





PERSISTENT N2 IS NOT A CONTRAINDICATION TO SURGERY FOR LUNG CANCER

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Surgical multimodal management of stage IIIA-N2 non-small cell lung cancer (NSCLC) has undergone a tremendous change with the introduction of chemo-immunotherapy in the neoadjuvant and perioperative setting. Molecular pathology and biomarkers gain increasing importance for determining personalized therapeutic strategies as well as for prognostic stratification.

Already prior to the introduction of immunotherapy in the curative setting subgroups of patients with persistent N2 disease were identified that showed a clear benefit from surgery. Superior long-term outcomes have repeatedly been demonstrated in patients with single station N2 compared to disease in multiple stations. This is also reflected in the current TNM9 staging with a refined N2 stage grouping in N2a (metastasis(es) in a single ipsilateral mediastinal or in the subcarinal nodal station) and N2b (metastases in multiple ipsilateral mediastinal nodal stations with or without involvement of the subcarinal nodal station).

The correlation between imaging-based mediastinal restaging after immunotherapy and pathological response rates is moderate. Thus, the radiographic judgment on persistent nodal disease is unreliable, and no clear guidelines on invasive mediastinal restaging after neoadjuvant chemo-immunotherapy have been established. Data on the prognostic impact of persistent nodal disease in modern multimodal treatment regimens are not yet consistent and sufficiently granular to allow definite conclusions. Current efforts are directed at obtaining a consensus definition of surgical resectability in stage III disease. In view of the excellent pathological response rates after neoadjuvant chemo-immunotherapy and the encouraging results of perioperative systemic treatment approaches persistent N2 disease should not routinely exclude patients from curative intent surgical resection.

References

Significantly favourable outcome for patients with non-small-cell lung cancer stage IIIA/IIIB and single-station persistent N2 (skip or additionally N1) disease after multimodality treatment. Stamatis G, Müller S, Weinreich G, Schwarz B, Eberhardt W, Pöttgen C, Aigner C. Eur J Cardiothorac Surg. 2022 Jan 24;61(2):269-276

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