Predictive Factors for Inaccurate Diagnosis of Swollen Lymph Nodes in Endoscopic Ultrasound-Guided Fine Needle Aspiration

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Background/Aims: This study aimed to identify the predictive factors for inaccurate endoscopic ultrasound-guided fine needle aspiration (EUS-FNA) diagnosis of swollen lymph nodes without rapid on-site cytopathological evaluation.

Methods: Eighty-three consecutive patients who underwent EUS-FNA for abdominal or mediastinal lymph nodes from January 2008 to June 2017 were included from a prospectively maintained EUS-FNA database and retrospectively reviewed. The sensitivity, specificity, and accuracy of EUS-FNA for the detection of neoplastic diseases were calculated. Candidate factors for inaccurate diagnosis (lymph node size and location, needle type, puncture route, number of passes, and causative disease) were evaluated by comparison between accurately diagnosed cases and others.

Results: The final diagnosis of the punctured lymph node was classified as neoplastic (65 cases: a metastatic lymph node, malignant lymphoma, or Crow-Fukase syndrome) or non-neoplastic (18 cases: a reactive node or amyloidosis). The sensitivity, specificity, and accuracy were 83%, 94%, and 86%, respectively. On multivariate analyses, small size of the lymph node was the sole predictive factor for inaccurate EUS-FNA diagnosis with a significant difference (odds ratios, 19.8; 95% confidence intervals, 3.15–124; \( p = 0.0015 \)).

Conclusions: The lymph node size of <16 mm was the only independent factor associated with inaccurate EUS-FNA diagnosis of swollen lymph nodes.

Key Words: Lymphadenopathy; Accuracy; Endosonography; Risk factors; Lymphoma

INTRODUCTION

Since endoscopic ultrasound-guided fine needle aspiration (EUS-FNA) for a pancreatic tumor was first reported in 1992,¹ it has been applied to many targets, such as subepithelial lesions of the gastrointestinal tract, mediastinal masses, biliary tumors, and ascites.²⁻²² EUS-FNA can also be useful for diagnosing internal lymphadenopathy, which often requires correct diagnosis to confirm the cancer stage, recurrence of surgically resected malignancy, and non-neoplastic lymph nodes. Although several studies have reported the usefulness of EUS-FNA,¹³⁻¹⁹ its sensitivity for lymphadenopathy seems relatively lower than that for other targets, such as pancreatic masses.²⁰⁻²²

The diagnostic accuracy of EUS-FNA can vary due to a few factors, including the target size and location, needle type, puncture route, number of needle passes, and causative disease.²³ Although rapid on-site evaluation (ROSE) during EUS-FNA has been reported to be associated with improved diagnostic yield,²⁴⁻²⁶ a recent randomized study has shown conflicting results regarding the impact of ROSE.²⁷ Moreover, that type of evaluation may be difficult to implement in most institutions due to the lack of a cytopathologist who is always available to perform EUS-FNA. If predictive factors for inaccurate results using EUS-FNA without ROSE can be iden-
tified, outcomes may improve with some countermeasures without on-site evaluation. Therefore, we sought to investigate these predictive factors in this study.

**PATIENTS AND METHODS**

**Patients**

All patients who underwent EUS-FNA for swollen lymph nodes, which were detected in cross-sectional imaging examinations, at Sendai City Medical Center between January 2008 and June 2017 were extracted from the prospectively maintained EUS-FNA database and retrospectively reviewed. A swollen lymph node was defined as a node with a long axis size of at least 8 mm. Lymph-node EUS-FNA was performed in cases of swollen nodes with unknown etiology, those for which a determination of cancer stage was necessary, those with the possibility of cancer recurrence after surgical treatment, and those where the original cancer was difficult to puncture. Patients without sufficient procedural records and those without definitive final diagnoses because of a lack of follow-up were excluded. There were no patients aged <18 years and no occurrences of untreated coagulopathy or pregnancy during the study period. All patients provided written informed consent to undergo EUS-FNA. This study was approved by the institutional review board of our hospital.

**EUS-FNA procedures**

EUS-FNA was performed under conscious or deep sedation with intravenous administration of midazolam, pentazocine, and/or propofol. Prophylactic antimicrobial agents were not administered. All patients were hospitalized to monitor possible adverse events for at least 1 night. After a swollen lymph node was visualized with a convex-arrayed echoendoscope (GF-UCT260, GF-UC240P; Olympus Co., Tokyo, Japan), an appropriate puncture route without intervening vessels was identified using the color Doppler mode. A 19-, 22-, or 25-G needle (EchoTip Ultra; Cook Japan Inc., Tokyo, Japan, EchoTip ProCore; Cook Japan Inc., Expect; Boston Scientific Japan K.K., Tokyo, Japan, EzShot 3 Plus; Olympus Co., SonoTip Pro Control; Medico’s Hirata Inc., Tokyo, Japan, or Acquire; Boston Scientific Japan K.K.) was inserted through the accessory channel of the scope and advanced into the target node. After the stylet was pulled out, 15 to 30 back-and-forth movements of the needle were performed by suction with a 10- or 20-mL syringe. The needle was then withdrawn. The aspirated material was expelled into a receptacle with formalin solution using saline flushing. The puncture procedures were repeated until macroscopically visible whitish material was obtained but were limited to a maximum of 5 passes if no material was obtained. ROSE was not conducted for any patient. All procedures were carried out by an expert endosonographer (KI, SK, YK and TO) who had performed >500 EUS examinations and >50 EUS-FNA procedures or by a trainee supervised by experts. The needle type and size were determined at the endosonographer’s discretion. Adverse events related to EUS-FNA were defined as newly emerged complications that required prolongation of the hospitalization or therapeutic intervention. All EUS-FNA samples were prepared using the cell block method for microscopic evaluation. Pathological evaluation was performed using hematoxylin–eosin staining, followed by additional immunochemistry and monoclonal antibody staining if desired.

**Outcome measurements and definitions**

The main outcome measurement of this study was the accuracy rate of EUS-FNA for swollen lymph nodes. Accuracy was defined as the precise diagnosis of neoplasm and non-neoplasm of the punctured node. Final diagnosis was classified into neoplastic and non-neoplastic lymph nodes, and the sensitivity and specificity for the detection of neoplastic lymph nodes were estimated. Neoplastic diseases included metastatic lymph nodes and hematopoietic/lymphoid neoplasms.

The final diagnosis was confirmed by pathological evaluation of a resected specimen when surgery was performed. For patients who did not undergo surgery, the diagnosis was defined as malignancy when pathological evaluation, such as EUS-FNA for the originating cancer and ascitic/pancreatic-juice cytology, confirmed the malignancy and the lymph node was larger on follow-up imaging evaluations. In patients without such pathological confirmation, the final diagnosis was defined as malignancy when the clinical course was sufficiently consistent, namely, when there were obvious findings in the imaging evaluation or when a patient’s condition continued to deteriorate, or they responded to antineoplastic treatment during the follow-up period of >12 months. The diagnosis was defined as non-neoplastic when the swollen lymph nodes regressed or disappeared during follow-up evaluation or were stable for >12 months.

The technical success of EUS-FNA was defined as the successful acquisition of macroscopically visible whitish material. The size and location of the punctured lymph node, needle type, needle diameter, puncture route, number of needle passes, and final diagnosis were evaluated to analyze the association between each factor and diagnostic accuracy using univariate and multivariate analyses.

**Statistical analysis**

The sensitivity, specificity, and accuracy rate of EUS-FNA
for detecting neoplastic diseases were calculated with 95% confidence intervals (CI).

The above-mentioned candidate factors were compared between cases with accurate diagnosis and others. Analyses of predictive factors were performed using the Chi-squared test. Continuous variables were divided into 2 groups by using a cut-off value defined with the receiver operating characteristic (ROC) curve for multivariate analysis. Factors with a p-value <0.20 in the univariate analyses were evaluated in the multiple logistic regression model, and odds ratios (OR) with 95% CI were calculated. A two-tailed p-value <0.05 was considered statistically significant. The analyses were performed using JMP Pro13 for Mac (JMP 13; SAS Institute Inc., Cary, NC, USA).

RESULTS

Patients’ characteristics

After elimination of 6 patients without final diagnosis, 83 patients (male, 47; median age, 68 years [range, 22–88 years]) were included in this study. The final diagnosis was classified as neoplastic lymphadenopathy in 65 patients and non-neoplastic lymphadenopathy in 18 patients (Table 1). The origin of the metastatic lymph nodes was extrahepatic bile duct cancer in 17 patients, pancreatic cancer in 14, neuroendocrine tumor in 8, intrahepatic cholangiocarcinoma in 3, hepatocellular carcinoma in 2, hepatocellular carcinoma in 2, colonic cancer in 1, duodenal cancer in 1, and an unknown cancer in 4. Details of the EUS-FNA procedures are shown in Table 2. Macroscopically visible material was obtained in all cases. No adverse events related to the procedure were observed.

Table 1. Characteristics of the Patients and Lesions

| Parameter                              | Number     |
|----------------------------------------|------------|
| Age, median (range), yr                | 68 (22–88) |
| Sex, male, n (%)                       | 47 (57)    |
| Diameter of lymph node, median (range), mm | 20 (8–56) |
| Location of lymph node, n (%)          |            |
| Peripancreatic and porta hepatis       | 48 (58)    |
| Para-aorta                             | 29 (35)    |
| Mediastinum                             | 6 (7)      |
| Final diagnosis, n (%)                 |            |
| Metastasis                             | 54 (65)    |
| Malignant lymphoma                     | 10 (12)    |
| Crow-Fukase syndrome                   | 1 (1.2)    |
| Reactive change                        | 17 (21)    |
| Amyloidosis                            | 1 (1.2)    |

Diagnostic yields of EUS-FNA

The obtained materials were adequate for pathological evaluation in 93% (77/83) of the cases. Additional immunohistochemical staining was performed in 60/83 of the patients. Fifty-five of the specimens were pathologically determined to be neoplastic, while 22 were non-neoplastic. The EUS-FNA diagnosis accurately concurred with the final diagnosis in 71 cases. Of the 6 cases with inaccurate diagnosis, the EUS-FNA results were false-positive in 1 and false-negative in 5. In the single false-positive case, atypical lymphocytes obtained in EUS-FNA were positive for both CD20 and leukocyte common antigen, resulting in incorrect diagnosis as malignant lymphoma. However, the swollen lymph nodes, up to 15 mm in size, were obviously regressed to an undetectable level during follow-up without treatment, and the final diagnosis was judged as non-neoplastic reactive change. The 5 cases with false-negative EUS-FNA results can be described as follows. Two false-negative cases were diagnosed as having malignant lymphoma by bone marrow aspiration cytology. One case of pancreatic cancer was diagnosed by using EUS-FNA on the main tumor. One case of duodenal cancer was diagnosed by imaging examinations and clinical follow-up. Finally, 1 case of Crow-Fukase syndrome was diagnosed by evaluating surgically sampled lymph nodes. The EUS-FNA specimens of these false-negative cases contained several cell aggregations, which

Table 2. Details of Endoscopic Ultrasound-Guided Fine Needle Aspiration Procedures

| Needle, n (%) | Expect | EzShot3 Plus | EchoTip | Acquire | SonoTip | Unknown |
|---------------|--------|--------------|---------|---------|---------|---------|
|               | 44 (53)| 8 (9.6)      | 6 (7.2) | 5 (6.0) | 2 (2.4) | 18 (22) |
| Needle size, n (%) |        |              |         |         |         |         |
| 19 G           | 8 (10) |              |         |         |         |         |
| 22 G           | 50 (60)|              |         |         |         |         |
| 25 G           | 15 (18)|              |         |         |         |         |
| Unknown        | 10 (12)|              |         |         |         |         |
| Puncture route, n (%) |     |              |         |         |         |         |
| Transesophageal| 6 (7.2)|              |         |         |         |         |
| Transgastric   | 44 (53)|              |         |         |         |         |
| Transduodenal  | 33 (40)|              |         |         |         |         |
| No. of needle passes, median (range) |     |              |         |         |         |         |
| 3 (1–5)        |       |              |         |         |         |         |
| Technical success, n (%) |     |              |         |         |         |         |
| 83 (100)       |       |              |         |         |         |         |
| Adverse events, n (%) |     |              |         |         |         |         |
| 0 (0)          |       |              |         |         |         |         |
did not show malignant findings. In 6 other cases, EUS-FNA samples did not contain a sufficient number of neoplastic cells despite evaluating the macroscopically detectable materials on site. The final diagnoses were as follows: biliary tract cancer in 3 cases, pancreatic cancer in 2, and neuroendocrine tumor in 1.

In all the 83 cases evaluated, the overall sensitivity, specificity, and accuracy rate for the detection of a neoplastic lymph node were 83%, 94%, and 86%, respectively (Table 3).

**Predictive factors for inaccurate diagnosis**

On univariate analyses comparing accurately diagnosed cases and others, the size of the punctured lymph node was the sole predictive factor with a significant difference (median [interquartile range], 20 [16–25] mm vs. 14 [12–15] mm, \( p < 0.0001 \)) (Table 4).

| Table 3. Diagnostic Yields of Endoscopic Ultrasound-Guided Fine Needle Aspiration for Lymph Nodes |
|---|---|---|
| **Sensitivity** | **83%** | **78–84** |
| **Specificity** | **94%** | **77–99** |
| **Accuracy** | **86%** | **78–88** |
| **CI**, confidence intervals. |

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**Table 4. Univariate Analysis for Predictive Factors Associated with Diagnostic Accuracy of Endoscopic Ultrasound-Guided Fine Needle Aspiration**

| Predictive Factor | Accurate diagnosis | Inaccurate diagnosis | \( p \)-value |
|---|---|---|---|
| Lymph node diameter (long axis) | | | <0.0001 |
| \( \geq 16 \) mm | 54 | 2 | |
| <16 mm | 17 | 10 | |
| Location of lymph node | | | 0.16 |
| Peripancreatic and porta hepatis | 44 | 4 | |
| Para-aorta | 22 | 7 | |
| Mediastinum | 5 | 1 | |
| Needle | | | 0.53 |
| Expect | 36 | 8 | |
| EzShot3 Plus | 8 | 0 | |
| EchoTip | 5 | 1 | |
| Acquire | 5 | 0 | |
| SonoTip | 2 | 0 | |
| Needle size | | | 0.83 |
| 19 G | 7 | 1 | |
| 22 G | 43 | 7 | |
| 25 G | 12 | 3 | |
| Puncture route | | | 0.53 |
| Transesophageal | 5 | 1 | |
| Transgastric | 36 | 8 | |
| Transduodenal | 30 | 3 | |
| No. of needle passes | | | 0.14 |
| \( \geq 4 \) | 12 | 1 | |
| <4 | 59 | 11 | |
| Final diagnosis | | | 0.06 |
| Metastasis | 46 | 8 | |
| ML | 8 | 2 | |
| Crow-Fukase syndrome | 0 | 1 | |
| Benign | 17 | 1 | |
| ML, malignant lymphoma. |
In the ROC curve, the sensitivity and specificity were 76% and 83%, respectively, with a cut-off size of 16 mm (Fig. 1). The area under the receiver operator curve (AUC) was 0.76.

Multiple logistic regression showed that the lymph node size of <16 mm was the significant predictive factor for inaccuracy in EUS-FNA diagnoses (OR, 19.8; 95% CI, 3.15–124; \( p = 0.0015 \)) (Table 5).

**DISCUSSION**

Accurate diagnosis of neoplastic lymph nodes using EUS-FNA appears to be relatively difficult in comparison with that of other neoplasms, such as pancreatic cancer, according to the results of previously reported studies\(^ {13-22} \) and the present study. If the predictive factors for unfavorable results of EUS-FNA can be identified, diagnostic performance may improve.

Hence, many previous studies have attempted to identify such factors.\(^ {26,27,29-32} \)

Some studies have reported that on-site evaluation of the obtained samples by a cytopathologist improves the diagnostic yield of EUS-FNA for lymph nodes as well as pancreatic tumors,\(^ {24-26,13-32} \) whereas Kappelle et al. recently reported that ROSE during EUS-FNA of lymph nodes did not improve the diagnostic yield in a multicenter, randomized trial.\(^ {27} \) In our study, the 6 cases (7.2%) without adequate specimens might have been successfully diagnosed via on-site evaluation. However, on-site evaluation is not available in most institutions due to limitations of human resources and costs.

The number of needle passes and the needle size are procedural factors that are associated with outcomes. When on-site evaluation is unavailable, the appropriate number of passes is always in question. Leblanc et al.\(^ {29} \) have reported that the number of needle passes (≥5) is the most significant influential factor on the diagnostic yields of EUS-FNA for lymph nodes without on-site cytopathologists. However, according to a randomized study by Wallace et al.,\(^ {30} \) more than 3 passes are adequate. Pellisé et al.\(^ {31} \) have demonstrated that efficacy of needle pass repetition reaches a plateau at the third pass in lymph node EUS-FNA. Clearly, the optimal number of needle passes remains controversial. Although the reason why the number of needle passes was not a factor associated with accuracy in our study is unknown; for most of the patients (53/83), the number of needle passes was 3 or more, i.e., there might be a beta error.

On the contrary, in another study, a larger needle size was found to improve tissue acquisition in lymph node EUS-FNA.\(^ {13} \) However, there have been no comparative studies in which the needle size was a significant predictive factor.\(^ {26,32-36} \)

It seems particularly difficult to diagnose lymphoproliferative disorders with EUS-FNA specimens. Some studies have reported high diagnostic yields using flow cytometry for malignant lymphoma.\(^ {13,14,37} \) Although most lymphoproliferative diseases (malignant lymphoma and Crow-Fukase syndrome) were accurately diagnosed in our study, selective use of flow cytometry could improve diagnostic outcomes.

This is the first study in which the small size of the punctured lymph node was a predictive factor for inaccurate

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**Fig. 1.** The receiver operating characteristic (ROC) curve showing the relationship between lymph node size and the diagnostic yields of endoscopic ultrasound-guided fine needle aspiration for lymph nodes. Area under the ROC curve=0.76.

**Table 5.** Logistic Regression Analysis for Predictive Factors Associated with Diagnostic Accuracy of Endoscopic Ultrasound-Guided Fine Needle Aspiration

|                                        | OR (95% CI)   | \( p \)-value |
|----------------------------------------|---------------|---------------|
| Lymph node diameter (≥16 mm)           | 19.8 (3.15–124) | 0.0015        |
| Location of lymph node                 | 2.48 (0.49–12.6) | 0.27          |
| No. of needle passes                    | 2.54 (0.18–35.7) | 0.49          |
| Final diagnosis                         | 5.08 (0.48–53.3) | 0.17          |

CI, confidence intervals; OR, odds ratios.
diagnoses. Metastatic cells in small lymph nodes, which are in an earlier period of metastasis, are thought to settle in a subcapsular sinus located in the marginal part of nodes. However, it is extremely difficult to arbitrarily puncture a marginal part of such small nodes. Therefore, selecting larger nodes among multiple swollen ones may improve outcomes. In patients without lymph nodes larger than 16 mm, certain countermeasures, such as increasing needle passes, may be effective. Moreover, puncturing nodes with metastatic findings may improve the diagnostic outcomes. Metastatic lymph nodes have been reported to be round, larger than 10 mm, hypoechoic, and well-demarcated. In addition, other studies have reported the usefulness of elastography and contrast enhancement in the harmonic mode to distinguish neoplastic from non-neoplastic nodes.

In addition, small lymph nodes can be reactively enlarged even if a neoplastic disease exists in another part of the body. The EUS-FNA results in this situation may have been false-negative in the present study due to our definition of a final diagnosis. Theoretically, the real accuracy could be revealed only in studies wherein the EUS-FNA results of the punctured lymph node are compared with the histological results of surgical sampling of the same node. However, such studies are extremely difficult to carry out because of ethical concerns. Performing surgery merely to sample lymph nodes should be avoided in patients with unresectable cancers due to metastatic lymph nodes or lymphoproliferative diseases. Instead, sufficient clinical follow-up for >12 months should be avoided in patients with unresectable cancers due to metastatic lymph nodes or lymphoproliferative diseases. More specifically, metastatic lymph nodes in patients with unresectable cancers due to metastatic lymph nodes or lymphoproliferative diseases with a lack of needle information in this study, this could be clarified in other studies wherein many different types of needles are not used. (4) The AUC value of 0.76 is relatively low. (5) Inadequate specimens led to 50% of the inaccurate diagnoses in this study. In conclusion, the smaller size of the lymph node (<16 mm) was the only independent predictive factor associated with inaccurate EUS-FNA diagnosis of swollen lymph nodes without on-site evaluation. Future prospective studies with a large sample size are required to confirm these results.

Conflicts of Interest
The authors have no financial conflicts of interest.

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