Prediction of lymph node metastasis in oral tongue squamous cell carcinoma using the neutrophil-to-lymphocyte ratio and platelet-to-neutrophil ratio

Bo Wang | Junwen Liu | Zhengrong Zhong

Abstract

Background: Lymph node metastasis in a variety of tumors is associated with systemic inflammatory markers. However, this association has not been reported in oral tongue squamous cell carcinoma (OTSCC). This study aimed to investigate how the preoperative neutrophil-to-lymphocyte ratio (NLR) and platelet-to-neutrophil ratio (PNR) in OTSCC patients correlated with the occurrence of OTSCC and lymph node metastasis.

Methods: The data of 73 patients with primary OTSCC who underwent surgical resection were retrospectively analyzed. Patients with other malignant tumors, patients who had received radiotherapy or chemotherapy before surgery, and patients with active inflammation were excluded. The enrolled patients were divided into groups N₀ (no early-stage lymph node metastasis) and N₁ (early-stage lymph node metastasis). Venous blood samples were collected before surgery and at the third week after surgery and subjected to complete blood counting in a blood analyzer. Eighty-seven healthy people were included as a control group. In addition, the NLR and PNR in OTSCC patients were compared with those in the controls, and the postoperative NLR and PNR of group N₀ were compared with those of group N₁.

Results: The NLR was significantly higher in the OTSCC patients than the controls (p < 0.05). The area under the receiver operating characteristic curve was 0.595. Further comparison of the NLR and PLR between group N₀ and group N₁ showed that when NLR was ≤1.622, and the probability of early-stage lymph node metastasis in OTSCC patients was 73.3%, and when PNR was >60.889, the probability was 86.7%. In re-examination 3 weeks postoperatively, the NLR and PNR were not significantly different between groups.

Conclusion: The NLR has certain reference value for the diagnosis of OTSCC. The preoperative NLR and PNR can be used to predict early-stage lymph node metastasis in patients with histopathologically confirmed OTSCC.

Keywords
neutrophil-to-lymphocyte ratio (NLR), oral tongue squamous cell carcinoma (OTSCC), platelet-to-neutrophil ratio (PNR)
INTRODUCTION

Oral tongue squamous cell carcinoma (OTSCC) is one of the most common oral malignancies, and its incidence has been rising in recent years. Because the tongue is rich in blood vessels and lymphatic vessels, it is prone to lymph node metastasis. Therefore, the prognosis of OTSCC is poor, and the 5-year survival rate is low. Although lymph node micrometastases sometimes cannot be detected by clinical examinations such as palpation, ultrasound, computed tomography (CT), and magnetic resonance imaging (MRI), they can be revealed by postoperative pathological examination. Such micrometastases are called occult lymph node metastases. Up to 33% of OTSCC cases may have occult lymph node metastasis that is currently undetectable by imaging techniques. In patients with OTSCC, the survival rate of patients with cervical lymph node metastasis is 50% lower than that of patients without cervical lymph node metastasis. Therefore, preoperative diagnosis of cervical lymph node metastasis in patients with OTSCC is difficult but important in clinical practice. In addition, patients without lymph node metastasis (N0) tend to suffer unnecessary complications such as intraoperative risk and shoulder joint dysfunction if they receive cervical lymph node dissection, whereas if patients with occult lymph node metastases do not undergo lymph node dissection, they tend to have a high risk of short-term local recurrence and poor prognosis.

The early diagnosis of OTSCC and risk assessment of lymph node metastasis still lack readily available and reliable molecular markers. In the past few years, the relationship between tumors and inflammation has become a research hotspot. Tumor cells promote systemic inflammatory responses and cause changes in white blood cell (WBC) and platelet (PLT) counts in peripheral blood. They mainly mediate these changes by releasing stress-related substances and proinflammatory cytokines, such as tumor necrosis factor α, which in turn can affect the occurrence and progression of tumors. Therefore, the neutrophil-to-lymphocyte ratio (NLR), the platelet-to-lymphocyte ratio (PLR), and the platelet-to-neutrophil ratio (PNR) of the peripheral blood obtained from a complete blood count have been used as indicators of systemic inflammation, and the diagnostic and prognostic value of NLR, PLR, and PNR for multiple tumors has been investigated. Current studies have found that NLR, PLR, and PNR have important reference value for the early diagnosis of lung cancer, liver cancer, breast cancer, and colorectal cancer as well as the prediction of lymph node metastasis. However, the relationships of NLR and PNR with the occurrence and development of OTSCC have not been reported.

This study aimed to evaluate the value of NLR and PNR in peripheral blood for the diagnosis of OTSCC and the prediction of cervical lymph node metastasis to provide a reference for the development of surgical plans for OTSCC patients.

MATERIALS AND METHODS

General information

A total of 73 patients with primary OTSCC who underwent surgery at the Department of Oral Head and Neck Oncology of the Ninth People’s Hospital affiliated to Shanghai JiaoTong University School of Medicine between January 2017 and December 2019 were enrolled. OTSCC was histopathologically confirmed in all patients. The exclusion criteria were as follows: (a) patients with other malignancies, (b) patients who had received radiotherapy or chemotherapy before surgery, (c) patients who could not be followed up, and (d) patients with active inflammation. At the same time, the information about age, sex, smoking history, drinking history, surgical records, pathological examination results, and tumor-node-metastasis (TNM) stage (according to the TNM classification of the American Joint Committee on Cancer (AJCC) in 2017) was collected. The OTSCC patients were divided into groups N0 (no early-stage lymph node metastasis) and N1 (early-stage lymph node metastasis). Early-stage refers to primary tumor in T1 stage (tumor ≤2 cm with depth of invasion DOI ≤5 mm) or T2 stage (tumor ≤2 cm, with DOI >5 mm and ≤10 mm; or tumor >2 cm and ≤4 cm, with DOI ≤10 mm). During the same period, 87 healthy individuals were included in the control group (group HC). This study was approved by the Ethics Committee of the Ninth People’s Hospital affiliated to Shanghai JiaoTong University School of Medicine, and all participants signed informed consent forms.

Complete blood count

Venous blood samples of each OTSCC patient were collected before surgery and 3 weeks after the surgery. Each sample was immediately mixed with K2-ethylenediaminetetraacetic acid (EDTA) in anticoagulation vacuum tubes. The sample was submitted to a complete blood count within 4 h of sample collection in a blood analyzer. Venous blood samples of patients in group HC were also collected for a complete blood count.

Statistical analysis

SPSS was used for statistical analysis. The independent-samples t test, the chi-squared test, and receiver operating characteristic (ROC) curve analysis were used for intergroup comparisons. p < 0.05 was considered statistically significant.

RESULTS

General information of patients

There was no significant difference in the general information between group N0 and group N1. The detailed results are shown in Table 1.

NLR level in OTSCC patients

The OTSCC patients had lower lymphocyte and PLT counts than the controls (p < 0.05), though their neutrophil counts were similar (p > 0.05). The PLR and PNR of OTSCC patients were not significantly different from those of the controls, while the NLR level
TABLE 1  General clinical information of group N₀ and group N₁

| Variable | HC group | OTSCC group | p  |
|----------|----------|-------------|----|
|          | (n = 87) | (n = 73)    |    |
| Age      | 50.9 ± 9.4 | 53.7 ± 17.3 | 0.303 |
| Gender   | Male 50 | 40          | 0.734 |
|          | Female 37 | 33         |    |
| Smoker   | Yes 14 | 20          | 0.082 |
|          | No 73 | 53         |    |
| Drinker  | Yes 5 | 7           | 0.358 |
|          | No 82 | 66         |    |

Note: Data are presented as numbers.

TABLE 2  Complete blood count parameters and calculated values of PLR, NLR, and PNR for groups N₀, N₁, and HC

| Groups | PLT (×10⁹/L) | LYMPH (×10⁹/L) | NEUT (×10⁹/L) | PLR | NLR | PNR |
|--------|-------------|----------------|---------------|-----|-----|-----|
|        |             |                |               |     |     |     |
| HC     | 87          | 226.44 ± 47.97 | 1.98 ± 0.42   | 3.30 ± 0.85 | 117.12 ± 30.32 | 1.72 ± 0.60 | 73.34 ± 22.07 |
| Pretreatment |       |                |               |     |     |     |
| N₀     | 43          | 210.23 ± 60.69 | 1.78 ± 0.60†  | 3.85 ± 1.29†  | 129.59 ± 48.66 | 2.33 ± 0.80†  | 58.69 ± 21.79†  |
| N₁     | 30          | 206.90 ± 67.79 | 1.84 ± 0.59†  | 2.84 ± 1.71†  | 122.31 ± 57.20 | 1.62 ± 1.00†  | 87.27 ± 38.26†  |
| N₀ + N₁ | 73          | 208.86 ± 63.26† | 1.80 ± 0.60†  | 3.44 ± 1.55   | 126.60 ± 52.08 | 2.03 ± 0.95†  | 70.43 ± 32.66   |
| Three month after treatment |       |                |               |     |     |     |
| N₀     | 25          | 229.68 ± 152.98 | 1.50 ± 1.38   | 8.08 ± 4.01   | 193.88 ± 134.64 | 6.73 ± 4.52   | 33.70 ± 20.04   |
| N₁     | 26          | 217.23 ± 69.69 | 1.2 ± 0.41    | 6.63 ± 3.24   | 202.65 ± 98.62 | 6.18 ± 4.23   | 40.77 ± 20.43   |

Note: Data are presented as numbers, and the data were calculated by different etiologies.

Abbreviations: NLR, neutrophil-to-lymphocyte ratio; PLR, platelet-to-lymphocyte ratio; PNR, platelet-to-neutrophil ratio.

†Compared with group HC, p < 0.05.
‡Compared with group HC, p < 0.01.
§Compared with group N₀, p < 0.01.
¶Compared with group N₁, p < 0.01.

4 | DISCUSSION

There is a controversy regarding the treatment of cervical lymph nodes in N₀ patients at the early stage of oral squamous cell carcinoma (OSCC). Some researchers advocate a wait-and-see strategy, which suggests that lymph node dissection should be performed only after confirmation of lymph node metastasis. However, this treatment regimen significantly reduces the survival rate of patients. Extended resection of the primary tumor(s) combined with cervical lymph node dissection has a high cure rate. Due to the extended scope of surgery, this may lead to early functional impairment or loss and thus affect the quality of life of the patients. Therefore, how to effectively assess noninvasive treatment for cervical lymph node–negative patients with enlarged cervical lymph nodes or a high risk for metastasis has been the focus of clinical studies.

This study first analyzed the NLR levels in the OTSCC group. Compared with group HC, the OTSCC group had a significantly higher NLR, mainly due to the lower absolute lymphocyte count and the (nonsignificantly) higher average absolute neutrophil count in the OTSCC group. According to the absence or presence of lymph node metastasis, the OTSCC group was divided into groups N₀ and N₁. This analysis showed that the absolute neutrophil count in group N₁ was significantly lower than that in group N₀ and that the higher PNR (p = 0.0001) (Table 2). Changes in NLR and PNR in groups N₀ and N₁ were analyzed using ROC curves. The NLR had the AUC of 0.769, the 95% CI of 0.649–0.889, the cutoff value of 1.6223, and the sensitivity of 73.3% in predicting early-stage lymph node metastasis. The PNR had the AUC of 0.762, the 95% CI of 0.649–0.875, the cutoff value of 60.8892, and the sensitivity of 86.7% in predicting early-stage lymph node metastasis (Figure 2, Table 3). However, at the third postoperative week, there was no significant difference in the NLR or PNR between groups N₀ and N₁ (Table 2).

3.3 | Evaluation of the predictive value of NLR and PNR for early-stage lymph node metastasis in patients with histopathologically confirmed OTSCC

Compared with group HC, group N₀ had a higher NLR (p = 0.0001) and lower PNR (p = 0.0001), and group N₁ had a higher PNR (p = 0.017). Compared with group N₀, group N₁ had a lower NLR (p = 0.001) and
absolute lymphocyte count and the absolute PLT count in group $N_1$ were not significantly different from those in group $N_0$. As a result, compared with group $N_0$, group $N_1$ had a significantly lower NLR, a significantly higher PNR, and a nonsignificantly different PLR. The causes for the differences between groups $N_0$ and $N_1$ might be that when the OTSCC did not metastasize (group $N_0$), the innate immunity was dominant, so the neutrophil count was elevated. Our results did show that the absolute neutrophil count in group $N_0$ was significantly higher than that in group HC.$^{16}$ After tumor metastasis (group $N_1$), the innate immunity was weakened, adaptive immunity began to play a dominant role, the neutrophil count dropped significantly lower than that before tumor metastasis (group $N_0$), and the mean lymphocyte count was (nonsignificantly) larger than that in group $N_0$, resulting in the lower NLR and the higher PNR in group $N_1$. $^{17,18}$

Further analysis of groups $N_0$ and $N_1$ showed that the NLR and PNR had a sensitivity of 73.3% and 86.7%, respectively, in predicting early-stage lymph node metastasis in patients with histopathologically confirmed OTSCC. Among OTSCC patients without lymph node metastasis, only 16.3% and 39.5% had "positive" NLR and PNR values, respectively. Therefore, NLR and PNR have good predictive value for early-stage lymph node metastasis in OTSCC patients.

No previous studies have reported a relationship between lymph node metastasis in tongue cancer and the PNR. Xian et al. $^{19}$ studied 174 patients with solid carcinoma-related cerebral venous sinus thrombosis and found that the higher PNR, the higher the probability of intracranial metastasis in these patients. This finding is similar to our finding that when PNR was $>60.889$, OTSCC patients were more prone to early-stage lymph node metastasis, their innate immunity was weakened, their neutrophil count was decreased, and their PLT count was unchanged, resulting in an elevated PNR. Wu et al. $^{20}$ found that a high NLR ($\geq 2.95$) was significantly correlated with occult lymph node metastasis in OSCC. That result is inconsistent with ours. The possible cause of the discrepancy is that although OTSCC is an OSCC, it has its own characteristics, as a special OSCC with unique characteristics. This suggests that future studies on OSCC should refine their results by subtype and that the general conclusions drawn about OSCC are not necessarily applicable to all subtypes.

The NLR and PNR can be easily determined through a standardized, inexpensive process. One limitation of this study is that other systemic diseases may affect the NLR and PNR because they reflect the inflammatory state of the body. Second, this study was a retrospective single-center study, so future multicenter studies are needed to verify our results. Third, the absolute counts of

### Table 3: Assessment of early-stage lymph node metastasis in OTSCC patients using the NLR and PNR

| Samples      | NLR | PNR |
|--------------|-----|-----|
|              | No. of cases | No. of positive | Positive rate (%) | No. of positive | Positive rate (%) |
| HC group     | 87  | 43  | 49.4 | 30  | 34.5 |
| OTSCC group  | 73  | 29  | 39.7 | 43  | 58.9 |
| $N_0$ group  | 43  | 7   | 16.3 | 17  | 39.5 |
| $N_1$ group  | 30  | 22  | 73.3 | 26  | 86.7 |

Note: Data are presented as numbers and percentages, and the data were calculated by different etiologies.
circulating neutrophils and lymphocytes were nonspecific, so there was no detailed information about the relationships of WBC subsets or activation status with OTSCC, such as whether a low activity of neutrophils or a high CD4/CD8 ratio is related to the progression of OTSCC. Nevertheless, our results strongly suggest that the preoperative NLR and PNR can be used as indicators to predict early-stage lymph node metastasis in patients with histopathologically confirmed OTSCC.

ACKNOWLEDGMENTS
The work was funded by the Natural Science Foundation of Anhui Province (No 1308085MH124) and the Natural Science Research of Educational Department Anhui Province (No. KJ2015ZD31).

CONFLICT OF INTEREST
The authors declare that they have no competing interests.

DATA AVAILABILITY STATEMENT
The raw/processed data required to reproduce these findings cannot be shared at this time as the data also forms part of an ongoing study.

ORCID
Bo Wang https://orcid.org/0000-0003-2693-0583

REFERENCES
1. Omori H, Nishio M, Masuda M, et al. YAP1 is a potent driver of the onset and progression of oral squamous cell carcinoma. Sci Adv. 2020;6(12):eaay3324.
2. Attaran N, Gu X, Coates PJ, et al. Downregulation of TAP1 in tumor-free tongue contralateral to squamous cell carcinoma of the oral tongue, an indicator of better survival. Int J Mol Sci. 2020;21(17):E6220.
3. Abbate V, Dell’Aversana Orabona G, Salzano G, et al. Pre-treatment neutrophil-to-lymphocyte ratio as a predictor for occult cervical metastasis in early stage (T1-T2 cN0) squamous cell carcinoma of the oral tongue. Surg Oncol. 2018;27(3):503-507.
4. Jiang Q, Tang A, Long S, et al. Development and validation of a nomogram to predict the risk of occult cervical lymph node metastases in cN0 squamous cell carcinoma of the tongue. Br J Oral Maxillofac Surg. 2019;57(10):1092-1097.
5. De Zinis LO, Bolzoni A, Piazza C, Nicolai P. Prevalence and localization of nodal metastases in squamous cell carcinoma of the oral cavity: role and extension of neck dissection. Eur Arch Otorhinolaryngol. 2006;263(12):1131-1135.
6. Lim SC, Zhang S, Ishii G, et al. Predictive markers for late cervical metastasis in stage I and II invasive squamous cell carcinoma of the oral tongue. Clin Cancer Res. 2004;10(1 Pt 1):166-172.
7. Dogan E, Cetinayak HO, Saraloglu S, Erdag TK, Ikiz AO. Patterns of cervical lymph node metastases in oral tongue squamous cell carcinoma: implications for elective and therapeutic neck dissection. J Laryngol Otol. 2014;128(3):268-273.
8. Itami Y, Miyake M, Tatsumi Y, et al. Preoperative predictive factors focused on inflammation-, nutrition-, and muscle-status in patients with upper urinary tract urothelial carcinoma undergoing nephroureterectomy. Int J Clin Oncol. 2019;24(5):533-545.
9. Balkwill F, Joffroy C. TNF: a tumor-suppressing factor or a tumor-promoting factor? Future Oncol. 2010;6(12):1833-1836.
10. Wang X, Cao L, Li S, Wang F, Huang D, Jiang R. Combination of PD-L1 expression and NLR as prognostic marker in patients with surgically resected non-small cell lung cancer. J Cancer. 2019;10(26):6703-6710.
11. Liu X, He L, Han J, et al. Association of neutrophil-lymphocyte ratio and T lymphocytes with the pathogenesis and progression of HBV-associated primary liver cancer. PLoS One. 2017;12(2):e0170605.
12. Moon G, Noh H, Cho IJ, Lee JI, Han A. Prediction of late recurrence in patients with breast cancer: elevated neutrophil to lymphocyte ratio (NLR) at 5 years after diagnosis and late recurrence. Breast Cancer. 2020;27(1):51-61.
13. Stojkovic Lalosevic M, Pavlovic Markovic A, Stankovic S, et al. Combined diagnostic efficacy of neutrophil-to-lymphocyte ratio (NLR), platelet-to-lymphocyte ratio (PLR), and mean platelet volume (MPV) as biomarkers of systemic inflammation in the diagnosis of colorectal cancer. Dis Markers. 2019;2019:6036979.
14. Lawrance S, Bui C, Mahindra V, Cooke-Yarborough C. Assessing a modified-AJCC TNM staging system in the New South Wales Cancer Registry, Australia. BMC Cancer. 2019;19(1):850.
15. Schmitz S, Machiels JP, Weynand B, Gregoire V, Hamoir M. Results of selective neck dissection in the primary management of head and neck squamous cell carcinoma. Eur Arch Otorhinolaryngol. 2009;266(3):437-443.
16. Magalhaes MA, Glogauer JE, Glogauer M. Neutrophils and oral squamous cell carcinoma: lessons learned and future directions. J Leukoc Biol. 2014;96(5):695-702.
17. Caldeira PC, Vieira ÉLM, Sousa AA, Teixeira AL, Aguilar MCF. Immunophenotype of neutrophils in oral squamous cell carcinoma patients. J Oral Pathol Med. 2017;46(9):703-709.
18. Quan H, Fang L, Pan H, et al. An adaptive immune response driven by mature, antigen-experienced T and B cells within the microenvironment of oral squamous cell carcinoma. Int J Cancer. 2016;138(12):2952-2962.
19. Xian Z, Chen Y, Chen L, et al. A clinical research on the potential pathogenesis of somatic cancer related cerebral venous sinus thrombosis. Medicine (Baltimore). 2019;98(19):e15134.
20. Wu CN, Chuang HC, Lin YT, Fang FM, Li SH, Chien CY. Prognosis of neutrophil-to-lymphocyte ratio in clinical early-stage tongue (cT1/ T2N0) cancer. Onco Targets Ther. 2017;10:3917-3924.

How to cite this article: Wang B, Liu J, Zhong Z. Prediction of lymph node metastasis in oral tongue squamous cell carcinoma using the neutrophil-to-lymphocyte ratio and platelet-to-neutrophil ratio. J Clin Lab Anal. 2021;35:e23684. https://doi.org/10.1002/jcla.23684