



Background

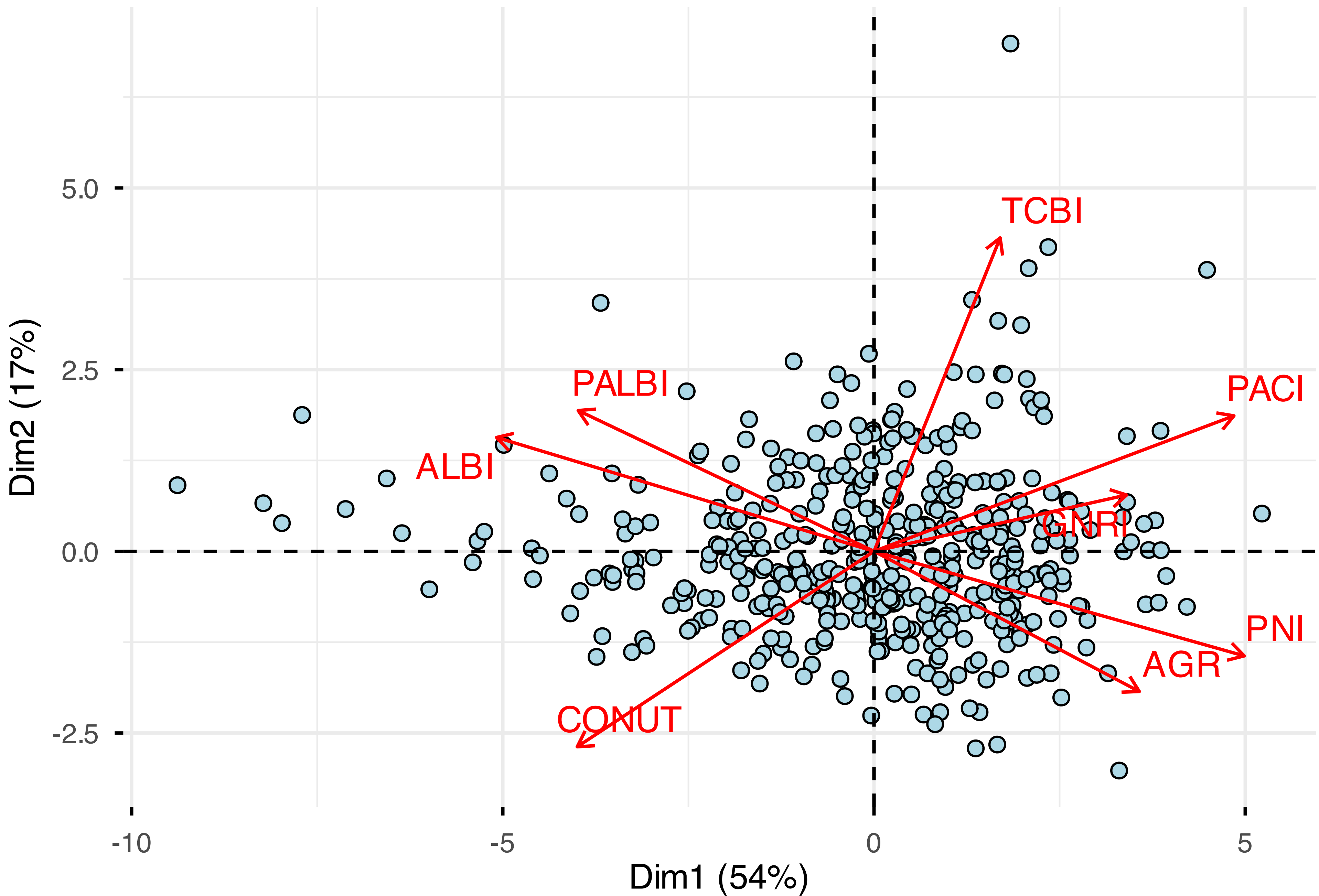
Pre-procedural nutritional status has been proposed as a prognostic marker in transcatheter aortic valve replacement (TAVR), although most studies rely on isolated nutrition scores rather than on a multidimensional assessment. Among the established serum-based indices, including the CONUT score, PNI, and GNRI, their comparative prognostic utility in TAVR populations remains largely unexplored. This study examined how multiple serum-based nutritional and inflammatory scores characterize patients with severe aortic stenosis and whether these scores influence post-TAVR survival.

Methods

Consecutive patients undergoing TAVR between December 2021 and December 2023 were retrospectively analyzed. Exclusion criteria were emergency procedures, endocarditis, active or prior cancer, immunocompromised status, redo aortic interventions, predominant aortic regurgitation, non-transfemoral access, concomitant coronary revascularization, and conversion to surgery. Eight serum biomarker-based scores were calculated for each patient (GNRI, PNI, CONUT, PALBI, TCBI, PACI, AGR, ALBI). Associations with overall survival were assessed with Kaplan–Meier curves and multivariable Cox regression adjusted for age, sex, and STS score. Relationships among scores were evaluated using principal component analysis (PCA).

Results

A total of 459 patients were included (mean age 81.7 ± 7.1 years; 45.8% female; median STS 2.8 [1.8–4.4]). The eight considered scores frequently classified the same patients into different risk strata.



Multivariable Cox regression

Score	OR (95% CI)	P-value
GNRI	0.977 [0.956; 0.999]	0.037
PNI	0.932 [0.896; 0.969]	<0.001
CONUT	1.349 [1.137; 1.599]	<0.001
PALBI	1.816 [0.991; 3.330]	0.054
TCBI	1.000 [0.999; 1.000]	0.090
PACI	0.942 [0.916; 0.969]	<0.001
AGR	0.243 [0.129; 0.456]	<0.001
ALBI	2.101 [1.346; 3.280]	0.001
PCA	0.833 [0.764; 0.909]	<0.001

Cox regression adjusted for: age, sex, BMI, and STS score

Conclusions

Despite capturing overlapping biological domains, serum-based nutritional scores inconsistently classify TAVR patients and show limited association with overall mortality in this cohort. Their prognostic value appears weak when adjusted for baseline risk. Principal component analysis suggests that a single underlying biomarker dimension explains most of the variability observed among scores, supporting the development of streamlined or composite indices rather than relying on individual metrics.