Active substances set

Search phrase: osimertinib

Below you will find a list of active substances registered by the European Medical Agency (EMA) in the last 15 years, recommended by the European Society of Clinical Oncology (ESMO) and their reimbursement status in the country.

Tracheal, bronchus, and lung cancer

Osimertinib	Osimertinib as monotherapy is indicated for: - the adjuvant treatment after complete tumour resection in adult patients with stage IB-IIIA non small cell lung cancer (NSCLC) whose tumours have epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 (L858R) substitution mutations. - the treatment of adult patients with locally advanced, unresectable NSCLC whose tumours have EGFR exon 19 deletions or exon 21 (L858R) substitution mutations and whose disease has not progressed during or following platinum-based chemoradiation therapy. - the first-line treatment of adult patients with locally advanced or metastatic NSCLC with activating EGFR mutations. - the treatment of adult patients with locally advanced or metastatic EGFR T790M mutation-positive NSCLC. Osimertinib is indicated in combination with: - pemetrexed and platinum-based chemotherapy for the first-line treatment of adult patients with advanced NSCLC whose tumours have EGFR exon 19 deletions or exon 21 (L858R) substitution mutations.	REIMBURSEMENT WITH RESTRICTIONS ESMO