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***Retrospective Study***

**Prognostic significance of preoperative lymphocyte to monocyte ratio in patients with signet ring gastric cancer**

Liu HL *et al*. Preoperative LMR in SRCC GC

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**Abstract**

BACKGROUND

The ratio of lymphocytes to monocytes (LMR) has been shown to be an effective predictor of gastric cancer prognosis. However, its predictive accuracy for signet ring gastric cancer is currently not well understood.

AIM

To evaluate the prognosis predictive accuracy of preoperative LMR in signet ring gastric cancer.

METHODS

A total of 212 signet ring gastric cancer patients admitted at the Xiangya Hospital of Central South University, Department of Gastrointestinal Surgery, from January 2012 to December 2016 were enrolled in the study. The prognosis predictive accuracy of preoperative LMR was explored based on the area under the receiver operating characteristic. Factors that significantly affect the survival of patients were identified using single factor analysis, and those that were independently associated with signet ring gastric cancer were identified through multivariate analysis.

RESULTS

The results of the single factor analysis revealed a strong correlation between the survival of signet ring gastric cancer patients and several factors, including tumor invasion (*χ*2 = 49.726; *P* < 0.001), lymph node metastasis (*χ*2 = 30.269; *P* < 0.001), pTNM stage (*χ*2 = 49.322; *P* < 0.001), surgical approach (*χ*2 = 8.489; *P* = 0.004), age (*t* = -2.213; *P* < 0.028), carcinoembryonic antigen (CEA) (Z = -3.265; *P* = 0.001), platelet-to-lymphocyte ratio (Z = -2.196; *P* = 0.028), LMR (Z = -2.226; *P* = 0.026), ALB (t = 3.284; *P* = 0.001), prognostic nutritional index (t = -3.789; *P* < 0.001) and FIB (Z = -3.065; *P* = 0.002). Furthermore, the multivariate analysis further demonstrated that age (HR: 0.563, 95%CI: 0.363-0.873), tumor invasion depth (HR: 0.226, 95%CI: 0.098-0.520), pTNM stage (HR: 0.444, 95%CI: 0.255-0.771), preoperative CEA level (HR: 0.597, 95%CI: 0.386-8.790), and preoperative LMR level (HR: 1.776, 95%CI: 1.150-2.741) were independent factors influencing the prognosis of signet ring gastric cancer.

CONCLUSION

In signet ring gastric cancer patients, a low preoperative LMR level predicts poor prognosis. The death risk ratio of the low LMR group compared to the high LMR group is 1.776.

**Key Words:** Gastric cancer; Signet ring cell carcinoma; Inflammation indexes; Coagulation indexes; Prognosis.

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**Core Tip:** Low preoperative lymphocytes to monocytes levels -predict poor prognosis of patients with signet ring gastric cancer, making it a valuable prognostic factor.

**INTRODUCTION**

Signet ring gastric cancer is a type of stomach cancer that is characterized by presence of cells filled with mucus, which push the nucleus to one side of the cell, giving it a ring-like appearance. This type of cancer is highly invasive, progresses rapidly, and has a high degree of malignancy. Although the incidence of gastric cancer has decreased in recent decades, cases of signet-ring cell carcinoma (SRCC) are increasingly being reported. Studies have demonstrated that SRCC accounts for 35 % to 45 % of all cases of gastric adenocarcinoma[1], and its incidence increased by tenfold from 1970 to 2000[2].

Currently, the prognosis of SRCC is not well understood. Given that SRCC is prone to lymph node and peritoneal metastasis, less responsive to chemotherapy, and most patients are diagnosed at an advanced cancer stage, patients with SRCC have a poor prognosis[2].

The occurrence and development of tumors are driven by several factors including inflammatory immune response of the host[3]. Numerous studies have explored the relationship between different inflammatory markers, chemotherapeutic effects, and prognosis in gastric cancer. Among the most easily available inflammatory markers obtained from the whole blood cell count are lymphocyte-to-monocyte ratio (LMR)[4], neutrophil-to-lymphocyte ratio (NLR)[5], platelet-to-lymphocyte ratio (PLR)[6], and systemic immune inflammation (SII)[7]. The prognostic nutritional index (PNI), a simple and easy detection index, has been widely used in clinical practice and shown to be associated with the prognosis of malignant gastric tumors[8,9]. Moreover, the development of tumors is accompanied by changes in the blood coagulation dynamics of the host[10]. Coagulation factor levels, such as platelet count, international standard ratio, fibrin degradation products, fibrinogen, and D-dimer levels, have been associated with tumor stage, metastasis, chemotherapeutic effect, and prognosis of patients with solid tumors[11,12]. Although many scientists have explored the relationship between various indicators and chemotherapy response and prognosis of gastric cancer, few studies have explored the prognostic value of these indicators in SRCC.

Against this background, we explored the relationship between the common inflammatory indicators, nutritional indicators, coagulation indicators and the prognosis of SRCC to identify prognostic predictors of SRCC.

**MATERIALS AND METHODS**

***Patients***

The retrospective study included 212 patients with gastric cancer admitted to the Department of Gastroenterology at Xiangya Hospital of Central South University from January 2012 to December 2016. To be included in the study, patients had to meet the following criteria: (1) Postoperative pathology revealed SRCC components greater than 10%; (2) Accepted to undergo radical gastrectomy; and (3) With complete clinical and follow-up data. Moreover, the exclusion criteria are as follows: (1) Preoperative radiotherapy, chemotherapy, targeted therapy, immunotherapy, and other anti-tumor treatments that may affect the patient's blood routine, liver and kidney function, and coagulation routine; (2) Preoperative examination indicated the presence of distant metastases such as liver, lung, and bone metastases; (3) Intraoperative detection of metastasis; (4) Comorbidity hematological diseases and other systemic malignancies; (5) Combined with severe infections, liver disease, kidney disease, and autoimmune diseases; (6) Emergency surgery due to perforation and bleeding of gastric cancer; and (7) Gastric stump cancer. This study was approved by the Ethics Committee of Xiangya Hospital of Central South University.

***Measurement of variables***

Patient-related results, including demographic data, clinic characteristics, and biochemical test results such as carcinoembryonic antigen (CEA), LMR [= lymphocyte count (× 109/L)/monocyte count (× 109/L)], NLR [= Neutrophil count (× 109/L)/Lymphocyte count (× 109/L)], PLR [= platelet count (× 109/L)/Lymphocyte coun (× 109/L)], SII [= Neutrophil count (× 109/L) × Platelet count (× 109/L)/Lymphocyte count (× 109/L)][13], coagulation index [activated partial thromboplastin time (APTT), fibrinogen degradation product (FDP), fibrinogen (FIB), prothrombin time (PT), thrombin time (TT), and international normalized ratio (INR)] were obtained from the medical database of the hospital. Additionally, albumin (ALB), globulin (GLB), albumin to globulin ratio [AGR = serum albumin (g/L)/serum globulin (g/L)], and prognostic nutritional index [PNI=5 × Lymphocyte count (× 109/L)+serum albumin (g/L)][14] were obtained through peripheral complete blood count and blood biochemistry before the surgery. Cut-off values for each variable were obtained from the receiver operating characteristic (ROC) curves.

***Statistical analysis***

Statistical analysis was performed using SPSS 25.0 and GraphPad Prism 8.0. The normality of the data was assessed using the Shapiro-Wilk test, and the data which did not fit the normal distribution was represented by M (P25, P75) and analyzed using the Mann-Whitney *U* test. Single factor analysis was performed using the independent-sample *t*-test for normally distributed data, and the counts were presented as percentages (%). The prognostic value of inflammation, blood coagulation and other indicators was evaluated using ROC curves, and the optimal cut-off point of patient survival was determined. The patients were then divided into high and low groups based on the medium levels of the above indicators. Kaplan-Meier survival analysis and the Log-rank tests were used to compare survival rates between the high and low-risk groups. Cox proportional hazards regression analysis was conducted to identify independent predictors of the prognosis of gastric cancer with SRCC by including indicators that significantly affected the survival status. Statistical significance was set at *P* < 0.05.

**RESULTS**

***Baseline data and pathological characteristics***

The study included 212 patients with SRCC, of whom 87 patients (41.04 %) died, and 125 patients (58.96 %) survived for the 5-year follow-up period. The mean age of the patients was 51.42 ± 11.27 years, 117 were males (55.19 %), and 95 were females (44.81 %). The tumor location was in the upper, middle and lower third of the stomach in 5 (2.36 %), 58 (27.36 %) and 149 (70.28 %) cases, respectively. Distal gastrectomy was performed in 167 (78.78 %) cases, while total gastrectomy was performed in 45 (21.22 %) cases. The tumor infiltration depth was pT1 or pT2 in 79 cases (37.26 %) and pT3 or pT4 in 133 cases (62.74 %). Lymph node metastasis was found in 118 (55.66 %) cases with pN1, pN2 or pN3, while 94 (44.34 %) cases were classified as N0. The pTNM stage was I or II in 117 cases (55.19 %) and III in 95 (44.81 %) cases (Table 1).

***ROC analysis of predictors***

Table 2 shows the cut-off value and area under the curve (AUC) for each index, and Figure 1 shows the ROC curves (AUC, 95%CI) of the predictors, including age, CEA, PLR, LMR, ALB, PNI and FIB. Overall, the results indicate good predictive values of CEA (0.632, 0.556-0.708), LMR (0.590, 0.512-0.669), ALB (0.618, 0.540-0.696), PNI (0.644, 0.567-0.721), and FIB (0.624, 0.547-0.701),

***Single factor analysis for prognostic factors***

Table 3 displays the results of the pathological features and the 5-year survival rate of the patients. The analysis revealed that depth of tumor invasion, lymph node metastasis, pTNM stage and resection range were associated with the survival rate, while gender and tumor location showed no correlation.

As shown in Table 4, firstly, the single factor analysis demonstrated that younger patients had a better survival rate compared with older patients. The survival rate was also significantly better in the lower CEA level group compared with the higher CEA level group. In addition, SRCC patients in the higher PLR value had poorer survival rates than those in the lower PLR value group, while SRCC patients in the higher LMR value group had a better survival rate than those in the lower LMR value group. Finally, the survival rate of patients with lower FIB was significantly better in those with higher FIB, whereas patients with lower ALB had a significantly lower survival rate compared with higher ALB group.

***Grouping of predictors***

The predictors, including age, CEA, ALB, PNI, LMR, PLR, and FIB, were grouped as shown in Table 5.

***Kaplan-Meier survival analysis***

The Kaplan-Meier survival curves are shown in Figure 2. The Log-rank test indicated no significant difference in survival rate between the low PLR group and the high PLR group (*P* = 0.147). In contrast, significant differences were observed in survival rate between different resection groups (*P* < 0.001), depth of invasion (*P* < 0.001), lymph node metastasis (*P* < 0.001), pTNM stage (*P* < 0.001), age (*P* = 0.004), LMR (*P* = 0.003), ALB (*P* = 0.008), PNI (*P* = 0.002) and FIB (*P* = 0.001).

***Multivariate and multiple-factor analysis***

To explore the independent factors affecting the prognosis of SRCC, indicators that demonstrated statistical differences by the Log-rank test were included in a Cox proportional hazards regression model for multivariate analysis. The results (HR, 95%CI) are presented in Figure 3. The independent factors for SRCC prognosis were age (0.563, 0.363-0.873, *P* = 0.010), depth of tumor invasion (0.226, 0.098-0.520), pTNM stage (0.444, 0.255-0.771), preoperative CEA level (0.597, 0.386-8.790), and preoperative LMR level (1.776, 1.150-2.741). Advanced age, high CEA level before surgery, low LMR level before surgery, deep tumor invasion, and late pTNM stage were all indicative of a relatively poor prognosis. Specifically, the risk of death in low LMR group before surgery was 1.776 times higher than that of the high LMR group.

**DISCUSSION**

Currently, surgery is the mainstay treatment for gastric cancer patients, especially those with SRCC. However, despite radical resection or adjuvant chemotherapy, the prognosis of SRCC patients, particularly those in advanced stages, is not optimistic. Therefore, it is crucial to elucidate the mechanism of tumor progression and identify independent prognostic factors to evaluate the overall condition of tumor patients and optimize diagnosis and treatment.

The correlation between inflammation and tumors was first proposed by Rudolf Virchow[14]. Research has shown that inflammation participates in tumor development[15,16]. Furthermore, inflammation can influence the prognosis of tumors by altering immune response[17,18].

Lymphocytes and monocytes play a crucial role in anti-tumor immune response[19]. The relationship between LMR and the prognosis of malignant tumors has been widely reported[20-22]. However, few studies have investigated the relationship between inflammatory markers and SRCC. Chengcheng Tong *et al*[23], reported that derived monocyte-to-lymphocyte ratio (dMLR) could independently predict lymph node metastasis of SRCC. Zhu *et al*[9] reported the relationship between SII and the prognosis of SRCC, but the relationship between LMR and SRCC has not been investigated.

Numerous studies have shown that high levels of peripheral blood lymphocyte count and TILs are associated with a good prognosis in gastric cancer[24-26]. Li *et al*[27] reported that patients with smaller tumors (< 5 cm) had higher counts of peripheral blood CD4 + T cells (*P* = 0.003) and CD8 + T cells (*P* = 0.002). In addition, patients with well-differentiated gastric cancer showed higher counts of CD4 + T cells (*P* = 0.029).

NK cells, which possess potent anti-tumor, anti-viral and antibacterial activity, are crucial in activating and regulating adaptive immune responses. In human peripheral blood, NK cells account for approximately 3%-5% of lymphocytes[28,29]. The anti-tumor activity of NK cells is mainly determined by a group of inhibitory and activating receptors[30]. Patients with gastric cancer exhibit lower expression of activating receptors such as NKG2D, NKp30, and NKp46, but higher PD-1 expression. Moreover, NK cells of patients with gastric cancer secrete lower cytokines (IFN-γ, IL-2, TNF-α, IL-12) and impaired ability to release perforin and granzyme. Meanwhile, gastric cancer cells express little MICA/B, ULBP, and B7H6, to evade NK cell-mediated innate immunity. Gastric cancer cells can also produce cytokines such as IL-10, TGF-β, and PGE2, which recruit MDSC and Treg cells to suppress NK cell function[31]. The proportion of apoptotic NK cells in patients with gastric cancer is elevated when receiving gastrectomy[32]. Collectively, these lines of evidence show that the number and function of NK cells decrease sharply with the progression of gastric cancer[31].

Monocytes, particularly those that differentiate into tumor-associated macrophages (TAMs), contribute to the development of gastric cancer[33]. M1 TAMs have anti-tumor effects, while M2 TAMs promote tumor growth[34]. Under the influence of cytokines and extracellular matrix secreted by tumour cells and lymphocytes, M1 TAMs can convert into M2 TAMs. In gastric cancer, M2 TAMs are highly expressed in SRCC, mucinous adenocarcinoma and diffuse gastric cancer[35].

Our study found that the 5-year survival rate of patients with low LMR before surgery was significantly lower than that with high LMR. In summary LMR affects the prognosis of SRCC due to reduction of lymphocytes during inflammatory response and increase of tumor-associated macrophages produced by circulating monocytes. However, this study has some limitations. First, this was a retrospective single-center study and no external validation was performed. In addition, studies have suggested that gastric cancer with different SRCC ratios may have varying biological characteristics, although we demonstrated a relationship between LMR and the prognosis of SRCC. Therefore, it is imperative to explore further the predictive value of various indicators in gastric cancer with different SRCC ratios.

**CONCLUSION**

In summary, this study shows that a low preoperative LMR level indicates a poor prognosis of signet ring gastric cancer. Particularly, compared with the high LMR group, the risk of mortality in the low LMR group is 1.776.

**ARTICLE HIGHLIGHTS**

***Research background***

The incidence of signet ring gastric carcinoma has increased among the past decades. Several inflammation indexes, including ratio of lymphocytes to monocytes (LMR), have been shown to be effective predictors of gastric cancer prognosis.

***Research motivation***

The predictive accuracy of ratio of LMR for signet ring gastric cancer is unclear now.

***Research objectives***

To assess the prognosis predictive accuracy of preoperative LMR for signet ring gastric cancer.

***Research methods***

Our research center conducted a retrospective analysis of clinical data from patients diagnosed with signet ring gastric carcinoma over the past 5 years, identifying factors that significantly affect patients’ survival by using single factor analysis, and deciding independent prognostic factors related to signet ring cell gastric cancer by using multivariate analysis.

***Research results***

The results of the single factor analysis indicated a strong correlation between the survival of signet ring gastric cancer patients and several factors, including tumour invasion, lymph node metastasis, pTNM stage, surgical approach, age, carcinoembryonic antigen (CEA), platelet-to-lymphocyte ratio (PLR), LMR, ALB, PNI and FIB. Furthermore, the multivariate analysis revealed that age, tumor invasion depth, pTNM stage, preoperative CEA level, and preoperative LMR level were independent factors related to the prognosis of signet ring gastric cancer.

***Research conclusions***

In signet ring gastric cancer patients, a low preoperative LMR level is indicative of a poor prognosis. The death risk ratio of the low LMR group compared to the high LMR group is 1.776.

***Research perspectives***

The study subjects were followed up for 5 years and divided into survival group and death group. Clinical data, pathological data, and prognosis of the two groups of patients were observed.

**REFERENCES**

1 **Bamboat ZM**, Tang LH, Vinuela E, Kuk D, Gonen M, Shah MA, Brennan MF, Coit DG, Strong VE. Stage-stratified prognosis of signet ring cell histology in patients undergoing curative resection for gastric adenocarcinoma. *Ann Surg Oncol* 2014; **21**: 1678-1685 [PMID: 24394986 DOI: 10.1245/s10434-013-3466-8]

2 **Henson DE**, Dittus C, Younes M, Nguyen H, Albores-Saavedra J. Differential trends in the intestinal and diffuse types of gastric carcinoma in the United States, 1973-2000: increase in the signet ring cell type. *Arch Pathol Lab Med* 2004; **128**: 765-770 [PMID: 15214826 DOI: 10.5858/2004-128-765-DTITIA]

3 **Singh N**, Baby D, Rajguru JP, Patil PB, Thakkannavar SS, Pujari VB. Inflammation and cancer. *Ann Afr Med* 2019; **18**: 121-126 [PMID: 31417011 DOI: 10.4103/aam.aam\_56\_18]

4 **Pan YC**, Jia ZF, Cao DH, Wu YH, Jiang J, Wen SM, Zhao D, Zhang SL, Cao XY. Preoperative lymphocyte-to-monocyte ratio (LMR) could independently predict overall survival of resectable gastric cancer patients. *Medicine (Baltimore)* 2018; **97**: e13896 [PMID: 30593200 DOI: 10.1097/MD.0000000000013896]

5 **Miyamoto R**, Inagawa S, Sano N, Tadano S, Adachi S, Yamamoto M. The neutrophil-to-lymphocyte ratio (NLR) predicts short-term and long-term outcomes in gastric cancer patients. *Eur J Surg Oncol* 2018; **44**: 607-612 [PMID: 29478743 DOI: 10.1016/j.ejso.2018.02.003]

6 **Zhang X**, Zhao W, Yu Y, Qi X, Song L, Zhang C, Li G, Yang L. Clinicopathological and prognostic significance of platelet-lymphocyte ratio (PLR) in gastric cancer: an updated meta-analysis. *World J Surg Oncol* 2020; **18**: 191 [PMID: 32731872 DOI: 10.1186/s12957-020-01952-2]

7 **Cao X**, Xue J, Yang H, Han X, Zu G. Association of Clinical Parameters and Prognosis with the Pretreatment Systemic Immune-inflammation Index (SII) in Patients with Gastric Cancer. *J Coll Physicians Surg Pak* 2021; **31**: 83-88 [PMID: 33546540 DOI: 10.29271/jcpsp.2021.01.83]

8 **Ma LX**, Taylor K, Espin-Garcia O, Anconina R, Suzuki C, Allen MJ, Honorio M, Bach Y, Allison F, Chen EX, Brar S, Swallow CJ, Yeung J, Darling GE, Wong R, Kalimuthu SN, Jang RW, Veit-Haibach P, Elimova E. Prognostic significance of nutritional markers in metastatic gastric and esophageal adenocarcinoma. *Cancer Med* 2021; **10**: 199-207 [PMID: 33295697 DOI: 10.1002/cam4.3604]

9 **Zhu Y**, Fan L, Geng X, Li J. The predictive value of the prognostic nutritional index to postoperative prognosis and nursing intervention measures for colorectal cancer. *Am J Transl Res* 2021; **13**: 14096-14101 [PMID: 35035753]

10 **Gil-Bernabé AM**, Lucotti S, Muschel RJ. Coagulation and metastasis: what does the experimental literature tell us? *Br J Haematol* 2013; **162**: 433-441 [PMID: 23691951 DOI: 10.1111/bjh.12381]

11 **Kanda M**, Tanaka C, Kobayashi D, Mizuno A, Tanaka Y, Takami H, Iwata N, Hayashi M, Niwa Y, Yamada S, Fujii T, Sugimoto H, Murotani K, Fujiwara M, Kodera Y. Proposal of the Coagulation Score as a Predictor for Short-Term and Long-Term Outcomes of Patients with Resectable Gastric Cancer. *Ann Surg Oncol* 2017; **24**: 502-509 [PMID: 27600621 DOI: 10.1245/s10434-016-5544-1]

12 **Jomrich G**, Paireder M, Kristo I, Baierl A, Ilhan-Mutlu A, Preusser M, Asari R, Schoppmann SF. High Systemic Immune-Inflammation Index is an Adverse Prognostic Factor for Patients With Gastroesophageal Adenocarcinoma. *Ann Surg* 2021; **273**: 532-541 [PMID: 31425286 DOI: 10.1097/SLA.0000000000003370]

13 **Onodera T**, Goseki N, Kosaki G. [Prognostic nutritional index in gastrointestinal surgery of malnourished cancer patients]. *Nihon Geka Gakkai Zasshi* 1984; **85**: 1001-1005 [PMID: 6438478]

14 **Balkwill F**, Mantovani A. Inflammation and cancer: back to Virchow? *Lancet* 2001; **357**: 539-545 [PMID: 11229684 DOI: 10.1016/S0140-6736(00)04046-0]

15 **Mantovani A**. Cancer: inflammation by remote control. *Nature* 2005; **435**: 752-753 [PMID: 15944689 DOI: 10.1038/435752a]

16 **Hanahan D**, Weinberg RA. Hallmarks of cancer: the next generation. *Cell* 2011; **144**: 646-674 [PMID: 21376230 DOI: 10.1016/j.cell.2011.02.013]

17 **Iyengar NM**, Hudis CA, Dannenberg AJ. Obesity and inflammation: new insights into breast cancer development and progression. *Am Soc Clin Oncol Educ Book* 2013; **33**: 46-51 [PMID: 23714453 DOI: 10.14694/EdBook\_AM.2013.33.46]

18 **Moore MM**, Chua W, Charles KA, Clarke SJ. Inflammation and cancer: causes and consequences. *Clin Pharmacol Ther* 2010; **87**: 504-508 [PMID: 20147899 DOI: 10.1038/clpt.2009.254]

19 **Goeppert B**, Frauenschuh L, Zucknick M, Stenzinger A, Andrulis M, Klauschen F, Joehrens K, Warth A, Renner M, Mehrabi A, Hafezi M, Thelen A, Schirmacher P, Weichert W. Prognostic impact of tumour-infiltrating immune cells on biliary tract cancer. *Br J Cancer* 2013; **109**: 2665-2674 [PMID: 24136146 DOI: 10.1038/bjc.2013.610]

20 **Mandaliya H**, Jones M, Oldmeadow C, Nordman II. Prognostic biomarkers in stage IV non-small cell lung cancer (NSCLC): neutrophil to lymphocyte ratio (NLR), lymphocyte to monocyte ratio (LMR), platelet to lymphocyte ratio (PLR) and advanced lung cancer inflammation index (ALI). *Transl Lung Cancer Res* 2019; **8**: 886-894 [PMID: 32010567 DOI: 10.21037/tlcr.2019.11.16]

21 **Trinh H**, Dzul SP, Hyder J, Jang H, Kim S, Flowers J, Vaishampayan N, Chen J, Winer I, Miller S. Prognostic value of changes in neutrophil-to-lymphocyte ratio (NLR), platelet-to-lymphocyte ratio (PLR) and lymphocyte-to-monocyte ratio (LMR) for patients with cervical cancer undergoing definitive chemoradiotherapy (dCRT). *Clin Chim Acta* 2020; **510**: 711-716 [PMID: 32919942 DOI: 10.1016/j.cca.2020.09.008]

22 **Li C**, Zhang H, Li S, Zhang D, Li J, Dionigi G, Liang N, Sun H. Prognostic Impact of Inflammatory Markers PLR, LMR, PDW, MPV in Medullary Thyroid Carcinoma. *Front Endocrinol (Lausanne)* 2022; **13**: 861869 [PMID: 35350101 DOI: 10.3389/fendo.2022.861869]

23 **Tong C**, Wang W, Xia Y, He C. A potential novel biomarker in predicting lymph node metastasis of gastric signet ring cell carcinoma: A derived monocyte to lymphocyte ratio. *Am J Surg* 2022; **223**: 1144-1150 [PMID: 34702491 DOI: 10.1016/j.amjsurg.2021.10.026]

24 **Zhang D**, He W, Wu C, Tan Y, He Y, Xu B, Chen L, Li Q, Jiang J. Scoring System for Tumor-Infiltrating Lymphocytes and Its Prognostic Value for Gastric Cancer. *Front Immunol* 2019; **10**: 71 [PMID: 30761139 DOI: 10.3389/fimmu.2019.00071]

25 **Feng F**, Zheng G, Wang Q, Liu S, Liu Z, Xu G, Wang F, Guo M, Lian X, Zhang H. Low lymphocyte count and high monocyte count predicts poor prognosis of gastric cancer. *BMC Gastroenterol* 2018; **18**: 148 [PMID: 30305076 DOI: 10.1186/s12876-018-0877-9]

26 **Lee HE**, Chae SW, Lee YJ, Kim MA, Lee HS, Lee BL, Kim WH. Prognostic implications of type and density of tumour-infiltrating lymphocytes in gastric cancer. *Br J Cancer* 2008; **99**: 1704-1711 [PMID: 18941457 DOI: 10.1038/sj.bjc.6604738]

27 **Li F**, Sun Y, Huang J, Xu W, Liu J, Yuan Z. CD4/CD8 + T cells, DC subsets, Foxp3, and IDO expression are predictive indictors of gastric cancer prognosis. *Cancer Med* 2019; **8**: 7330-7344 [PMID: 31631566 DOI: 10.1002/cam4.2596]

28 **Vivier E**, Tomasello E, Baratin M, Walzer T, Ugolini S. Functions of natural killer cells. *Nat Immunol* 2008; **9**: 503-510 [PMID: 18425107 DOI: 10.1038/ni1582]

29 **Vivier E**, Nunès JA, Vély F. Natural killer cell signaling pathways. *Science* 2004; **306**: 1517-1519 [PMID: 15567854 DOI: 10.1126/science.1103478]

30 **Hinshaw DC**, Shevde LA. The Tumor Microenvironment Innately Modulates Cancer Progression. *Cancer Res* 2019; **79**: 4557-4566 [PMID: 31350295 DOI: 10.1158/0008-5472.CAN-18-3962]

31 **Du Y**, Wei Y. Therapeutic Potential of Natural Killer Cells in Gastric Cancer. *Front Immunol* 2018; **9**: 3095 [PMID: 30719024 DOI: 10.3389/fimmu.2018.03095]

32 **Wang Z**, Si X, Xu A, Meng X, Gao S, Qi Y, Zhu L, Li T, Li W, Dong L. Activation of STAT3 in human gastric cancer cells via interleukin (IL)-6-type cytokine signaling correlates with clinical implications. *PLoS One* 2013; **8**: e75788 [PMID: 24116074 DOI: 10.1371/journal.pone.0075788]

33 **Rihawi K**, Ricci AD, Rizzo A, Brocchi S, Marasco G, Pastore LV, Llimpe FLR, Golfieri R, Renzulli M. Tumor-Associated Macrophages and Inflammatory Microenvironment in Gastric Cancer: Novel Translational Implications. *Int J Mol Sci* 2021; **22** [PMID: 33916915 DOI: 10.3390/ijms22083805]

34 **Biswas SK**, Mantovani A. Macrophage plasticity and interaction with lymphocyte subsets: cancer as a paradigm. *Nat Immunol* 2010; **11**: 889-896 [PMID: 20856220 DOI: 10.1038/ni.1937]

35 **Räihä MR**, Puolakkainen PA. Tumor-associated macrophages (TAMs) as biomarkers for gastric cancer: A review. *Chronic Dis Transl Med* 2018; **4**: 156-163 [PMID: 30276362 DOI: 10.1016/j.cdtm.2018.07.001]

**Footnotes**

**Institutional review board statement:** Institutional review board statement: The study was reviewed and approved by the Xiangya Hospital, Central South University Institutional Review Board [Approval No. Xykyll-lx-2019030510].

**Informed consent statement:** All study participants, or their legal guardian, provided informed written consent prior to study enrollment.

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**Figure Legends**

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**Figure 1 Receiver operating characteristic curves of all predictors.** A: Age; B: Carcinoembryonic antigen; C: Albumin; D: Prognostic nutritional index; E: Lymphocyte to monocyte ratio; F: Platelet to lymphocyte ratio; G: Fibrinogen.



**Figure 2 The survival curves of all predictors.** A: Resection scope; B: Infiltration depth; C: Lymph node metastasis; D: pTNM staging;



**Figure 3 Survival of patients with different characteristics in Cox regression model.** CEA: Carcinoembryonic antigen; LMR: Lymphocytes to monocytes.

**Table 1 Baseline data and pathological features of the study participants**

|  |  |
| --- | --- |
| **Characteristics** | ***n* (%)** |
| Gender |  |
| Male | 117 (55.19) |
| Female | 95 (44.81) |
| Tumor site |  |
| Upper third | 5 (2.36) |
| Middle third | 58 (27.36) |
| Lower third | 149 (70.28) |
| pT |  |
| pT1-2 | 79 (37.26) |
| pT3-4 | 133 (62.74) |
| pN |  |
| pN0 | 94 (44.34) |
| pN1-3 | 118 (55.66) |
| pTNM stage |  |
| Ⅰ/Ⅱ | 117 (55.19) |
| Ⅲ | 95 (44.81) |
| Resection scope |  |
| Distal stomach | 167 (78.78) |
| Total stomach | 45 (21.23) |
| Survival status |  |
| Survival | 125 (58.96) |
| Dead | 87 (41.04) |

**Table 2 Receiver operating characteristic curve analysis results of predictors**

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Variables** | **AUC (95%CI)** | ***P* value** | **Cut-off value** | **Sensitivity** | **Specificity** |
| Age | 0.589 (0.509, 0.669) | 0.028 | 55.5 | 0.494 | 0.728 |
| PLR | 0.589 (0.511, 0.666) | 0.028 | 124.63 | 0.586 | 0.608 |
| LMR | 0.590 (0.512, 0.669) | 0.025 | 3.83 | 0.54 | 0.632 |
| ALB | 0.618 (0.540, 0.696) | 0.004 | 38.95 | 0.414 | 0.792 |
| PNI | 0.644 (0.567, 0.721) | ＜ 0.001 | 49.85 | 0.632 | 0.608 |
| FIB | 0.624 (0.547, 0.701) | 0.002 | 3.115 | 0.655 | 0.648 |
| CEA | 0.632 (0.556, 0.708) | 0.001 | 1.455 | 0.471 | 0.744 |

PLR: Platelet to lymphocyte ratio; LMR: Lymphocyte to monocyte ratio; ALB: Albumin; PNI: Prognostic nutritional index; FIB: Fibrinogen; CEA: Carcinoembryonic antigen; AUC: Area under the curve.

**Table 3 Relationship between baseline data, pathological features and 5-year survival rate**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Group** | **Survival status** | **5-OS (%)** | ***χ*2** | ***P* value** |
|  | **Survival(n)** | **Dead(n)** |  |  |  |
| Gender |  |  |  | 1.27 | 0.26 |
| Male | 73 | 44 | 62.39 |  |  |
| Female | 52 | 43 | 54.74 |  |  |
| Tumor site |  |  |  | 3.556 | 0.169 |
| Upper third  | 1 | 4 | 20 |  |  |
| Middle third | 33 | 25 | 56.9 |  |  |
| Lower third | 91 | 58 | 61.07 |  |  |
| pT |  |  |  | 49.726 | ＜ 0.001 |
| pT1-2 | 71 | 8 | 89.87 |  |  |
| pT3-4 | 54 | 79 | 40.6 |  |  |
| pN |  |  |  | 30.269 | ＜ 0.001 |
| pN0 | 75 | 19 | 79.79 |  |  |
| pN1-3 | 50 | 68 | 42.37 |  |  |
| pTNM stage |  |  |  | 49.322 | ＜ 0.001 |
| Ⅰ/Ⅱ | 94 | 23 | 80.34 |  |  |
| Ⅲ | 31 | 64 | 32.63 |  |  |
| Resection scope |  |  |  | 8.489 | 0.004 |
| Distal stomach | 107 | 60 | 64.07 |  |  |
| Total stomach | 18 | 27 | 40 |  |  |

OS: Overall survival.

**Table 4 Relationship between indicators and survival status**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Variables** | **Survival** | **Dead** | **t/Z value** | ***P* value** |
| Age | 50 ± 10.82 | 53 ± 11.65 | -2.213 | 0.028 |
| CEA | 0.90 (0.58, 1.48) | 1.21 (0.75, 2.35) | -3.265 | 0.001 |
| NLR | 1.82 (1.30, 2.43) | 2.05 (1.45, 2.71) | -1.668 | 0.095 |
| PLR | 118.33 (99.41, 150.18) | 130.59 (108.10, 162.00) | -2.196 | 0.028 |
| LMR | 4.20 (3.40, 5.67) | 3.80 (2.80, 5.00) | -2.226 | 0.026 |
| SII | 392.36 (275.38, 565.21) | 448.00 (311.67, 600.71) | -1.391 | 0.164 |
| PT | 12.40 (11.96, 12.90) | 12.30 (12.00, 12.70) | -1.114 | 0.265 |
| INR | 0.97 (0.93, 1.02) | 0.95 (0.93, 0.99) | -1.728 | 0.084 |
| APTT | 32.71 ± 4.16 | 31.90 ± 3.98 | 1.425 | 0.156 |
| TT | 17.55 ± 1.73 | 17.66 ± 1.85 | -0.424 | 0.672 |
| FIB | 2.93 (2.56, 3.42) | 3.29 (2.81, 3.80) | -3.065 | 0.002 |
| ALB (g/L) | 41.79 ± 3.83 | 40.09 ± 4.72 | 2.854 | 0.005 |
| GLB (g/L) | 26.00 (23.80, 29.40) | 25.20 (22.80, 28.50) | -1.356 | 0.175 |
| AGR | 1.58 (1.41, 1.80) | 1.56 (1.40, 1.75) | -0.716 | 0.474 |
| PNI | 51.11 ± 4.95 | 48.21 ± 6.15 | 3.789 | ＜ 0.001 |

CEA: Carcinoembryonic antigen; NLR: Neutrophil to lymphocyte ratio; PLR: Platelet to lymphocyte ratio; LMR: Lymphocyte to monocyte ratio; SII: Systemic immune inflammation index; PT: Prothrombin time; INR: International normalized ratio; APTT: Activated partial thromboplastin time; TT: Thrombin time; FIB: Fibrinogen; ALB: Albumin; GLB: Globulin; AGR: Albumin to globulin ratio; PNI: Prognostic nutritional index.

**Table 5 Grouping of predictors**

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
|  |  | **Survival(n)** | **Dead(n)** | **5-OS (%)** | **Total** |
| Age | <56 | 91 | 44 | 67.41 | 135 |
|  | ≥ 56 | 34 | 43 | 44.16 | 77 |
| CEA (ng/mL) | < 1.455 | 93 | 46 | 66.91 | 139 |
|  | ≥ 1.455 | 32 | 41 | 43.84 | 73 |
| PLR | < 124.63 | 76 | 36 | 67.86 | 112 |
|  | ≥ 124.63 | 49 | 51 | 49 | 100 |
| LMR | < 3.83 | 81 | 30 | 72.97 | 111 |
|  | ≥ 3.83 | 44 | 57 | 43.56 | 101 |
| ALB (g/L) | < 38.95 | 26 | 36 | 41.94 | 62 |
|  | ≥ 38.95 | 99 | 51 | 66 | 150 |
| PNI | < 49.85 | 49 | 55 | 47.12 | 104 |
|  | ≥ 49.85 | 76 | 32 | 70.37 | 108 |
| FIB (g/L) | < 3.115 | 46 | 47 | 49.46 | 93 |
|  | ≥ 3.115 | 79 | 40 | 66.39 | 119 |

OS: Overall survival; CEA: Carcinoembryonic antigen; PLR: Platelet to lymphocyte ratio; LMR: Lymphocyte to monocyte ratio; ALB: Albumin; PNI: Prognostic nutritional index; FIB: Fibrinogen.



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