

Assessment of malnutrition with bioimpedance analysis (BIA) and its prognostic impact in patients with ischemic stroke

Ilaria De Negri¹, Luca Antonelli¹, Simone Dal Bello¹, Laura Ceccarelli¹, Yan Tereshko², Gianluigi Gigli¹, Mariarosaria Valente¹, Giovanni Merlino²

¹Clinical Neurology, Udine University Hospital and DMED, University of Udine, Udine, Italy.

²Stroke Unit, Udine University Hospital, Department of Head, Neck, and Neurosciences, Udine University Hospital, Udine, Italy.

Background and Aims

The relationship between malnutrition and poor outcomes in stroke patients is well-established.

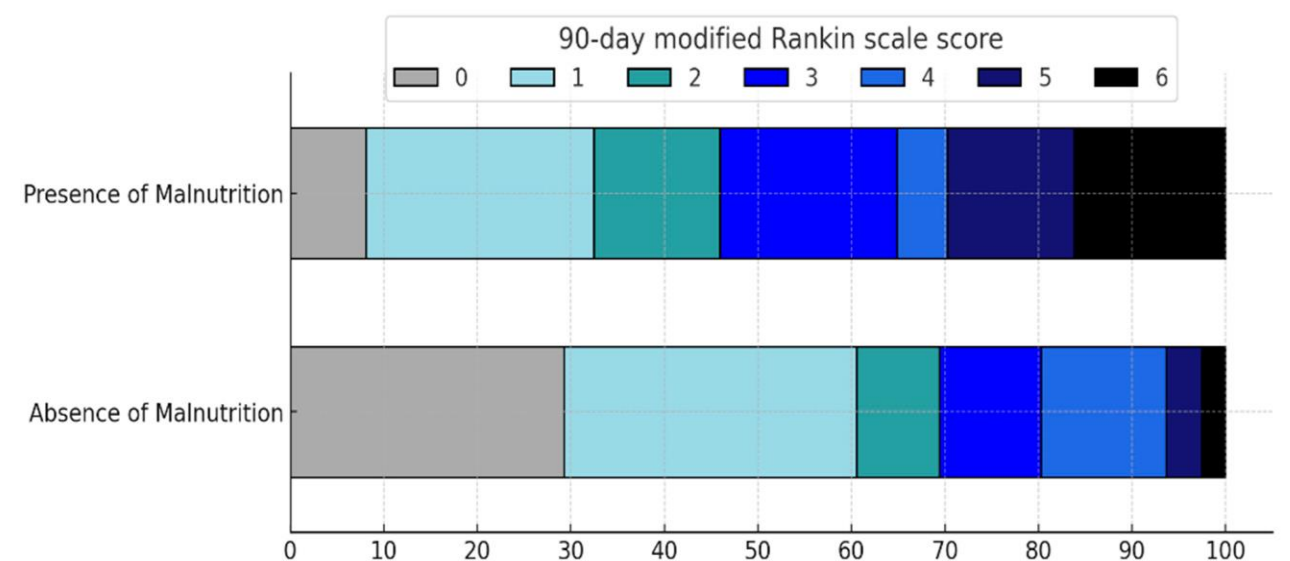
Nutritional status assessment has traditionally relied on composite lab-based scores. Bioimpedance analysis (BIA) and bioelectrical impedance vector analysis (BIVA) offer non-invasive, cost-effective, and rapid evaluations of body composition, nutritional, and hydration status. This study aimed to compare the ordinal distribution of mRS scores 90 days after ischemic stroke between malnourished and non-malnourished patients according to BIVA parameters.

Methods

We conducted a single-center prospective observational study on patients admitted for ischemic stroke between April 1 and September 30, 2024. Using inverse probability weighting (IPW) and ordinal logistic regression, we analyzed the association between malnutrition and mRS outcomes.

Weighted ordinal logistic regression according to the mRS scale at 90 days post-ischemic cerebrovascular event (Legend: mRS = modified Rankin Scale; NIHSS = National Institutes of Health Stroke Scale; IVT = intravenous thrombolysis; LDL = low-density lipoprotein).

mRS Shift (Univariate)			
Predictors	Common Odds Ratio	CI	p
Presence of malnutrition	3.34	1.74–6.41	0.001
mRS Shift (Multivariate)			
Predictors	Adjusted Common Odds Ratio	CI	p
Presence of malnutrition	2.79	1.37–5.70	0.005
Female sex	0.85	0.48–1.51	0.580
NIHSS at admission (per unitary increase)	1.19	1.11–1.28	<0.001
IVT	0.28	0.15–0.52	<0.001
Mechanical thrombectomy	0.41	0.14–1.18	0.097
Lymphocytes (per unitary increase)	1.01	1.00–1.02	0.027
Albumin (per unitary increase)	0.82	0.75–0.89	<0.001
Total cholesterol (per unitary increase)	0.99	0.97–1.02	0.875
LDL (per unitary increase)	1.01	0.98–1.03	0.742



Results

Among 195 patients, 37 (19%) were malnourished upon Stroke Unit admission based on BIVA parameters. Malnutrition was associated with higher mRS scores at 90 days (cOR 3.34, 95% CI 1.74–6.41; $p=0.001$). Adjusted analyses confirmed malnutrition as an independent predictor of unfavorable outcomes (acOR 2.79, 95% CI 1.37–5.70; $p=0.005$). Other significant predictors included NIHSS score (acOR 1.19, 95% CI 1.11–1.28; $p<0.001$), intravenous thrombolysis (acOR 0.28, 95% CI 0.15–0.52; $p<0.001$), lymphocyte count (acOR 1.01, 95% CI 1.00–1.02; $p=0.027$), and albumin concentration (acOR 0.82, 95% CI 0.75–0.89; $p<0.001$).

Conclusion

BIVA-identified malnutrition is a significant predictor of poor post-stroke outcomes. Implementing BIVA in clinical practice may enhance early risk stratification and guide tailored nutritional interventions to improve recovery trajectories.

Disclosures: Ilaria De Negri: nothing to disclose; Luca Antonelli: nothing to disclose; Simone Dal Bello: nothing to disclose; Laura Ceccarelli: nothing to disclose; Yan Tereshko: nothing to disclose; Gianluigi Gigli: nothing to disclose; Mariarosaria Valente: nothing to disclose; Giovanni Merlino: nothing to disclose.

Young Investigator Award Application: No

References:

- Liu, P.; Tian, H.; Ji, T.; Zhong, T.; Gao, L.; Chen, L. Predictive Value of Malnutrition, Identified via Different Nutritional Screening or Assessment Tools, for Functional Outcomes in Patients with Stroke: A Systematic Review and Meta-Analysis. *Nutrients* 2023, 15, 3280.
- Piccoli, A.; Pillon, L.; Dumler, F. Impedance vector distribution by sex, race, body mass index, and age in the United States: Standard reference intervals as bivariate Z scores. *Nutrition* 2002, 18, 153–167.
- Piccoli, A. Identification of operational clues to dry weight prescription in hemodialysis using bioimpedance vector analysis. *Kidney Int.* 1998, 53, 1036–1043.