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***Observational Study***

**Evaluation of the value of multiparameter combined analysis of serum markers in the early diagnosis of gastric cancer**

Zhang ZG *et al*. Serum markers panel for detection of GC

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**Abstract**

BACKGROUND

In early gastric cancer (GC), tumor markers are increased in the blood. The levels of these markers have been used as important indexes for GC screening, early diagnosis and prognostic evaluation. However, specific tumor markers have not yet been discovered. Diagnosis based on a single tumor marker has limited significance. The detection rate of GC is still very low

AIM

To improve the diagnostic value of blood markers for GC.

METHODS

We used a multiparameter joint analysis of 77 indexes of malignant GC and gastric polyp (GP), 64 indexes of GC and healthy controls (Ctrls).

RESULTS

By analyzing the data, there are 27 indexes in the final Ctrls *vs* GC with *P* values < 0.01, the area under the curve (AUC) of albumin is the largest in Ctrls *vs* GC, and the AUC was 0.907. 30 indexes in GP *vs* GC have *P* values < 0.01. Among them, the D-dimer showed an AUC of 0.729. The 27 indexes in Ctrls *vs* GC and 30 indexes in GP *vs* GC were used for binary logistic regression, discriminant analysis, classification tree analysis and artificial neural network analysis model. For the ability to distinguish between Ctrls *vs* GC, GP *vs* GC, artificial neural networks had better diagnostic value when compared with classification tree, binary logistic regression, and discriminant analysis. When compared Ctrl and GC, the overall prediction accuracy was 92.9%, and the AUC was 0.992 (0.980, 1.000). When compared GP and GC, the overall prediction accuracy was 77.9%, and the AUC was 0.969 (0.948, 0.990).

CONCLUSION

The diagnostic effect of multi-parameter joint artificial neural networks analysis is significantly better than the single-index test diagnosis, and it may provide an assistant method for the detection of GC.

**Key words:** Gastric cancer; Gastric polyp; Serum; Artificial neural network; Detection

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**Core tip:** In this study, we aimed to improve the diagnostic value of blood markers for gastric cancer. By comparing the binary logistic regression, discriminant analysis, classification tree and artificial neural network analysis, finally, artificial neural networks had better diagnostic value. When compared healthy control and gastric cancer, gastric polyp and gastric cancer, the area under the curve was 0.992 (0.980, 1.000) and 0.969 (0.948, 0.990), respectively. Based on artificial neural network and serum index, a novel diagnostic model for gastric cancer may be provided for clinical practice.

**INTRODUCTION**

According to estimates by the World Health Organization, nearly 7 million people worldwide die from cancer every year, and this number is increasing every year. Gastric cancer (GC) is a common malignant tumor that endangers human health. GC ranks second in cancer-related deaths. In China, GC is one of the most malignant tumors with high morbidity and mortality[1]. GC deaths account for approximately 25% to 30% of all cancer-type deaths[2]. The pathogenesis of GC involves physical aging, eating habits and psychological factors[3-5]. The development and progression of GC is a multistage process involving multiple changes at the gene and molecular levels. In the early stage of GC, there are precancerous lesions, most of which remain unchanged, and a small part of which develop into cancer. The Correa cascade is the most common pattern of GC[6]. In current clinical practice, the main treatment for GC is surgical treatment. The 5-year survival rate is very low[7]; however, if GC is detected early, then the 5-year survival rate can be as high as 90%[8]. In developed countries, such as Japan, where the early diagnosis of GC reached 50%, the five-year survival rate reached 90%[9]. The early diagnosis and treatment of GC are extremely important for patients with GC.

Currently, many methods for diagnosing GC are used in scientific research and clinical practice[10]. Among these methods, plasma biomarker detection is an important detection method. The most commonly used tumor markers for early GC detection include carcinoembryonic antigen (CEA), carbohydrate antigens (CA): CA19-9, CA72-4, CA125, CA24-2, CA50, and pepsinogen and alpha-fetoprotein (AFP)[11]. However, these tumor biomarkers are poorly specific and sensitive, and thus far, they have not been used alone for the diagnosis of GC[11,12]. In early GC, tumor markers, such as CEA and CA-724, are increased in the blood. The levels of these markers have been used as important indexes for GC screening, early diagnosis and prognostic evaluation[13]. However, specific tumor markers have not yet been discovered. Diagnosis based on a single tumor marker has limited significance[14]. The detection rate of GC is still very low.

In this study, to distinguish between healthy controls (Ctrls) *vs* GC, gastric polyp (GP) and GC, we analyzed the routine blood detection indexes of GC diagnosis by using binary logistic regression, discriminant analysis, classification tree and artificial neural network. We aimed to use multiparameter joint analysis to improve diagnostic sensitivity and specificity and provide a new potential method for the early diagnosis of GC in clinical practice.

**MATERIALS AND METHODS**

***Patient sample***

The serum samples of the patients involved in this study were obtained from the blood samples of patients admitted to the Beijing Daxing District People’s Hospital from April 2016 to April 2019 and confirmed by imaging and pathology. Sample collection and data screening were approved by the Ethics Committee of Beijing Daxing District People’s Hospital.

The inclusion criteria of the disease group were complete clinical and pathological data of the patient, with clear imaging and pathological diagnosis, and no radiotherapy, chemotherapy or other immunotherapy before surgery. The exclusion criteria for the disease group were patients with major diseases associated with the study, combined with other types of tumors, or individuals that had received radiotherapy, chemotherapy, or other immunotherapy before surgery. As shown in Table 1, this study included 144 GP and 253 GC patients. A total of 370 healthy controls were examined for tumor markers and imaging examinations. There were no diseases associated with this study, and both tumor markers and imaging examinations were qualified.

All subjects involved in the study provided early morning fasting peripheral blood samples. EDTA was used as an anticoagulant, and after centrifugation at 3500 r/min for 7 min, the patient serum was collected in a new Eppendorf tube. The serum was then dispensed into 3 tubes and labeled and immediately stored in a -80°C. During the collection process, it is necessary to pay attention to the removal of serum samples of hemolysis or lipemia and avoid repeated freezing and thawing during the test. When testing, directly remove the thawed test samples.

***Data analysis***

Using SPSS 22.0 statistical software, 77 indexes of GC and GP, 64 indexes of GC and Ctrls were analyzed. The serum levels of each index of GC and GP, Ctrls of GC were compared by an independent samples *T* test[15]. The diagnostic value was evaluated by the area under curve (AUC) of the receiver operating characteristic (ROC), and the cutoff value was determined by the Youden index. The combination of indexes was analyzed by statistical methods, such as binary logistic regression analysis, discriminant analysis, classification tree and artificial neural network[16-21]. *P* < 0.01 was considered statistically significant.

**RESULTS**

***Significant analysis of Ctrls vs GC, GP vs GC and ROC analysis***

There were significant differences in 40 indexes between Ctrls *vs* GC, and 24 indexes had no significant difference; 39 indexes of GP *vs* GC were significantly different, and 38 indexes had no significant difference. The ROC were generated for 40 indexes with significant differences in Ctrls *vs* GC and 39 indexes with significant differences between GP *vs* GC. Among these indexes, the largest AUC in Ctrls *vs* GC was and ALB, with values of 0.907. When the ALB cutoff value was 42.05, the sensitivity and specificity were 93.0% and 79.1%, respectively. In GP *vs* GC, the largest AUC was for D-dimer. The AUC value was 0.729. When the D-dimer cutoff value was 0.435, the sensitivity and specificity were 55.3% and 81.2%, respectively.

***Binary logistic******regression and discriminant analysis results of Ctrls vs GC, GP vs GC***

The 27 indexes in Ctrls *vs* GC and 30 indexes in GP *vs* GC were used to establish a binary logistic regression analysis model (70% of the data). As shown in Figure 1A, the AUC for Ctrls *vs* GC was 0.989 (0.982, 0.995). When the cutoff value was 0.675, the sensitivity and specificity were 93.4% and 95.5%, respectively. As shown in Figure 1B, the AUC of GP *vs* GC was 0.929 (0.901, 0.958), when the cutoff value was 0.477, the sensitivity and specificity were 85.1% and 87.6%, respectively. Binary logistic regression analysis is significantly better than the distinction between Ctrls *vs* GC for distinguishing GP *vs* GC. As shown in Figure 1C, the AUC of Ctrls *vs* GC was 0.971 (0.957, 0.985), and when the cutoff value was 0.470, the sensitivity and specificity were 86.4% and 97.3%, respectively. As shown in Figure 1D, the GP *vs* GC AUC was 0.914 (0.882, 0.946), and when the cutoff value was 0.462, the sensitivity and specificity were 78.0% and 92.1%, respectively. Discriminant analysis is significantly better than the distinction between Ctrls *vs* GC for distinguishing GP *vs* GC.

***Classification tree analysis and artificial neural network results of Ctrls vs GC, GP vs GC***

The 27 indexes in Ctrls *vs* GC and 30 indexes in GP *vs* GC were used to establish a classification tree analysis model. As shown in Figure 2A, the AUC of Ctrls *vs* GC was 0.863 (0.826, 0.900), and when the cutoff value was 0.520, the sensitivity and specificity were 74.0% and 76.3%, respectively. The prediction accuracy rate of the Ctrls was 100%, the prediction accuracy rate of the mGC was 48.2%, and the overall prediction accuracy rate was 76.8%. As shown in Figure 2B, the AUC of GP *vs* GC was 0.739 (0.680, 0.799), and when the cutoff value was 0.290, the sensitivity and specificity were 85.8% and 75.3%, respectively. The predictive accuracy rate of the GP was 62.1%, the correct rate of the GC was 67.8%, and the overall prediction accuracy rate was 65.9%. As shown in Figure 2C, the AUC of Ctrls *vs* GC was 0.992 (0.980, 1.000). When the cutoff value is 0.837, the sensitivity and specificity were 96.0% and 99.6%, respectively; the prediction accuracy rate of the Ctrls was 97.5%, the prediction accuracy rate of the GC was 84.8%, and the overall prediction accuracy rate was 92.9%. As shown in Figure 2D, the AUC of bGC *vs* mGC was 0.969 (0.948, 0.990); when the cutoff value was 0.970, the sensitivity and specificity were 94.9% and 96.0%, respectively. The predictive accuracy rate of GP was 71.0%, the predictive accuracy rate of GC was 82.6%, and the overall prediction accuracy rate was 77.9%.

**DISCUSSION**

Through saliency analysis and ROC curve analysis, there were 27 indexes in the final Ctrls *vs* GC with a *P* value of < 0.01 and 30 indexes in the GP *vs* GC with a *P* value of < 0.01. Among these indexes, the maximum AUC of Ctrls *vs* GC is ALB, and the AUC values were 0.907. The maximum AUC of GP *vs* GC is D-dimer, and the AUC was 0.729. Pre-ALB levels had been demonstrated to correlate with the outcomes of surgical patients[22,23]. It was usually used to assess the nutritional status. Lots of studies demonstrated that the poor postoperative nutritional status of GC may be related to worse prognosis[24,25]. In our study, we found that it was related to the development of GC. D-dimer is a widely used biomarker for evaluating the ability of coagulation and fibrinolysis, and involved in the progression of cancers[26]. Plasma D-dimer levels was significantly increased in GC patients with distant metastases, and it may be a promising biomarker of detection of GC[27]. In addition, high plasma D-dimer level may also predict poor prognosis in gynecological tumor[28].

With the rapid development of molecular technology, kinds of molecular detection methods had been explored[29-33]. Many statistical methods currently used in the multi-index joint detection analysis of cancer[15,21,34-36], such as binary logistic regression, discriminant analysis, classification tree and artificial neural network, have achieved good results[16-20]. For example, the artificial neural network model was applied in lung cancer-assisted diagnosis, and the effects of back-propagation neural network and Fisher discriminant model on lung cancer screening were compared by the joint detection of four biomarkers. The results showed that the back-propagation neural network predicts lung cancer model better than the Fisher discriminant analysis, which can provide excellent and intelligent diagnostic tools for lung cancer[37]. Li *et al*[38] used binary logistic regression analysis to analyze various cytokines in serum for the early detection of GC. Feng *et al*[39] used the ANN model established by six serum tumor markers to distinguish lung cancer, to identify not only benign lung diseases and normal people but also three common gastrointestinal cancers. These results showed that the artificial neural network model may be an excellent intelligent system to distinguish lung cancer[39]. Su *et al*[40] applied a classification decision tree model to distinguish between GC and healthy controls. This model is able to distinguish between GC patients and healthy volunteers. The sensitivity in the training set is 95.6%, and the specificity is 92.0%. In the blinded group, this model was able to distinguish GC samples from other samples with a specificity of 88.0%, a sensitivity of 85.3%, and an accuracy of 86.4%. By measuring serum CEA and CA19-9 together, these values were higher than those obtained in the parallel analysis. Therefore, a decision tree analysis demonstrating a serum proteomics model is likely to be used for the diagnosis of GC[40].

For distinguishing Ctrls *vs* GC, binary logistic regression, discriminant analysis, classification tree analysis and artificial neural network were significantly better than GP *vs* GC. Binary logistic regression, discriminant analysis and artificial neural network analysis of the ROC curve AUC and the maximum cutoff value corresponding to the sensitivity and specificity were greater than the AUC maximum single index. Therefore, the diagnostic effect of multiparameter joint analysis is significantly better than that of the single-index test. Through the comparison of these four methods, we have the ability to distinguish Ctrls *vs* mGC, bGC *vs* mGC, artificial neural network > binary logistic regression > discriminant analysis > classification tree. However, the results may be effected because of the relatively little sample size and lack of independent validation of the model which was built in our study. We propose that the artificial neural network analysis method has good prospects for the multi-index joint detection of tumors, and further research in this area should be carried out in the future.

**ARTICLE HIGHLIGHTS**

***Research background***

Tumor markers are increased in the blood in early gastric cancer (GC). The levels of these markers have been used as important indexes for GC screening, early diagnosis and prognostic evaluation.

***Research motivation***

Specific tumor markers have not yet been discovered. Diagnosis based on a single tumor marker has limited significance. The detection rate of GC is still very low.

***Research objectives***

In this study, we aimed to improve the diagnostic value of blood markers for GC.

***Research methods***

In this study, to distinguish between healthy controls (Ctrls) *vs* GC, gastric polyp (GP) and GC, we analyzed the routine blood detection indexes of GC diagnosis by using binary logistic regression, discriminant analysis, classification tree and artificial neural network.

***Research results***

By analyzing the data, there are 27 indexes in the final Ctrls *vs* GC with *P* values < 0.01, the area under the curve (AUC) of albumin is the largest in Ctrls *vs* GC, and the AUC was 0.907. For 30 indexes in GP *vs* GC have *P* values < 0.01. Among them, the D-dimer showed an AUC of 0.729. The 27 indexes in Ctrls *vs* GC and 30 indexes in GP *vs* GC were used for binary logistic regression, discriminant analysis, classification tree analysis and artificial neural network analysis model. The overall prediction accuracy was 92.9%, and the AUC was 0.992 (0.980, 1.000).

***Research conclusions***

The diagnostic effect of multi-parameter joint artificial neural networks analysis is significantly better than the single-index test diagnosis, and it may provide an assistant method for the detection of GC.

***Research perspectives***

We propose that the artificial neural network analysis method has good prospects for the multi-index joint detection of tumors, and further research in this area should be carried out in the future.

**REFERENCES**

1 **Ning FL**, Zhang CD, Wang P, Shao S, Dai DQ. Endoscopic resection versus radical gastrectomy for early gastric cancer in Asia: A meta-analysis. *Int J Surg* 2017; **48**: 45-52 [PMID: 28987558 DOI: 10.1016/j.ijsu.2017.09.068]

2 **Zheng Q**, Chen C, Guan H, Kang W, Yu C. Prognostic role of microRNAs in human gastrointestinal cancer: A systematic review and meta-analysis. *Oncotarget* 2017; **8**: 46611-46623 [PMID: 28402940 DOI: 10.18632/oncotarget.16679]

3 **Kalisperati P**, Spanou E, Pateras IS, Korkolopoulou P, Varvarigou A, Karavokyros I, Gorgoulis VG, Vlachoyiannopoulos PG, Sougioultzis S. Inflammation, DNA Damage, *Helicobacter pylori* and Gastric Tumorigenesis. *Front Genet* 2017; **8**: 20 [PMID: 28289428 DOI: 10.3389/fgene.2017.00020]

4 **Maleki SS**, Röcken C. Chromosomal Instability in Gastric Cancer Biology. *Neoplasia* 2017; **19**: 412-420 [PMID: 28431273 DOI: 10.1016/j.neo.2017.02.012]

5 **Sunakawa Y**, Lenz HJ. Molecular classification of gastric adenocarcinoma: translating new insights from the cancer genome atlas research network. *Curr Treat Options Oncol* 2015; **16**: 17 [PMID: 25813036 DOI: 10.1007/s11864-015-0331-y]

6 **Futawatari N**, Fukuyama T, Yamamura R, Shida A, Takahashi Y, Nishi Y, Ichiki Y, Kobayashi N, Yamazaki H, Watanabe M. Early gastric cancer frequently has high expression of KK-LC-1, a cancer-testis antigen. *World J Gastroenterol* 2017; **23**: 8200-8206 [PMID: 29290656 DOI: 10.3748/wjg.v23.i46.8200]

7 **Ahn S**, Park DY. Practical Points in Gastric Pathology. *Arch Pathol Lab Med* 2016; **140**: 397-405 [PMID: 27128297 DOI: 10.5858/arpa.2015-0300-RA]

8 **Beeharry MK**, Liu WT, Yan M, Zhu ZG. New blood markers detection technology: A leap in the diagnosis of gastric cancer. *World J Gastroenterol* 2016; **22**: 1202-1212 [PMID: 26811658 DOI: 10.3748/wjg.v22.i3.1202]

9 **Pasechnikov V**, Chukov S, Fedorov E, Kikuste I, Leja M. Gastric cancer: prevention, screening and early diagnosis. *World J Gastroenterol* 2014; **20**: 13842-13862 [PMID: 25320521 DOI: 10.3748/wjg.v20.i38.13842]

10 **Uedo N**, Yao K. Endoluminal Diagnosis of Early Gastric Cancer and Its Precursors: Bridging the Gap Between Endoscopy and Pathology. *Adv Exp Med Biol* 2016; **908**: 293-316 [PMID: 27573777 DOI: 10.1007/978-3-319-41388-4\_14]

11 **Tsai MM**, Wang CS, Tsai CY, Huang HW, Chi HC, Lin YH, Lu PH, Lin KH. Potential Diagnostic, Prognostic and Therapeutic Targets of MicroRNAs in Human Gastric Cancer. *Int J Mol Sci* 2016; **17**: 945 [PMID: 27322246 DOI: 10.3390/ijms17060945]

12 **Tong W**, Ye F, He L, Cui L, Cui M, Hu Y, Li W, Jiang J, Zhang DY, Suo J. Serum biomarker panels for diagnosis of gastric cancer. *Onco Targets Ther* 2016; **9**: 2455-2463 [PMID: 27217769 DOI: 10.2147/OTT.S86139]

13 **Feng F**, Tian Y, Xu G, Liu Z, Liu S, Zheng G, Guo M, Lian X, Fan D, Zhang H. Diagnostic and prognostic value of CEA, CA19-9, AFP and CA125 for early gastric cancer. *BMC Cancer* 2017; **17**: 737 [PMID: 29121872 DOI: 10.1186/s12885-017-3738-y]

14 **Zhang Q**, Qu H, Sun G, Li Z, Ma S, Shi Z, Zhao E, Zhang H, He Q. Early postoperative tumor marker responses provide a robust prognostic indicator for N3 stage gastric cancer. *Medicine (Baltimore)* 2017; **96**: e7560 [PMID: 28796039 DOI: 10.1097/MD.0000000000007560]

15 **Xu W**, Zhao Y, Nian S, Feng L, Bai X, Luo X, Luo F. Differential analysis of disease risk assessment using binary logistic regression with different analysis strategies. *J Int Med Res* 2018; **46**: 3656-3664 [PMID: 29882459 DOI: 10.1177/0300060518777173]

16 **Bicciato S**. Artificial neural network technologies to identify biomarkers for therapeutic intervention. *Curr Opin Mol Ther* 2004; **6**: 616-623 [PMID: 15663326]

17 **Berger RP**, Beers SR, Richichi R, Wiesman D, Adelson PD. Serum biomarker concentrations and outcome after pediatric traumatic brain injury. *J Neurotrauma* 2007; **24**: 1793-1801 [PMID: 18159990 DOI: 10.1089/neu.2007.0316]

18 **Lin KC**, Wu HP, Huang CY, Lin CY, Chang CF. Discriminant analysis of serum inflammatory biomarkers which differentiate pediatric appendicitis from other acute abdominal diseases. *Acta Paediatr Taiwan* 2007; **48**: 125-130 [PMID: 17912983]

19 **Navaglia F**, Fogar P, Basso D, Greco E, Padoan A, Tonidandel L, Fadi E, Zambon CF, Bozzato D, Moz S, Seraglia R, Pedrazzoli S, Plebani M. Pancreatic cancer biomarkers discovery by surface-enhanced laser desorption and ionization time-of-flight mass spectrometry. *Clin Chem Lab Med* 2009; **47**: 713-723 [PMID: 19426140 DOI: 10.1515/CCLM.2009.158]

20 **Liu Z**, Lin S, Tan MT. Sparse support vector machines with Lp penalty for biomarker identification. *IEEE/ACM Trans Comput Biol Bioinform* 2010; **7**: 100-107 [PMID: 20150672 DOI: 10.1109/TCBB.2008.17]

21 **Zhang P**, Zou M, Wen X, Gu F, Li J, Liu G, Dong J, Deng X, Gao J, Li X, Jia X, Dong Z, Chen L, Wang Y, Tian Y. Development of serum parameters panels for the early detection of pancreatic cancer. *Int J Cancer* 2014; **134**: 2646-2655 [PMID: 24615168 DOI: 10.1002/ijc.28584]

22 **Saito H**, Kono Y, Murakami Y, Shishido Y, Kuroda H, Matsunaga T, Fukumoto Y, Osaki T, Ashida K, Fujiwara Y. Prognostic Significance of the Preoperative Ratio of C-Reactive Protein to Albumin and Neutrophil-Lymphocyte Ratio in Gastric Cancer Patients. *World J Surg* 2018; **42**: 1819-1825 [PMID: 29270656 DOI: 10.1007/s00268-017-4400-1]

23 **Saito H**, Kono Y, Murakami Y, Shishido Y, Kuroda H, Matsunaga T, Fukumoto Y, Osaki T, Ashida K, Fujiwara Y. Postoperative Serum Albumin is a Potential Prognostic Factor for Older Patients with Gastric Cancer. *Yonago Acta Med* 2018; **61**: 72-78 [PMID: 29599625]

24 **Jin Y**, Yong C, Ren K, Li D, Yuan H. Effects of Post-Surgical Parenteral Nutrition on Patients with Gastric Cancer. *Cell Physiol Biochem* 2018; **49**: 1320-1328 [PMID: 30205371 DOI: 10.1159/000493410]

25 **Wu M**, Pan Y, Jia Z, Wang Y, Yang N, Mu J, Zhou T, Guo Y, Jiang J, Cao X. Preoperative Plasma Fibrinogen and Serum Albumin Score Is an Independent Prognostic Factor for Resectable Stage II-III Gastric Cancer. *Dis Markers* 2019; **2019**: 9060845 [PMID: 31781312 DOI: 10.1155/2019/9060845]

26 **Dai H**, Zhou H, Sun Y, Xu Z, Wang S, Feng T, Zhang P. D-dimer as a potential clinical marker for predicting metastasis and progression in cancer. *Biomed Rep* 2018; **9**: 453-457 [PMID: 30402229 DOI: 10.3892/br.2018.1151]

27 **Repetto O**, De Re V. Coagulation and fibrinolysis in gastric cancer. *Ann N Y Acad Sci* 2017; **1404**: 27-48 [PMID: 28833193 DOI: 10.1111/nyas.13454]

28 **Xu L**, He F, Wang H, Gao B, Wu H, Zhao S. A high plasma D-dimer level predicts poor prognosis in gynecological tumors in East Asia area: a systematic review and meta-analysis. *Oncotarget* 2017; **8**: 51551-51558 [PMID: 28881667 DOI: 10.18632/oncotarget.17936]

29 **Gao W**, Long L, Tian X, Xu F, Liu J, Singh PK, Botella JR, Song C. Genome Editing in Cotton with the CRISPR/Cas9 System. *Front Plant Sci* 2017; **8**: 1364 [PMID: 28824692 DOI: 10.3389/fpls.2017.01364]

30 **Guo J**, Li K, Jin L, Xu R, Miao K, Yang F, Qi C, Zhang L, Botella JR, Wang R, Miao Y. A simple and cost-effective method for screening of CRISPR/Cas9-induced homozygous/biallelic mutants. *Plant Methods* 2018; **14**: 40 [PMID: 29872452 DOI: 10.1186/s13007-018-0305-8]

31 **Lei KJ**, Lin YM, An GY. miR156 modulates rhizosphere acidification in response to phosphate limitation in Arabidopsis. *J Plant Res* 2016; **129**: 275-284 [PMID: 26659856 DOI: 10.1007/s10265-015-0778-8]

32 **Sun Q**, Qiao J, Zhang S, He S, Shi Y, Yuan Y, Zhang X, Cai Y. Changes in DNA methylation assessed by genomic bisulfite sequencing suggest a role for DNA methylation in cotton fruiting branch development. *PeerJ* 2018; **6**: e4945 [PMID: 29915693 DOI: 10.7717/peerj.4945]

33 **Yu J**, Zhang Y, Liu J, Wang L, Liu P, Yin Z, Guo S, Ma J, Lu Z, Wang T, She Y, Miao Y, Ma L, Chen S, Li Y, Dai S. Proteomic discovery of H2O2 response in roots and functional characterization of PutGLP gene from alkaligrass. *Planta* 2018; **248**: 1079-1099 [PMID: 30039231 DOI: 10.1007/s00425-018-2940-8]

34 **Liu MM**, Wen L, Liu YJ, Cai Q, Li LT, Cai YM. Application of data mining methods to improve screening for the risk of early gastric cancer. *BMC Med Inform Decis Mak* 2018; **18**: 121 [PMID: 30526601 DOI: 10.1186/s12911-018-0689-4]

35 **Mohammadzadeh F**, Noorkojuri H, Pourhoseingholi MA, Saadat S, Baghestani AR. Predicting the probability of mortality of gastric cancer patients using decision tree. *Ir J Med Sci* 2015; **184**: 277-284 [PMID: 24626962 DOI: 10.1007/s11845-014-1100-9]

36 **Zhang Y**, Liu Y, Zhang J, Wu X, Ji X, Fu T, Li Z, Wu Q, Bu Z, Ji J. Construction and external validation of a nomogram that predicts lymph node metastasis in early gastric cancer patients using preoperative parameters. *Chin J Cancer Res* 2018; **30**: 623-632 [PMID: 30700931 DOI: 10.21147/j.issn.1000-9604.2018.06.07]

37 **Duan X**, Yang Y, Tan S, Wang S, Feng X, Cui L, Feng F, Yu S, Wang W, Wu Y. Application of artificial neural network model combined with four biomarkers in auxiliary diagnosis of lung cancer. *Med Biol Eng Comput* 2017; **55**: 1239-1248 [PMID: 27766520 DOI: 10.1007/s11517-016-1585-7]

38 **Li J**, Xu L, Run ZC, Feng W, Liu W, Zhang PJ, Li Z. Multiple cytokine profiling in serum for early detection of gastric cancer. *World J Gastroenterol* 2018; **24**: 2269-2278 [PMID: 29881236 DOI: 10.3748/wjg.v24.i21.2269]

39 **Feng F**, Wu Y, Wu Y, Nie G, Ni R. The effect of artificial neural network model combined with six tumor markers in auxiliary diagnosis of lung cancer. *J Med Syst* 2012; **36**: 2973-2980 [PMID: 21882004 DOI: 10.1007/s10916-011-9775-1]

40 **Su Y**, Shen J, Qian H, Ma H, Ji J, Ma H, Ma L, Zhang W, Meng L, Li Z, Wu J, Jin G, Zhang J, Shou C. Diagnosis of gastric cancer using decision tree classification of mass spectral data. *Cancer Sci* 2007; **98**: 37-43 [PMID: 17052262 DOI: 10.1111/j.1349-7006.2006.00339.x]

**Footnotes**

**Institutional review board statement:** The study was reviewed and approved by the Beijing Daxing District People’s Hospital review board.

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

**Conflict-of-interest statement:** We declare that we have no financial or personal relationships with other individuals or organizations that can inappropriately influence our work and that there is no professional or other personal interest of any nature in any product, service and/or company that could be construed as influencing the position presented in or the review of the manuscript.

**Data sharing statement:** No additional data are available.

**STROBE statement:** The authors have read the STROBE Statement – checklist of items, and the manuscript was prepared and revised according to the STROBE Statement – checklist of items.

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**Figure Legends**



**Figure 1 Binary logistic analysis and discriminant analysis results of normal control *vs* gastric cancer, gastric polyp *vs* gastric cancer.** A: Receiver operating characteristic (ROC) of the binary logistic regression analysis of normal control *vs* gastric cancer (GC); B: ROC of the binary logistic regression analysis of gastric polyp *vs* GC; C: ROC of the discriminant analysis of normal control *vs* GC; D: ROC of the discriminant analysis of gastric polyp *vs* GC.

**Figure 2 Artificial neural network analysis and classification tree analysis results of normal control *vs* gastric cancer, gastric polyp *vs* gastric cancer.** A: Receiver operating characteristic (ROC) of the classification tree analysis of normal control *vs* gastric cancer (GC); B: ROC of the classification tree analysis of gastric polyp *vs* GC; C: ROC of the artificial neural network analysis of normal control *vs* GC; D: ROC of the artificial neural network analysis of gastric polyp *vs* GC.

**Table 1 Clinical characteristics of samples in our study, *n* (%)**

|  |  |  |  |
| --- | --- | --- | --- |
| **Test variables** | **Gastric cancer** | **Benign** | **Normal** |
| Sex |
| Male | 188 (74.31) | 69 (47.92) | 232 (62.70) |
| Female | 65 (25.69) | 75 (52.08) | 138 (37.30) |
| Age (yr) |
| < 40 | 10 (3.95) | 16 (11.11) | 45 (12.16) |
| 40-60 | 93 (36.76) | 74 (51.39) | 272 (73.51) |
| ≥ 60 | 150 (59.29) | 54 (37.50) | 53 (14.33) |
| T |
| 1 | 5 (1.98) |  |  |
| 1a | 9 (3.56) |  |  |
| 1b | 15 (5.93) |  |  |
| 2 | 45 (17.79) |  |  |
| 3 | 77 (30.43) |  |  |
| 4 | 2 (0.79) |  |  |
| 4a | 17 (6.72) |  |  |
| 4b | 1 (0.40) |  |  |
| is | 6 (2.37) |  |  |
| N |
| 0 | 93 (36.76) |  |  |
| 1 | 24 (9.49) |  |  |
| 1a | 4 (1.58) |  |  |
| 1b | 4 (1.58) |  |  |
| 2 | 12 (4.74) |  |  |
| 2a | 5 (1.98) |  |  |
| 2b | 20 (7.91) |  |  |
| 3a | 9 (3.56) |  |  |
| 3b | 6 (2.37) |  |  |
| M |
| 0 | 175 (69.17) |  |  |
| 1 | 43 (17.00) |  |  |
| TNM |
| 0 | 6 (2.37) |  |  |
| I | 57 (22.53) |  |  |
| II | 50 (19.76) |  |  |
| III | 67 (26.48) |  |  |
| IV | 43 (17.00) |  |  |
| Unknown  | 30 (11.86) |  |  |

TNM: Tumor node metastasis.