.8- 3.4)	0.163	5.9(2.2- 15.4)	0.000	0.051	0.002	287.403	344.132
PG-SGA	1.4(0.7- 3.2)	0.365	6.8(2.9– 16.1)	0.000	0.078	<0.001	280.7383	337.467

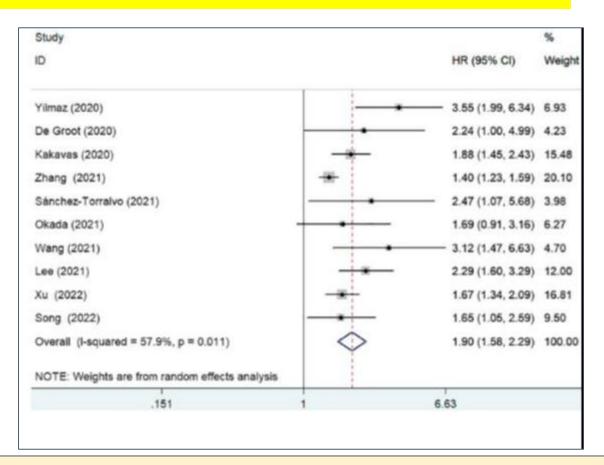
Clinical Nutrition

Volume 41, Issue 4, April 2022, Pages 855-861

GLIM predicitve validity (cancer patients)



Twelve article reporting on 11 studies including 6799 cancer patients were identified.



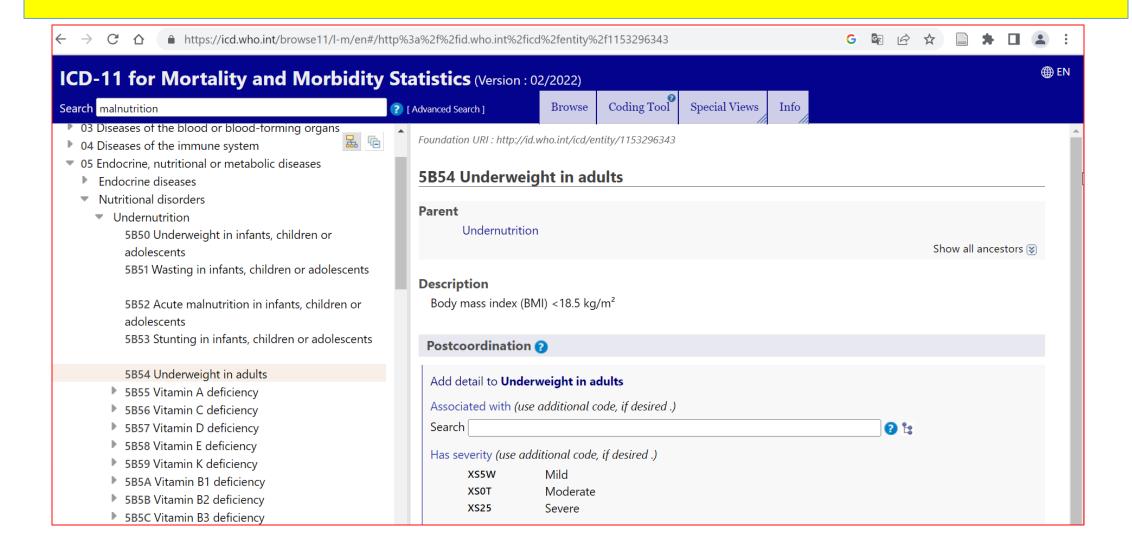
Meta-analysis indicated that malnutrition defined by the GLIM was associated with poor overall survival (HR 1.90; 95% CI 1.58-2.29) and disease-free survival (HR 1.51; 95% CI 1.27-1.79), respectively.

From NRS to GLIM?

Score	ВМІ		3 months weight loss	6 months weight loss	Weight loss beyond 6 months			Disease severity
NRS-2002	✓	√	✓	-	-	-	✓	✓
Glim	√	✓	✓ (included in the 6 months)	✓	✓	-	✓	✓

GLIM and malnutrition codification

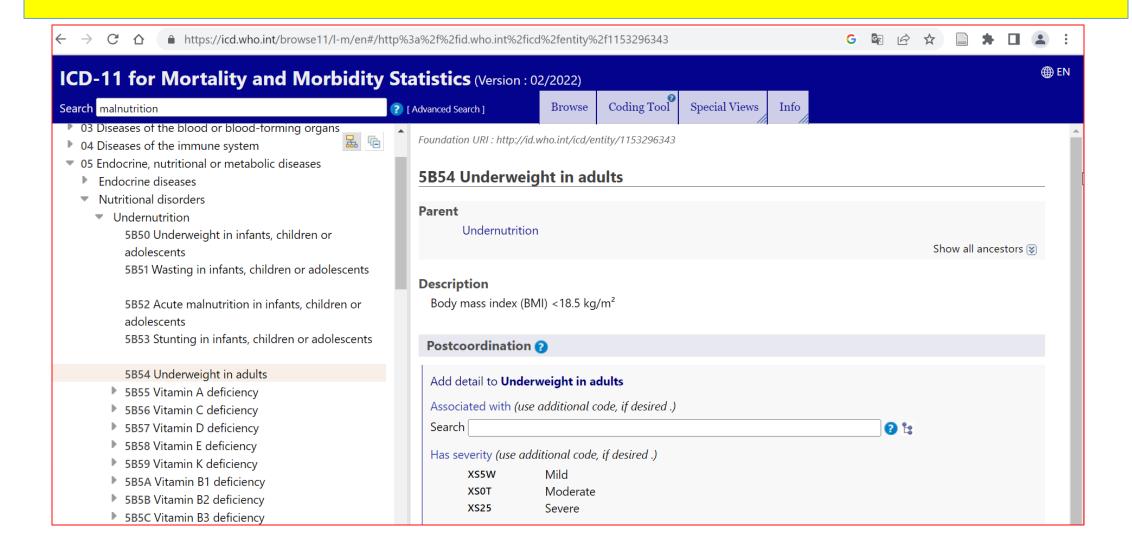
Toward a better codification of malnutrition



In switzerland

	Minimal (E44.1)	Moderate (E44.0)	Severe (E43)
NRS (mandatory)	3	4	≥5
ВМІ	-	18.5-20.5	<18.5kg
Weight loss	> 5% in 3 months	>5% in 2 months	>5% in 1 month
Food intake	50-75%	25-50%	0-25%

Toward a better codification of malnutrition



© Serdi and Springer-Verlag International SAS, part of Springer Nature

A Clinically Relevant Diagnosis Code for "Malnutrition in Adults" Is Needed in ICD-11

T. Cederholm¹, E. Rothenberg², R. Barazzoni³

1. Clinical Nutrition and Metabolism, Department of Public Health and Caring Sciences, Uppsala University, Uppsala, Theme Inflammation & Aging, Karolinska University Hospital, Stockholm, Sweden; 2. Facutly of Health Sciences, Kristianstad University, Kristianstad, Sweden; 3. Department of Medical, Surgical and Health Sciences, University of Trieste, Trieste, Italy

Corresponding Author: Tommy Cederholm, Clinical Nutrition and Metabolism, Department of Public Health and Caring Sciences, Uppsala University, Uppsala, Sweden, tommy.cederholm@pubcare.uu.se

Introduction

alnutrition is a devastating condition with adverse outcomes in terms of complications, prognosis and quality of life (1-5). It is a complex condition with many aetiologies, that evolves separately or from the interaction between food deprivation and catabolic processes linked to disease-related inflammation.

Textbooks define malnutrition in adults as "a state resulting from lack of intake or uptake of nutrition that leads to altered body composition (decreased fat free mass) and body cell mass leading to diminished physical and mental function and impaired clinical outcome from disease". Caring for malnutrition in clinical settings requires attention to prevention, diagnosis and management of nutritional and metabolic alterations.

International Classifications of Diseases

Urgent need of a clinically relevant ICD-11 diagnosis code for malnutrition in adults

Thus, the clinical picture of malnutrition is characterized by a complex combination of various etiologic factors, like food deprivation and inflammatory conditions, and phenotypic expressions like weight loss and muscle wasting. For this reason, the clinical nutrition community needs an ICD coding of "Malnutrition in adults" that reflects the current malnutrition perception, and that may support clinical decision-making in every day practice.

The pending ICD-11 in its present edition does not meet this requirement. Currently, there are two ICD-11 codes relating to malnutrition in adults. "Underweight in adults" (5B54) appears to be the major malnutrition concept for adults, that is described by BMI<18.5 kg/m². Malnourished adults in healthcare in most regions of the world now rarely fulfil this criterion, at least not until late disease stages. On the contrary, many malnourished

ICD-11 new nomenclature proposal (Cederholm et al)

- Proposal of 4 codes that reflect disease related malnutrition:
 - 5B72 Malnutrition in adults: When malnutrition is confirmed by the presence of a combination of phenotypic criteria; e.g. weight loss, low body mass index or reduced muscle mass, and etiologic criteria; e.g. reduced food assimilation (intake/digestion/absorption) or disease burden/inflammation.
 - 5B72.0 Malnutrition in adults related to acute or chronic disease, injury or infection with moderate to severe inflammation
 - 5B72.1 Malnutrition in adults related to disease with <u>non-discernible or low level of inflammation</u>
 - 5B72.2 Malnutrition in adults related to pure starvation

In conclusion

- GLIM criteria are intended to improve screening, diagnosis and grading of malnutrition.
- By composing with phenotypic and etiologic criteria they allow a large possibilities for the diagnosis of malnutrition
- They translate well the concept of disease related malnutrition to clinical practice
- Body composition parameters may allow to diagnose malnutrition in certain circumstances
- Validation studies prove that GLIM is compatible with clinical practice with interesting accuracy
- GLIM should support a better codification of malnutrition compatible with the concept of disease related malnutrition.





et de médecine

Thank you for your



attention



Phenotypic Criteria			Etiologic Criteria		
	Weight loss (%)	Low body Mass Index (kg/m²)	Reduced muscle mass/muscle function	Reduced food intake	Severe disease/inflammation
SGA ⁴	'				
Moderate/Stage B	5–10% past 6 mo	NA	Mild to moderate deficits in function or muscle mass	"Definite decrease"	Yes
Severe/Stage C	>10% past 6 mo	NA	Severe deficit in function and muscle mass	"Severe deficit"	Yes
Evans 2008 ⁵	·				
Cachexia	>5% in <12 mo	<20	Low FFMI, decreased muscle strength	"Anorexia"	Increased CRP/IL6, low serum albumin (<3.2 g/l)
PEW 2008 ²³		<u>'</u>	'		
Protein-energy wasting	>5% in 3 mo, or > 10% in >6 mo	<23	Muscle mass down by 5% last 3 mo, or > 10% in >6 mo. Reduced MAC	Energy intake <25 kcal/kg BW/d for >2 mo	Chronic kidney disease, Serum albumin <3.8 g/dl
Fearon 2011 ⁶	·		<u> </u>		
Precachexia	<5%	NA	NA	"Anorexia"	Metabolic change
Cachexia	>5% in 6 mo (>2%)	<20 (when WL > 2%)	Sarcopenia - ASMI 7.26/5.45 kg/m ² (m/w) when WL >2%	"Often reduced food intake"	Cancer with catabolic drive (systemic inflammation)
ASPEN/AND 2012 ⁷		·			
Moderate	1–2% in 1 w to 20% in 1 y	NA	Mild muscle loss	<75% of ER for 7 d-3 mo	Yes
Severe	>2% in 1 week to >20% in 1 year	NA	Moderate to severe muscle loss, or reduced grip strength	e <50% of ER for >5 d- < 1 mo	Yes
ESPEN 2015 ⁸					
Malnutrition	>5% past 3 mo, or > 10%	<18.5, or <20 (<70v)/ <22(>70v)	FFMI <15 kg/m ² (f), <17 kg/m ² (m)	According to any validated tool	NA