Qualitative role of endobronchial elastography with endobronchial ultrasound in differentiating malignant and benign lesions: a retrospective single-center study from India
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Introduction
Endobronchial ultrasound (EBUS) has gained a lot of importance in recent years and has been an integral part of the diagnostic armamentarium in the evaluation of mediastinal lymphadenopathy. It has expanded its scope from a mere diagnostic procedure to even therapeutic lymph node (LN) decompressions and implanting fiducial markers [1,2]. Lung cancer management requires careful mediastinal LN evaluation. EBUS and endoscopic ultrasound are complementary methods as in combination virtually all mediastinal and hilar nodal stations can be visualized and sampled. Selection of LNs to be sampled is a crucial step, affecting diagnostic accuracy and preventing undue sampling.

Elastography provided to the new generation EBUS scopes have enabled mapping of the elasticity of LNs in the form of colors and strain ratio leading to better guidance while taking samples [3].

Elastography is an ultrasonography-associated technology that measures tissue compressibility.

Endobronchial elastography can be performed in real time and the elastic responses to external compression may be converted to a color format of red, green, and blue images to allow for qualitative assessment of the soft tissue concerned.

We aim to discuss the potential role of endobronchial elastography in identifying LN features for better characterization of LNs and subsequently better selection of biopsies.

Patients and methods
This study was performed retrospectively on consecutive patients who were admitted for EBUS-transbronchial needle aspiration (TBNA) at our department for a period of 6 months. Retrospective study done, all necessary consents obtained. The

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elastography evaluation was done in addition of the EBUS-TBNA for clinical indications independent of the purpose of the study. EBUS-TBNA was done for patients with mediastinal lymphadenopathy, taken as an enlargement of LNs of more than or equal to 1 cm in short-axis diameter on the computed tomography chest. The patients gave written informed consent before the EBUS-TBNA and to use records for the study. Patient confidentiality was maintained throughout the study.

All procedures were executed by the same physician. The procedure was performed on conscious sedation with midazolam and fentanyl with the addition of local lidocaine sprays. The EBUS convex probe (CP-EBUS, BF-UC260FW; Olympus, Tokyo, Japan) was introduced through the oral route, with spray-as-you-go instillation of 2 ml aliquot doses of 2% lidocaine.

Elastography was performed on mediastinal LNs that were seen to be candidates for EBUS-TBNA. The colors correlated with soft, firm, and hard tissues were yellow/red, green, and blue, respectively. The B-mode images and elastography images were displayed simultaneously on the screen alongside. Elastography evaluation was done on every node sampled. The predominant color was documented as interpreted by two independent bronchoscopists after the images were obtained. Subsequently, classification of the EBUS elastography images was done using the classification protocol used by Izumo et al. [4].

Type 1 group: largely nonblue (green, yellow, and red). Type 2 group: part blue, part nonblue (green, yellow, and red). Type 3 group: largely blue.

The final result was confirmed from the histopathological examination of EBUS-TBNA specimens made as cell blocks by pathologists who were not aware of the results of EBUS elastography. Subsequently, elastography color patterns were correlated with biopsy results.

Statistical analysis
The statistical analysis was carried out using SPSS software (18.0; SPSS Inc., Chicago, Illinois, USA). The statistical significance between groups was decided by $\chi^2$ test.

Results
A total of 80 patients were included in the study while 105 LNs were studied.

The patients’ demographics and characteristics are given in Table 1.

<table>
<thead>
<tr>
<th>Elastography type</th>
<th>Number of benign LNs/total number [n/N (%)]</th>
<th>Number of malignant LNs/total number [n/N (%)]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Type 1 (N=46)</td>
<td>42/46 (91.30)</td>
<td>4/46 (8.70)</td>
</tr>
<tr>
<td>Type 2 (N=30)</td>
<td>24/30 (80)</td>
<td>6/30 (20)</td>
</tr>
<tr>
<td>Type 3 (N=29)</td>
<td>13/29 (44.82)</td>
<td>16/29 (55.14)</td>
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</table>

LN, lymph node.

A qualitative classification based on the patterns and colors of LNs was performed. Elastographic patterns/colors were described according to the dominant colors and their distribution within the target LN.

The following representation formed the premise for the classification of elastographic types:

Type 1 group: largely nonblue (green, yellow, and red). Type 2 group: part blue, part nonblue (green, yellow, and red). Type 3 group: largely blue.

The elastography colors and patterns were compared with the eventual pathologic diagnosis (Table 2).

The nodes which were labeled as type 1 on EBUS elastography were benign in 42/46 (91.30%) and malignant in 4/46 (8.70%); 40/46 were of size less than 20 mm; for type 2 LNs, 24/30 (80%) were benign and 6/20 (20%) were malignant; 10/30 were of size less than 20 mm; type 3 nodes were found to be of benign pathology in 13/29 (44.82%) and malignant in 16 (55.14%), 4/29 were of size less than 20 mm.
On labeling type 1 and type 2 as ‘benign,’ and type 3 as ‘malignant,’ the sensitivity, specificity, positive predictive value, negative predictive value, and diagnostic accuracy rates were 83.54, 61.54, 86.84, 55.17, and 78.10%, respectively. Only LNs with adequate lymphocytes or those with a definitive diagnosis were included for data analysis (Figs 1–3).

![Figure 1](image1)

Type 1 can be seen on the elastography image.

![Figure 2](image2)

Type 2 can be seen on the elastography image.
Discussion

EBUS elastography complements the conventional EBUS without significant prolongation of procedure, cost, or added complications. It can be performed in real-time during the procedure and can give important information that can impact the final diagnosis.

Elastography can be considered an imaging equivalent of the ancient skill of palpation. It is based on the principle that softer parts of tissues can deform easier under compression than the harder parts, allowing an objective and quantified determination of tissue consistency, showing differences in hardness between a diseased and normal tissue [5,6].

A cancerous lesion is usually hard and dense than the surrounding tissues, with comparatively less deformation under the surrounding pressures of oppression. Elastography can reflect the elastic properties of tissues and hence can judge the relative hardness of the lesion, facilitating the characterization and diagnosis of mediastinal lesions [7].

Elastography can be divided into two major divisions: qualitative one is based on the tissue response to an internal or external generated force, called strain elastography; quantitative one, based on measuring shear waves generated by a ‘push-pulse’ of low frequency, called shear wave elastography [8,9].

Strain ratio is a way to semiquantify information from strain images and can be calculated by measuring tissue stiffness in a targeted area comparing with an area outside of a region representing the normal tissue. The strain ratio value can then be calculated as the quotient between the two regions [10]. It is a semiquantitative method and has been found to give favorable results that have been reported for mediastinal LN analysis in patients with esophageal cancer.

This technology was initially used in the field of breast ultrasound and for evaluation of the thyroid, prostate, and liver [11–14].

Trosini-Désert et al. [15] in a small sample size, for the first time, showed that EBUS elastography was feasible and that the bronchial tracheal cartilage did not affect the collection of elastic data and concluded that elastography improved the diagnostic ability of EBUS.

One of the first initial reports on the use of EBUS elastography is by Izumo and colleagues, who analyzed...
75 LNs and classified the findings according to the color distribution: the nodes were classified in the same manner as in our study. On an evaluation of the LNs, 42 were malignant, 33 were found to be benign. The nodes that were classified as type 1 were benign by 100%; for type 2 LNs, 46.9% were found to be benign and 57.1% were malignant; and for type 3 LNs, 5.4% were benign (2/37) and 94.6% were malignant [4].

We had deviated results which could be accounted to more benign population in the study which led to increased fibrotic components in LNs which can cause ambiguity for the operator.

EBUS elastographic analysis can assist in guiding the puncture site in a non-necrotic part of the suspicious LN and hence improving the diagnostic accuracy and quality for further diagnostic testing.

Mean stiff ratios can also be calculated and used in analysis as was done by Nakajima et al. [16]. The group evaluated 49 LNs (16 malignant) in 21 patients by EBUS elastography. Mean stiff area ratios were found to be significantly high for the malignant nodes (0.478) than for benign LNs (0.216; P=0.0002). The group was able to show a sensitivity and specificity for predicting malignant disease of 0.81 and 0.85 using a cutoff value of 0.311 for stiff area ratio. The group did not use any color distribution for an additional analysis.

In a group of 40 lung cancer patients, a Chinese group, He and colleagues evaluated in a group of 40 lung cancer patients, the accuracy of B-mode features, elastographic strain ratio and a qualitative elastographic score to predict metastatic LN involvement. The elastographic score proved to have more sensitivity and specificity in determining the malignant LNs than all B-mode EBUS criteria. Further, the combination of B-mode criteria and elastography improved the diagnostic accuracy of EBUS to differentiate between benign and metastatic mediastinal LNs [17].

However, EBUS elastography is not meant currently to replace EBUS-fine needle aspiration for mediastinal LN diagnosis but it may be useful as a supplemental method to reduce the number of punctures and improve the yield of fine needle aspiration [18].

The potential application for elastography in EBUS would be targeting higher risk areas within normal looking sized LNs. It can assist in guiding the puncture site in the non-necrotic part of a suspicious LN when the necrotic tissue is present, as seen in advanced cancers. The real challenge would be in the evaluation of benign lesions where the fibrotic component would be a hindrance to a good mapping quality.

It can become an important tool in selecting nodes for sampling, thus improving the overall yield.

Several limitations should be highlighted in the present study. First, the number of LNs studied was limited. Second, previous studies suggested that strain ratio of elastography may be an objective indicator for the classification of LNs; however, we did not take this feature as we wanted a fast and practical way of predicting the characteristic of the LN.

Conclusion

In conclusion, our study shows a significant practical clinical utility of the addition of elastography mode for classifying and guiding TBNA in suspicious mediastinal LNs with a high positive predictive value and diagnostic accuracy.

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Nil.

Conflicts of interest

There are no conflicts of interest.

References


