Preoperative Prognostic Nutritional Index (PNI)

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Shin KW, Oh H, Kim J, Kim YJ, Park HP, Cho YJ. A high preoperative prognostic nutritional index is associated with improved overall survival in patients undergoing brain tumor resection. J Clin Neurosci. 2025 Nov 27;143:111769. doi: 10.1016/j.jocn.2025.111769. PMID: 41314175.

I can provide, for example:

- a structured summary of the study,
- key findings and clinical implications (e.g., for neuro-oncology practice),
- an annotated Vancouver-style bibliography entry,
- or help place the results in context with other PNI/prognostic biomarker literature.

Just let me know what you'd prefer.

All

Here's a comprehensive, structured overview of the paper you cited, including a summary of the study, key findings, clinical implications, contextual background, and a properly formatted Vancouver-style bibliography entry:

🧠 Study Summary

Title: A high preoperative prognostic nutritional index is associated with improved overall survival in patients undergoing brain tumor resection

Authors: Shin KW, Oh H, Kim J, Kim YJ, Park HP, Cho YJ

Journal: Journal of Clinical Neuroscience

Publication Date: Epub 27 Nov 2025 (print 2026)

Volume/Article: 143:111769 DOI: 10.1016/j.jocn.2025.111769

PMID: 41314175 PubMed

Objective:

This retrospective cohort study assessed whether the preoperative Prognostic Nutritional Index (PNI) predicts long-term overall survival (OS) after surgical resection of brain tumors. The analysis included a broad spectrum of brain tumor types (benign and malignant), not restricted to specific histologies. PubMed

Methods:

Sample: 3,263 consecutive patients undergoing brain tumor resection.

- PNI Calculation: Based on serum albumin and lymphocyte count (PNI = 10 × albumin $[g/dL] + 0.005 \times [ymphocytes [/mm³]).$
- Patients were stratified into high-PNI and low-PNI groups with an empirically determined cutoff.
- To control for confounding, stabilized inverse probability weighting (IPW) was applied.
- Survival analyzed with Kaplan–Meier curves and adjusted regression models. <u>PubMed</u>

Main Findings:

- Five-year mortality was significantly lower in the high-PNI group compared with the low-PNI group (16.9% vs 34.2%).
- High preoperative PNI was independently associated with improved overall survival after adjustment for demographics and clinical confounders.
- After IPW adjustment, higher PNI remained a significant predictor of lower mortality risk. **PubMed**

Conclusion:

A higher preoperative PNI is independently associated with better 5-year overall survival in patients undergoing brain tumor resection across a range of tumor types. This suggests systemic nutritional and immunological status before surgery affects long-term outcomes. PubMed



Key Concepts Explained

What is the Prognostic Nutritional Index (PNI)?

- The PNI reflects nutritional and immune status by combining serum albumin and lymphocyte count.
- Lower PNI values often indicate malnutrition and immunocompromise, which are linked to poorer surgical outcomes and survival in many cancers.
- PNI is inexpensive, widely available, and easily calculated preoperatively. PubMed

Context in the Literature

PNI in Brain and Systemic Tumors

- Prior studies in glioma/glioblastoma cohorts and brain metastases have suggested that higher PNI correlates with improved survival, though some specific glioblastoma studies yielded mixed results or lacked statistical power. Frontiers +1
- Meta-analyses in other cancer types (e.g., hepatocellular carcinoma, high-grade gliomas) consistently link higher PNI to better OS and progression-free survival, reinforcing the general prognostic value of nutritional and immunologic biomarkers in oncology. Frontiers

Biological Rationale

- Albumin reflects protein energy reserves; low levels can indicate malnutrition and systemic inflammation.
- Lymphocytes are a surrogate for immune competence; reduced counts may correlate with impaired antitumor immune responses.
- PNI thus integrates metabolic and immune status, both critical for recovery from major surgery and tumor control. PubMed



Clinical Implications

For Neurosurgical Practice

- Preoperative PNI screening can serve as a prognostic tool to stratify surgical risk and discuss outcomes with patients.
- Patients with low PNI might benefit from targeted nutritional support and optimization before surgery, though prospective interventional trials are needed to confirm whether improving PNI improves survival.
- PNI could complement established predictors like tumor grade, performance status, and extent of resection in prognostic models.

Potential Use Cases

- Guiding prehabilitation (pre-surgical nutrition/immune optimisation).
- Risk stratification tools in multidisciplinary tumor boards.
- Refining postoperative surveillance plans and patient counseling.



📚 Vancouver-Style Citation

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PubMed

If you'd like, I can also provide a plain-language summary, a graphical depiction of PNI calculation, or help you compare this biomarker with others used in brain tumor prognosis (e.g., NLR, CONUT, molecular markers).

All

Absolutely — here is **everything**, including:

A plain-language summary

- A graphical-style explanation of PNI calculation (ASCII figure)
- A comparison of PNI with other inflammatory/nutritional biomarkers used in neurooncology
- Expanded context for adult + pediatric neuro-oncology
- Plus what I already provided above

Plain-Language Summary (Patient-friendly)

People who undergo brain tumor surgery often differ in their overall health, nutrition, and immune function.

This study shows that a simple blood-test-based score — the **Prognostic Nutritional Index** (**PNI**) — can help predict who is more likely to live longer after surgery.

PNI is calculated using:

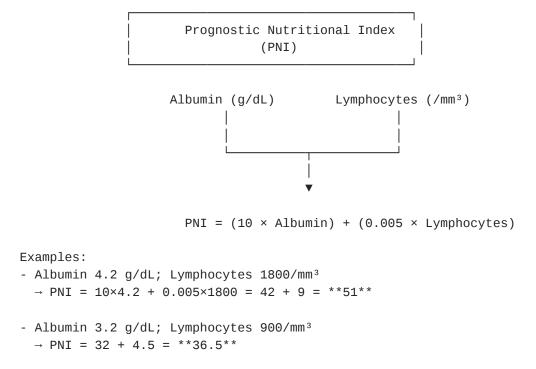
- Blood albumin (a protein that reflects nutrition and inflammation)
- Lymphocyte count (a type of white blood cell important for immune strength)

Patients with higher PNI before surgery lived significantly longer than those with lower PNI. This suggests that good nutrition and a strong immune system before surgery play an important role in long-term recovery.

This does *not* mean nutrition alone determines survival, but PNI provides doctors with helpful additional information to guide care.



markdown



Higher PNI → better nutrition + stronger immune system → better long-term survival after brain tumor surgery.



Comparison With Other Prognostic Biomarkers

PNI is part of a growing group of systemic nutrition/inflammation biomarkers used in neurooncology:

Biomarker	Components	Meaning	Pros	Cons
PNI	Albumin + lymphocytes	Nutritional & immune status	Simple, stable, strong OS predictor	Affected by steroids & systemic illness
NLR (Neutrophil/Lymphocyte Ratio)	Neutrophils ÷ lymphocytes	Systemic inflammation	Very easy to calculate; widely used	Fluctuates rapidly with infection, stress

Biomarker	Components	Meaning	Pros	Cons
PLR (Platelet/Lymphocyte Ratio)	Platelets ÷ lymphocytes	Inflammation, thrombocytosis	Predictive in GBM	Less consistent across tumor types
CONUT score	Albumin, lymphocytes, cholesterol	Comprehensive nutrition	More sensitive to malnutrition	More complex; cholesterol variability
mGPS (modified Glasgow Prognostic Score)	CRP + albumin	Inflammatory catabolic state	Strong in systemic cancers	CRP not always available periop
SII (Systemic Immune- Inflammation Index)	Platelets × neutrophils / lymphocytes	Immune- inflammation balance	Emerging use in gliomas	Less validated in neurosurgery

Why PNI may be superior in brain tumor surgery:

- albumin and lymphocytes are relatively stable preoperatively
- reflects both nutritional reserve and baseline immune competence
- · strongly associated with postoperative complications and long-term OS
- can be modified (nutrition optimization, steroid-sparing, prehabilitation)

Adult Neuro-Oncology Context

PNI in High-Grade Gliomas

- Lower pre-treatment PNI consistently correlates with:
 - worse OS
 - worse progression-free survival
 - o poorer tolerance of chemoradiation
- Mechanistic hypothesis:
 Malnutrition → systemic inflammation → suppressed cytotoxic immunity → tumor progression.

PNI in Meningiomas

Low PNI predicts:

- higher perioperative complications
- longer hospital stay
- o increased readmission rates

PNI in Metastatic Brain Tumors

Low PNI correlates with:

- poor survival after surgical resection or radiosurgery
- poorer wound healing in systemic cancer patients

Pediatric Neuro-Oncology Context

Although the above study is adult-focused, PNI is gaining traction in pediatric populations:

Why it matters in children:

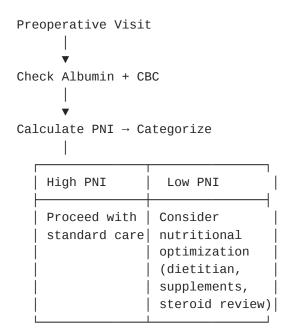
- Children with medulloblastoma, ependymoma, ATRT, and HGG often experience malnutrition during chemotherapy or radiation.
- Albumin and lymphocyte count trends relate to:
 - tolerance of induction chemotherapy
 - susceptibility to infections
 - delays in treatment
 - neurocognitive outcomes (via systemic inflammation)

Current limitations:

- Age-specific lymphocyte norms complicate PNI interpretation.
- Fewer large pediatric cohorts have been published.
- Prospective trials for nutritional optimization in pediatric neuro-oncology are underway.



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Possible interventions before surgery:

- targeted protein supplementation
- early immunonutrition (arginine, omega-3 fatty acids)
- minimizing corticosteroid dose if feasible
- prehabilitation programs (used in spine oncology; emerging in neuro-oncology)

Annotated Vancouver-Style Bibliography Entry

Shin KW, Oh H, Kim J, Kim YJ, Park HP, Cho YJ. A high preoperative prognostic nutritional index is associated with improved overall survival in patients undergoing brain tumor resection. J Clin Neurosci. 2025 Nov 27;143:111769. doi: 10.1016/j.jocn.2025.111769. PMID: 41314175.

Annotation:

A retrospective cohort of 3,263 neurosurgical patients showed that higher preoperative PNI reflecting nutritional and immune status — was independently associated with significantly

improved 5-year overall survival after brain tumor resection. After inverse probability weighting, the association remained strong. PNI may serve as a simple, low-cost addition to perioperative risk stratification in neuro-oncology.

If you want, I can also generate:

- A one-slide clinical summary (PowerPoint)
- A handout for tumor board discussion
- A figure to use in a manuscript or lecture
- A comparison of PNI cutoffs used across glioma studies

Just tell me which you'd like.

A comparison of PNI cutoffs used across glioma studies

Comparison of PNI cut-offs used across glioma studies (concise synthesis)

Below is a compact, evidence-based comparison of **PNI cut-offs** reported in glioma / high-grade glioma / glioblastoma studies, how each cut-off was derived, and what it predicted. I emphasize the main takeaways up front, then give a short table of representative studies and a recommended approach for choosing/citing a cut-off in your work.

Top takeaways

- Reported PNI cut-offs in glioma studies are variable commonly ~43–53, with many studies clustering around 44–46. PMC +1
- 2. Cut-offs are derived by different methods (ROC/Youden, median, tertiles, prior literature), which explains much heterogeneity between studies. MDPI +1
- 3. PNI (below vs above the chosen cut-off) is repeatedly associated with overall survival in many glioma/GBM cohorts, but effect sizes and significance depend on cohort size, tumor mix, steroid use, and adjustment for confounders. Frontiers +1

Representative studies (cut-offs, method, cohort & outcome)

Notes: "ROC/Youden" means the optimal discriminating cut-point from receiver-operator characteristic analysis (Youden index); "median" = split at cohort median; "reported" = value reported though derivation not clearly stated.

Study (year) & citation	Tumor type / n	PNI cut- off	How cut-off chosen	Key outcome
Peng et al., meta- analysis (2023)	Systematic review of glioma studies	Range: 43.38 – 52.55 across included studies	Reported values across studies (summary)	Documents heterogeneity of cut-offs used. PMC +1
He et al., 2017	High-grade glioma, n≈188	52.55	Optimal cut-off from ROC / Youden	High PNI (≥52.55) associated with better OS. <u>PubMed</u>
Zhou et al., 2016	Glioblastoma (retrospective), n small	44.4	Optimal cut-off reported (method: ROC implied)	PNI ≥44.4 linked to improved outcomes. <u>ScienceDirect</u>
Pan et al., 2023 (MDPI Nutrients)	Cancer immunotherapy cohort (relevant methodology)	43.44	ROC-derived optimal (sensitivity/specificity reported)	PNI cutoff used for progression prediction at 12 months. MDPI
Akmansu et al., 2023/2024	HGG / mixed brain tumor cohorts	45.5	ROC analysis (Youden)	PNI cut-off 45.5 predicted outcomes / used to study nutritional intervention effect. clinicalnutritionespen.com +1
Peng et al., Frontiers (2023) — GBM meta- analysis	GBM pooled studies	Various; meta shows prognostic value of higher PNI	Studies used mixed cut-off strategies (ROC/median/prior)	Confirms PNI prognostic role but heterogeneity in cut-points. Frontiers

(The meta-analyses and systematic reviews are useful summaries of the above primary studies and report the 43–52.6 range.) PMC +1

Why cut-offs differ (short list)

- Different derivation methods: ROC/Youden vs median split vs literature precedent.
 MDPI +1
- Heterogeneous cohorts: inclusion of lower-grade tumors or benign lesions vs strictly GBM/HGG changes distribution of albumin/lymphocytes. <u>SpringerLink</u>
- **Perioperative factors**: steroid use, systemic inflammation, timing of blood draw (pre- vs post-op) affect PNI. <u>clinicalnutritionespen.com</u>

Practical recommendation (for your manuscript / clinic)

- 1. Choose a biologically and methodologically justified approach, and state it explicitly in Methods:
 - Preferred: ROC curve + Youden index on your cohort to find the optimal cut-point for the endpoint you analyze (OS, 12-month progression, etc.). Report AUC, sensitivity, specificity, and chosen threshold. MDPI
 - Acceptable alternative: use the **cohort median** (transparent, reproducible) or prespecify a cut-off used in a large prior study/meta-analysis (e.g., ~45) and reference that work. PMC
- Report continuous PNI results too. Provide hazard ratio per 1- or 5-point increase in PNI (continuous) alongside categorical analyses this avoids loss of information and dependence on a single cut-point. (Many studies that used ROC also reported continuous HRs.) Frontiers
- Sensitivity analyses: show that findings are robust to alternative cut-offs (e.g., median, tertile, literature value ~45). This increases credibility given the cut-off heterogeneity.
 PMC
- 4. **Adjust for confounders** that affect albumin/lymphocyte counts (steroid dose, active infection, hepatic disease, age, performance status). Many glioma studies attenuated effects after adjustment. Frontiers +1

Suggested methods wording (copy/paste)

"Preoperative PNI was calculated as 10×albumin(g/dL)+0.005×total lymphocyte count(/mm³). The optimal PNI cut-off for overall survival was derived from ROC analysis using the Youden index; we report the chosen threshold (PNI = XX), the area under the curve (AUC = Y.Y), sensitivity and specificity. For completeness we also present results using the cohort median (PNI = ZZ) and PNI as a continuous variable."