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Mediastinum and Systematic Nodal Dissection



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EBUS/EUS FOR RESTAGING VALIDATED BY TEMPLA

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Real-time endobronchial ultrasound-guided transbronchial needle aspiration (EBUS-TBNA) and endoscopic ultrasound-guided fine needle aspiration by use of bronchoscope (EUS-b-FNA) are recently introduced methods of biopsy of the mediastinal, hilar and interlobar lymph nodes. Although widely accepted in non-small cell lung cancer (NSCLC) staging, still is lacking data about the role of both endosonographic techniques in lung cancer restaging after neo-adjuvant therapy. The actual data show quite high specificity and accuracy of both methods, but low negative predictive value (NPV) and sensitivity. It has been suggested that negative results of EBUS-TBNA and EUS-b-FNA in restaging should be confirmed surgically before thoracotomy. There are currently still no accepted standards regarding mediastinal restaging, and many strategies, based on radiological, minimally invasive and surgical techniques like transcervical extended mediastinal lymphadenectomy (TEMLA) are advocated. If the restaging endosonographic biopsies confirm persistent N2 disease, non-surgical treatment is indicated in most of patients. The use of the EBUS-TBNA and EUS-b-FNA in NSCLC restaging is gaining increasing acceptance, due to its efficiency and low invasiveness, particularly as the initial procedure. As the risk of complications related to the EBUS-TBNA and EUS-b-FNA is very low, these procedures may be performed even in the outpatient settings. However sensitivity of both endosonographic methods are reported to be 75-78%, specificity and PPV — 100% and accuracy — 74-77% and NPV — 20-78%. The patients undergoing restaging with endosonography followed by TEMPLA the most accurate referral test for EBUS-TBNA and EUS-b-FNA. All mediastinal nodal stations (according to the Mountain-Dresler map), except for the pulmonary ligament nodes (station 9) are removed during TEMPLA procedure. In our series in patients who underwent TEMPLA preceded by negative EBUS-TBNA and EUS-b-FNA sensitivity for endosonography was 72%-64%, depending on prevalence of mediastinal lymph node metastases 44% and 19% respectively and NPV was high 83%. In our studies, the mean number of biopsied nodes using EBUS-TBNA, EUS-b-FNA and a combination of both techniques were 2.1, 2.4, and 3.7, respectively in comparison with 27.9 as a mean number of nodes removed with the surrounding mediastinal fatty tissue during restaging TEMPLA. Such a big difference in the number of examined nodes was the main reason for the difference in the diagnostic yields, especially in sensitivity and NPV among endosonographic methods and TEMPLA (64% vs 96%

and 82 vs 98%, respectively). Eventhough the results promote maximally invasive surgical techinques as TEMPLA over endosonography for NSCLC restaging, EBUS-TBNA and EUS-b-FNA or both seem to be the method of the first choice. In the era of induction immunotherapy combined with chemotherapy very little is known about utility of all invasive techniques for restaging purposes.

References

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