Identification of molecular biomarkers for the diagnosis of gastric cancer and lymph-node metastasis

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Abstract

Background and objective: Biomarkers are important tools for prompt diagnosis of cancer. This study aimed to identify reliable biomarkers for clinical applications in the diagnosis of gastric cancer and lymph-node (LN) metastasis.

Methods: Between 1 December 2014 and 31 December 2015, we prospectively collected samples of gastric-cancer tissues, corresponding matched-pair normal gastric mucosa, and their peri-gastric metastatic and non-metastatic LNs to identify quantitatively reliable genes using quantitative real-time polymerase chain reaction. Relative quantity (RQ) was used to calculate the mRNA expression levels of our target genes. Statistics were calculated using one-way analysis of variance (ANOVA) and Tukey’s multiple comparison test. Analytical graphs were plotted using GraphPad Prism.

Results: Of nine assessed genes, the mRNA levels of inhibin beta A (INHBA) and secreted phosphoprotein 1 (SPP1) were most consistently highly expressed in tumor tissues by 15.4- and 15.6-fold, respectively, as compared with normal tissues (P < 0.001), with 91.3% sensitivity and 95.7% specificity (receiver operating characteristic [ROC] curve area = 0.974) for the former and 82.6% sensitivity and 87.0% specificity (ROC curve area = 0.924) for the latter. Further analysis revealed no differentiating significance of SPP1 mRNA expression between metastatic and non-metastatic LNs (P = 0.470). In contrast, the INHBA mRNA level was up-regulated 4.1-fold in metastatic LNs (P < 0.001), with 80.0% sensitivity and 81.5% specificity (ROC curve area = 0.857), and was also able to successfully differentiate between more severe disease conditions, T3 and T4 (P = 0.003), M0 and M1 (P = 0.043) and different histological variants (intestinal type vs diffuse type, P = 0.019).
Conclusions: Our results showed that INHBA was the most optimally reliable biomarker for diagnosing gastric cancer and LN metastasis.

Key words: Diagnosis; gastric cancer; lymph node; metastasis; molecular biomarker

Introduction

Gastric cancer, which was once the most common cancer worldwide, is responsible for nearly 1 million new cases and the death of more than 700,000 patients annually [1]. Although gastric cancer is ranked the third leading cause of cancer-related death in both sexes [2], it still has a dismal prognosis due to its late diagnosis and early forms of metastases. Lymph-node (LN) metastasis, which can occur in early stages, is the most commonly observed form of metastasis and has been repeatedly demonstrated as an independent risk factor for survival of gastric cancer [3, 4]. Thereby, identifying prognostic biomarkers to accurately diagnose gastric cancer and LN metastasis is crucial for the proper classification of patients, as they can help to optimize treatment options and result in improved outcomes [5].

Hematoxylin and eosin (H&E) staining is commonly employed in histological examination, for which only the largest cut-cross-sectional dimension of LNs are usually stained and examined [6]. However, as cancer spread inside LNs tend to be randomly distributed, these cancerous cells may at times be missed by conventional H&E staining and lead to pathologically misclassified staging [6, 7]. Consequently, more detailed histological examination might be necessary for improving results, though such an effort would involve the preparation and examination of more slides, increase the overall workload of pathologists and be time-consuming. Rapid- and automated quantitative real-time polymerase chain reaction (qRT-PCR) can examine the entire LNs for cancerous cells and has been shown as a promising adjunct to conventional histological examination for minimizing this effect [8].

Further, recent developments in molecular biology have facilitated the identification of genomic drivers for differentiating between metastatic and non-metastatic LNs and there are numerous studies reporting differential gene expression in gastric cancer using techniques such as DNA micro-array and transcriptome sequencing [9, 10]. However, there is an ever-increasing gap between biomarker discovery and quantitative validation, especially for gastric cancer [11].

Accordingly, to identify reliable gene biomarkers, we collected clinical research data from previous research publications and selected nine highly expressed genes related to gastric cancer, namely H19 [9, 12], osteopontin (OPN) or secreted phosphoprotein 1 (SPP1) [9, 13], chitinase-3-like protein 1 (CHI3L1) [14], inhibin beta A (INHBA) [15, 16], keratin 17 (KRT17) [17, 18], growth factor receptor-bound protein 7 (GRB7) [9], stress-induced phosphoprotein 1 (STIP1) [9], collagen type IV alpha 2 (COL4A2) [9], and procollagen C-endopeptidase enhancer (PCOLCE) [9], to examine their mRNA expression levels in clinical samples using qRT-PCR.

As the expressions of these genes in gastric-cancer tissues have to be further validated and their relation to LN metastasis has not yet been studied, this study aimed to evaluate the diagnostic reliability of the above-mentioned genes to diagnose gastric cancer and peri-gastric metastatic LNs.

Methods

Patients and tissues collected

After receiving approval from the ethics board of Sun Yat-sen University Cancer Center (Guangzhou, China), patients who had pre-operative pathological confirmation of gastric cancer at the Department of Gastric Surgery, Sun Yat-sen University Cancer Center between 1 December 2014 and 31 December 2015 were prospectively enrolled for this study. Signed consent was obtained from all patients. All patients enrolled in the present study underwent radical or palliative surgery, and all cases were staged according to the 7th American Joint Committee on Cancer (AJCC) pathological tumor-node-metastasis (TNM) classification system for gastric cancer. None of the patients had received any form of pre-operative anti-cancer treatment. The samples were collected during surgery from patients who met the following inclusion criteria: (i) tumor size larger than 3 cm (probability of obtaining metastatic LNs would be higher) [19]; (ii) at least one macroscopically identified/hypothesized metastatic LN [20]; (iii) sufficient remnant stomach for retrieval of normal gastric mucosa at least 5 cm away from the primary tumor and/or within the surgical margin proximity; and (iv) absence of Bormann IV cancers. Each LN was dissected into equal symmetrical halves, with one-half being sent for pathological examination and the other half being used for this research study. Using the post-operative pathological report as a reference guide, LNs with and without cancer infiltration were classified as metastatic (LN+) and non-metastatic (LN-) LNs, respectively. All collected specimens were embedded immediately after removal with RNAlater solution and then preserved at −80°C until RNA extraction.

Sample homogenization and RNA extraction

All collected specimens were used for RNA extraction. Each tissue sample, weighing approximately 100 mg, was homogenized in liquid nitrogen using a mortar and pestle; 1 mL of TRIzol (Invitrogen, Carlsbad, CA, USA) was added and the material was collected into a centrifuge tube. RNA was then extracted according to the manufacturer’s instructions.

RNA quality and quantity assessments

RNA quantification was performed using a spectrophotometric NanoDrop instrument (Thermo Fisher Scientific Inc., Waltham, MA, USA), and only RNA samples with an A260/280 ratio greater than 1.8 and an A260/230 close to 2.0 were chosen for gene-expression analysis. The RNA quality and integrity were monitored by the appearance of two clear bands, indicating the presence of 18S and 28S rRNA under ultra-violet light, on 1.2% agarose gels stained with 1% ethidium bromide. RNA samples with serious degradation were discarded.
qRT-PCR

qRT-PCR was performed to analyse H19, SPP1, CHI3L1, INHBA, KRT17, GRB7, STIP1, COL4A2 and PCsLCE mRNA expression levels. In brief, 1 µL of the total RNA prepared as described above was used for cDNA synthesis using QuantiTect Reverse Transcription Kit (Qiagen, Hilden, Germany), generating a 20-µL cDNA solution diluted 5-fold before qRT-PCR. The first-strand cDNA was then synthesized by reverse transcription according to the manufacturer's instructions, followed by qRT-PCR for amplification.

For each reaction, 200-nmol/L forward and reverse primers were used. qRT-PCR was performed using the Applied Biosystem Fast 7500 (Foster City, CA, USA) instrument to detect amplification products in real-time PCR using GoTaq™ qPCR Master Mix (Madison, WI, USA) and all reactions were performed in triplicate 96-well plates (Bioplastics, Landgraaf, Germany). For the nontemplate control, RNase-free water was added instead of cDNA to the cells assigned to each corresponding primer pair to ensure that no nucleic acid contaminations or primer dimers were present. The PCR cycling conditions were as follows: one cycle at 95°C for 10 min, followed by 40 cycles of melting at 95°C for 15 s and annealing/extension at 60°C for 1 min. Fluorescence data were collected after each PCR cycle to generate an amplification plot for determination of the cycle threshold (Ct) value.

The PCR primers used in this study are listed in Table 1. All primers were designed to cross intron–exon boundaries to ensure that PCR products were generated mainly from the cDNA template instead of residual genomic DNA. For all amplification products, the melting curves were examined by measuring the decrease in fluorescence from 95 to 60°C, and a single peak (data not shown) indicated that no primer dimers interfered with the fluorescence detected.

### Table 1. Sequences of the primers used for quantitative real-time polymerase chain reaction

| Marker  | Primer                     | Tm (°C) | Product length (bp) |
|---------|----------------------------|---------|---------------------|
| ACTB    | F: 5'-TTCTTCCCTGGCATGGAGTCCT-3' | 59      | 171                 |
|         | R: 5'-TGCAGGGCTAGTACTGCT-3'    | 60      |                     |
| CHI3L1  | F: 5'-TGGATACCAAGGCAGCAAGA-3'  | 60      | 112                 |
|         | R: 5'-AAGGAGCTGTGACTGATCT-3'   | 62      |                     |
| H19     | F: 5'-TTTGTACGCTCTGCTCTTTGCT-3' | 60      | 131                 |
|         | R: 5'-CAACGGATGCAAAATGACTTAG-3' | 60      |                     |
| INHBA   | F: 5'-TGGAGATCATACGTTGTTG-3'   | 60      | 154                 |
|         | R: 5'-TGACTTGGTCCTGCTGCT-3'    | 62      |                     |
| SPP1    | F: 5'-CAAGAATACACCCAGTGCGT-3'  | 60      | 105                 |
|         | R: 5'-TACATGTTCTTCTCCAGAGA-3'  | 60      |                     |
| KRT17   | F: 5'-ATGGAAGACCCGCTGCTG-3'    | 60      | 109                 |
|         | R: 5'-ACCTAGCGGTGGTCACGGTT-3'  | 60      |                     |
| GRB7    | F: 5'-ATAGCCGCTGCTGCTTTCC-3'   | 61      | 129                 |
|         | R: 5'-GGTCTGCTGATGACAG-3'      | 60      |                     |
| STIP1   | F: 5'-CGAGCAGAATAACCCCTCA-3'   | 60      | 123                 |
|         | R: 5'-CTGGAGATCTCCTCAGATT-3'   | 60      |                     |
| COL4A2  | F: 5'-GGAGAGAAGGGAGACAGAG-3'   | 60      | 119                 |
|         | R: 5'-TCTGGAATCTCCCTTTGCTC-3'  | 60      |                     |
| PCsLCE  | F: 5'-AGGGTTCGCTCACTGCTC-3'    | 63      | 142                 |
|         | R: 5'-AAGACCTCCAGGACACGTCA-3'  | 60      |                     |

OPN, osteopontin; SPP1, secreted phosphoprotein 1; CHI3L1, chitinase-3-like protein 1; INHBA, inhibin beta A; KRT17, keratin 17; GRB7, growth factor receptor-bound protein 7; STIP1, stress-induced phosphoprotein 1; COL4A2, collagen type IV alpha 2; PCsLCE, procollagen C-endopeptidase enhancer; F: forward primer; R: reverse primer. Reference gene: ACTB (beta-actin).

### Statisticlal analysis

Relative quantity (RQ) was used to represent the relative mRNA expression level of the target genes measured by qRT-PCR. RQ represents the level of expression of a target gene (H19, SPP1, CHI3L1, INHBA, KRT17, GRB7, STIP1, COL4A2 and PCsLCE) relative to a reference gene beta-Actin (ACTB), which was calculated according to the following formula: $RQ = 2^{-Ct \text{ reference gene} - Ct \text{ target gene}}$.

IBM SPSS software (version 21.0, Chicago, IL, USA) and GraphPad Prism (La Jolla, CA, USA) were used for statistical analysis using one-way analysis of variance (ANOVA) and Tukey's multiple comparison test to investigate correlations between the mRNA expression level of the target genes and clinicopathological features, including depth of tumor invasion (T), nodal spread (N), distant metastasis (M), TNM classification and histological subtype. All analytical graphs were plotted using GraphPad Prism. A P-value of <0.05 was considered statistically significant.

### Results

#### Patient characteristics

In total, we collected 132 gastric tissues and 104 peri-gastric LNs. After pathological confirmation of the collected specimens, Table 2. Patient characteristics and clinicopathological features of the collected specimen

| Basic characteristic | Patient/gastric tissue | Patient/lymph node |
|----------------------|------------------------|--------------------|
| **Age (years)**      | 60.2 ± 9.2             | 58.4 ± 11.8        |
| **Sex**              |                         |                    |
| Male                 | 47 (71.2%)             | 28 (56.0%)         |
| Female               | 19 (28.8%)             | 22 (44.0%)         |
| **Lauren classification** |                    |                    |
| Intestinal           | 19 (28.8%)             | 9 (18.0%)          |
| Diffuse              | 33 (50.0%)             | 29 (58.0%)         |
| Mixed                | 14 (21.2%)             | 12 (24.0%)         |
| **7th AJCC pathological T category** |                |                    |
| T2                   | 6 (9.1%)               | 3 (6.0%)           |
| T3                   | 34 (51.5%)             | 13 (26.0%)         |
| T4                   | 26 (39.4%)             | 34 (68.0%)         |
| **7th AJCC pathological N category** | |                    |
| N0                   | 14 (21.2%)             | 0 (0.0%)           |
| N1                   | 10 (15.2%)             | 0 (0.0%)           |
| N2                   | 16 (24.2%)             | 11 (22.0%)         |
| N3a                  | 13 (19.7%)             | 20 (40.0%)         |
| N3b                  | 13 (19.7%)             | 19 (38.0%)         |
| **7th AJCC pathological M category** | |                    |
| M0                   | 61 (92.4%)             | 43 (86.0%)         |
| M1                   | 5 (7.6%)               | 7 (14.0%)          |
| **7th AJCC pathological TNM classification** | |                    |
| IIA                  | 11 (16.7%)             | 1 (2.0%)           |
| IIB                  | 9 (13.6%)              | 0 (0.0%)           |
| IIIA                 | 20 (30.3%)             | 5 (10.0%)          |
| IIIB                 | 10 (15.1%)             | 14 (28.0%)         |
| IIC                  | 11 (16.7%)             | 24 (48.0%)         |
| IV                   | 5 (7.6%)               | 6 (12.0%)          |

*Except for age, other values are presented as the number of patients followed by the percentage in parentheses.

AJCC, American Joint Committee on Cancer; TNM, tumor-node-metastasis.

Note: The clinicopathological characteristics of patients with negative lymph nodes (LNs) are identical to those with positive LNs because at least one positive LN and one negative LN were collected from each patient, for which when suspicious LNs (macroscopic estimation of LN status was difficult) we retrieved one or two additional LNs.
we amassed 66 pairs of fresh primary gastric-cancer tissues and their matched adjacent normal gastric tissues (>5 cm away from tumor), 50 metastatic LNs and 54 non-metastatic LNs. The clinicopathological information of the patients is summarized in Table 2.

Expression levels of the target genes in gastric tissues

Preliminary tests analysing the mRNA expression levels of the nine investigated genes were first performed. We found that the expression of five genes, namely KRT17, GRB7, STIP1, COL4A2 and PCOLCE, demonstrated no significant difference in expression between the cancerous samples and their matched normal gastric tissues (Supplementary Figure 1). These genes were therefore excluded from further investigation.

Next, qRT-PCR was performed to assess the mRNA expression levels of H19, CHI3L1, INHBA and SPP1 in 66 pairs of gastric tissues. The expression level of H19 showed no significant difference between normal and cancer tissues ($P = 0.079$; Figure 1A). On the other hand, despite CHI3L1,
demonstrating high expression in gastric-cancer tissues ($P = 0.002$; Figure 1B), INHBA and SPP1 were the main genes to show consistent significant statistical association between gastric-cancer tissues and normal gastric mucosa ($P < 0.001$; Figure 1C and D, respectively) and were therefore used for further investigation.

The mean expression level of INHBA in tumor tissues was 15.4-fold higher than that in normal tissues, with an area under the receiver operating characteristic (ROC) curve of 0.974, indicating high accuracy of the test (Figure 2A). Based on the INHBA mRNA level indicated by the ROC curve, the cut-off RQ for cancer detection was set to 0.001 and the sensitivity and specificity for the detection of gastric cancer were found to be 91.3 and 95.7%, respectively.

Regarding SPP1, it similarly had a high mean mRNA expression level in gastric-cancer tissues, which was 15.6-fold higher...
Figure 4. Association between SPP1 mRNA expression level and clinicopathological features in gastric-cancer tissues. SPP1 expression level was compared with (A) tumor depth/T category (normal vs T3 and T4, \( P \leq 0.026 \) and \( 0.002 \), respectively); (B) nodal status/N category (normal vs N3a and N3b, \( P = 0.031 \) and \( 0.040 \), respectively); (C) distant metastasis/M category (normal vs M0, \( P < 0.001 \)); (D) TNM classification (normal vs IIC, \( P = 0.029 \)); and (E) Lauren classification (normal vs intestinal type, \( P = 0.008 \)). The graphs illustrate the mean relative quantity (RQ) value of each group; * \( P < 0.05 \); ** \( P < 0.01 \); *** \( P < 0.001 \). Columns that did not demonstrate statistical correlation between cancer and normal tissues are not labeled. SPP1, secreted phosphoprotein 1.

Figure 5. mRNA level of INHBA and SPP1 in 54 LN− and 50 LN+ samples. (A) INHBA can be seen to be significantly up-regulated in LN+ (\( P < 0.001 \)), whereas (B) no significant difference for SPP1 were found (\( P = 0.470 \)). The graphs illustrate the mean relative quantity (RQ) value of each group; LN−, non-metastatic lymph nodes; L+, metastatic lymph nodes; n.s., no significance; *** \( P < 0.001 \). INHBA, inhibin beta A; SPP1, secreted phosphoprotein 1; LN, lymph node.
than that in normal gastric tissues, with an area under the ROC curve of 0.924 (Figure 2B). The cut-off RQ for cancer detection as indicated by the ROC curve was set to 0.001 and therefore showed a sensitivity of 82.6% and a specificity of 87.0%.

The accuracy of the combined expression levels of these two biomarkers for the diagnosis of gastric cancer was also investigated. Considering that samples with an expression level of either of the two genes above or expression levels of both genes below the cut-off RQ were considered to be positive and negative for cancer detection, the sensitivity and specificity were found to be 92.3 and 84.6%, respectively.

**Association between mRNA expression and clinicopathological features in gastric tissues**

The mean expression of INHBA was increased with the increase in invasive tumor depth in gastric tissues; the expression of INHBA was significantly lower in normal gastric tissues than in T3 and T4 gastric-cancer tissues (P < 0.001; Figure 3A), but not in T2 gastric-cancer tissues (P = 0.447). In addition, the expression level of INHBA in gastric-cancer tissues was significantly associated with nodal status (Figure 3B), distant metastasis (Figure 3C), TNM classification (except for stage IV; Figure 3D) and Lauren classification (Figure 3E).

The RQ of mean SPP1 expression was increased with the increase in tumor depth (normal vs T3, P = 0.026 and normal vs T4, P = 0.002; Figure 4A); however, SPP1 was less consistently overexpressed between the different subgroups of N, M, TNM classification and Lauren subtypes as compared to INHBA (Figure 4B–E).

**Expression level of INHBA and SPP1 in peri-gastric LNs**

The INHBA expression level was significantly up-regulated at an average of 4.1-fold in metastatic LNs (P < 0.001; Figure 5A), with an area under the ROC curve of 0.857 (Figure 6), whereas no difference in the expression level of SPP1 was found between positive and negative LNs (P = 0.470; Figure 5B). Further, the sensitivity and specificity of INHBA mRNA for differentiating between metastatic and non-metastatic LNs were found to be 80.0 and 81.5%, respectively, when the RQ of INHBA expression was 0.0007.

**Association between mRNA expression of INHBA and clinicopathological features in peri-gastric LNs**

Comparing the expression levels of INHBA between metastatic and non-metastatic LNs, we found that INHBA expression was positively associated with tumor depth/T category (LN– vs T4, P < 0.001; T3 vs T4, P = 0.003), nodal status/N category (LN– vs N2 and N3b, P = 0.001 and 0.042, respectively), distant metastasis/M category (LN– vs M0 and M1, P = 0.014 and 0.001, respectively; M0 vs M1, P = 0.043), TNM classification (LN– vs IIIB and IIIC, P = 0.033 and 0.040, respectively) and Lauren classification (LN– vs intestinal type and mixed variant, P < 0.001, and 0.020, respectively; intestinal vs diffuse, P = 0.019) (Figure 7A–E), except for N3a, stage IIIA and IV. Of note, because of the limited number of cases, we were unable to demonstrate the association of INHBA expression with T2 (n = 3), N1 (n = 0), IIA (n = 1) and IIIB (n = 0).

**Discussion**

In this study, we have demonstrated that the genes KRT17, GRB7, STIP1, COL4A2 and PCOLCE were not consistently highly expressed in our sets of gastric-cancer tissues and that the expression levels of both H19 and CHI3L1 were lower than those of SPP1 and INHBA. In addition, the mRNA levels of INHBA and SPP1 were significantly higher in gastric-cancer tissues than in non-tumorous tissues (both P < 0.001) and demonstrated high sensitivity of 91.3 and 82.6%, respectively, and specificity of 95.7 and 87.0%, respectively. As a combination, the sensitivity and specificity of INHBA and SPP1 reached 92.3 and 84.6%, respectively, for the diagnosis of gastric cancer. Many publications regarding SPP1 and INHBA mainly associated their expression levels in serum with prognosis [21–24], but only a few studies examined the expression of SPP1 or INHBA in gastric-cancer tissues [22, 25]. To our knowledge, there is no study that reported the expressions of both SPP1 and INHBA and their sensitivity and specificity for the diagnosis of gastric cancer and metastatic LNs.

Further analysis demonstrated that the mRNA expression level of INHBA was superior to that of SPP1 for diagnosing metastatic LNs (P < 0.001). This may be due to the existing different splice variants of SPP1, since different variants have been reported to have different clinicopathological and biological functions in gastric cancer [26]. In addition, the expression level of INHBA may reveal potential LN metastasis with high sensitivity and specificity (80.0 and 81.5%, respectively) and differentiate between different disease conditions, e.g. T3 and T4 (P = 0.003), M0 and M1 (P = 0.043), and different histological subtypes (intestinal type vs diffuse type, P = 0.019). Based on these results, we identified INHBA as a promising biomarker for the diagnosis of LN metastasis in gastric cancer.

The clinical significances of our findings in the present study are as follows. First, detecting the expressions of molecular biomarkers before surgery may optimize the diagnosis of inconclusive biopsies, especially regarding small, non-characteristic gastric-cancer lesions (dual applicability of SPP1 and INHBA demonstrated high sensitivity and specificity) and abnormally enlarged suspicious LNs (INHBA had superior sensitivity and specificity), as recent qRT-PCR technology has been shown to be superior to routine pathological examination at revealing minor tumor deposits because it can analyze a larger number of tissue samples with greater sensitivity and specificity [27]. For such cases, molecular diagnosis may help to select the better...
treatment between surveillance, endoscopic mucosal/submucosal resection and even radical gastrectomy [28–30]. Second, detecting INHBA expression during surgery may yield more accurate frozen-section diagnosis [31] and therefore select the superior modality between radical gastrectomy and the extent of lymphadenectomy, such as for uncertain enlarged LNs beyond the scope D2 resection. This is because the analysis of frozen sections during operation has some inherent problems, since it uses less than 1% of tissues in a node for analysis and therefore may result in comparatively low to moderate diagnostic sensitivity (52.2%), specificity (88.8%) and overall accuracy (73.8%) [32]. In addition, the use of qRT-PCR was once limited because it takes a long time to obtain the results of qRT-PCR. However, recent advances including quicker temperature change and faster enzymes without sacrificing accuracy have enabled more efficient analysis (results obtained within 30–60 min) [33–35]. As such, Ferris et al. [8] have demonstrated that, for an average time of 35 min, rapid automated qRT-PCR was able to detect LNs metastasis with higher accuracy than intra-operative pathological examination.

Third, determining a threshold level of INHBA overexpression might facilitate differentiating between early-stage (stage IA cases, for which adjuvant therapy might not be necessary), middle-stage (stages IB to IIIA, which may not require neo-adjuvant treatment) and advanced (stages IIIB–IIIC) gastric cancer, thereby helping to recognize cases for which more aggressive therapies would be most beneficial. Thus, molecular biomarkers

![Figure 7. Association between INHBA mRNA expression level and clinico-pathological features with peri-gastric lymph nodes. The INHBA expression level was compared with (A) tumor depth/T category (LN– vs T4, \( P < 0.001; T3 vs T4, P = 0.003\)); (B) nodal status/N category (LN– vs N2 and N3b, \( P = 0.001 \) and 0.042, respectively); (C) distant metastasis/M category (LN– vs M0 and M1, \( P = 0.014 \) and 0.001, respectively; M0 vs M1, \( P = 0.043 \)); (D) TNM classification (LN– vs IIIB and IIIC, \( P = 0.033 \) and 0.019, respectively); and (E) Lauren classification (LN– vs intestinal type and mixed variant, \( P < 0.001 \) and 0.010, respectively; intestinal vs diffuse, \( P = 0.019 \)). The graphs illustrate the mean relative quantity (RQ) value of each group. LN–, non-metastatic lymph nodes; LN+, metastatic lymph nodes; *\( P < 0.05 \); **\( P < 0.01 \); ***\( P < 0.001 \). Columns that did not demonstrate statistical correlation between tumoral and normal tissues are not labeled. INHBA, inhibin beta A; LN, lymph node.]
such as INHBA can be regarded as potential biomarkers for accelerating the leap in surgical oncology progress.

Despite the novel findings of the present study, there are some limitations that should be mentioned. First, the limited sample size may have resulted in the lack of statistical associations of SPP1 and INHBA with certain clinicopathological features of gastric-cancer tissues and nodes. However, the sample size can be explained as follows. It was relatively cumbersome to macroscopically differentiate between large metastatic and non-metastatic nodes, considering that only a few (only one or two) LNs per patient could be collected from the overall retrieved nodes for the following reasons: (i) only enlarged LNs (at least 1 cm in diameter) were used to increase the diagnostic accuracy and reliability between each pair of halves sent for pathology and collected for this study and (ii) collection of a greater number of LNs would have resulted in additional financial burden on the patients. Second, we mainly enrolled cancer patients with a tumor size of at least 3 cm in diameter, and thus no data for category T1 patients could be retrieved and analysed. Third, since the tissue samples had been recently collected, the associations of INHBA and SPP1 with overall survival and cancer recurrence could not be analysed.

Conclusions

Our present study showed that both SPP1 and INHBA are reliable biomarkers for the diagnosis of gastric cancer. The findings also demonstrated that INHBA has a high accuracy for diagnosing metastatic LNs and may thus be considered a promising biomarker.

Supplementary Data

Supplementary data is available at Gastroenterology Report online.

Funding

This work was supported by the Natural Science Foundation of Guangdong Province (Grant number 2015A030313089, 2018A030313631), Guangzhou University-Institute-Industry Collaborative Innovation Major Projects (Grant number 201508030042, 201604020038), Center for Nasopharyngeal Carcinoma Research, Hong Kong (Grant number AoEM-06/08) and Shenzhen Dept. of Science and Information (Grant number JCYJ20130329110752138).

Author contributions

C.L., S.R.S. and Z.W.Z. conceived of and designed this study. Z.W.Z. and C.L. provided administrative support. Z.W.Z. and C.L. provided study materials or patients. S.R.S., W.Y.C., J.W., T.W. and J.Q.L. performed qRT-PCR experiments. S.R.S., C.F., W.W. and Z.M.L. collected and assembled data. S.R.S., C.F. and W.W. performed data analysis and interpretation. All authors contributed to the writing and final approval of the manuscript.

Conflict of interest statement: none declared.

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