NATIONAL INSTITUTE FOR HEALTH AND CARE EXCELLENCE

Health Technology Evaluation

Alectinib for adjuvant treatment of ALK-positive non-small-cell lung cancer [ID6368]

Final scope

Remit/evaluation objective

To appraise the clinical and cost effectiveness of alectinib within its marketing authorisation as adjuvant treatment of ALK-positive non-small-cell lung cancer (NSCLC).

Background

Lung cancer is the third most common cancer and the most common cause of cancer death in the UK, accounting for 13% of all new cancer cases and 21% of all cancer deaths between 2017 and 2019. Most lung cancers are diagnosed at an advanced stage when the cancer has spread to lymph nodes and other organs in the chest (locally advanced disease; stage 3) or to other parts of the body (metastatic disease; stage 4). Around 30% of lung cancers are diagnosed at an early stage (stage 1 or 2) and around 10% diagnosed at stage 3a.²

In 2021, 91% (around 31,000) of people diagnosed with lung cancer in England had NSCLC.² Of these people, 17% (5,333) had surgical treatment for their cancer.² Despite the curative intent of treatment for early-stage lung cancer, survival is poor, with only about 57% people with stage 1, 34% with stage 2 and 13% with stage 3 surviving for 5 years after diagnosis.³ It is estimated that around 2 to 6% of NSCLCs have an ALK fusion genetic alteration. This alteration inhibits processes which stop lung cells dividing and can lead to cancer.⁴,5,6 ALK fusions can occur in any type of NSCLC, but are most likely to occur in adenocarcinoma histology.⁵

The treatment pathway for NSCLC can be divided into interconnected decision points based on the number staging system and line of therapy. Treatment choices are influenced by the presence of biological markers (including programmed cell death 1 ligand PD-L1 status), oncogenic driver genetic alterations, histology (squamous or non-squamous) and previous treatment. NICE's Technology Appraisal Pathway Pilot scope for treatments for non-small-cell lung cancer outlines in more detail the NSCLC treatment pathway.

NICE guideline 122 (NG122) '<u>Lung cancer: diagnosis and management</u>' recommends surgery, radiotherapy, chemoradiotherapy or a combination of these for stage 1 to 2 NSCLC. People may be offered a neo-adjuvant (before surgical removal of cancerous tumour) treatment which could be platinum based chemotherapy, or nivolumab with chemotherapy as recommended by NICE <u>TA876</u>. Neoadjuvant chemotherapy has shown equivalent outcomes in terms of survival to adjuvant chemotherapy.⁶

For stage 3 NSCLC, surgery is carried out if the surgeon deems the tumour to be resectable. Before surgery, chemoradiotherapy (chemotherapy with radiotherapy) may be used or surgery may potentially be followed by chemotherapy. If well enough,

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people may be offered a cisplatin-based chemotherapy (adjuvant treatment) after surgery.

People who have had surgery may have an adjuvant treatment. NICE <u>TA761</u> recommends osimertinib in the Cancer Drugs Fund as adjuvant treatment for people whose cancer has an EGFR exon 19 deletion or an exon 21 (L858R) substitution mutation. For people whose cancer does not have an EGFR mutation, platinum chemotherapy may be offered as adjuvant treatment. NICE <u>TA823</u> recommends atezolizumab in the Cancer Drugs Fund as an option for maintenance treatment after complete tumour resection in adults with stage 2 to 3a NSCLC and adjuvant chemotherapy, but this is not available in routine commissioning.

People with confirmed ALK-fusion positive NSCLC also have several treatment options available for untreated advanced and metastatic disease. These include crizotinib (<u>TA406</u>), ceritinib (<u>TA500</u>), alectinib (<u>TA536</u>) and brigatinib (<u>TA670</u>).

The technology

Alectinib (Alecensa, Roche) does not currently have a marketing authorisation in the UK for adjuvant treatment of ALK-fusion positive NSCLC. It is being studied in comparison with platinum based chemotherapy in adults with resected ALK-positive NSCLC.

Alectinib monotherapy is currently licenced for other indications including as first-line treatment of ALK positive advanced NSCLC.

Intervention(s)	Alectinib
Population(s)	Adults with ALK-positive NSCLC who have undergone surgical resection
Subgroups	If the evidence allows it, the following subgroups will be considered:
	Disease stage
	• Ethnicity
Comparators	Established adjuvant clinical management without alectinib, which may include:
	 Platinum-based chemotherapy
	Active monitoring
	 Pembrolizumab (subject to NICE appraisal)

Outcomes The outcome measures to be considered include: disease-free survival overall survival adverse effects of treatment health-related quality of life. **Economic analysis** The reference case stipulates that the cost effectiveness of treatments should be expressed in terms of incremental cost per quality-adjusted life year. The reference case stipulates that the time horizon for estimating clinical and cost effectiveness should be sufficiently long to reflect any differences in costs or outcomes between the technologies being compared. Costs will be considered from an NHS and Personal Social Services perspective. The availability of any commercial arrangements for the intervention, comparator and subsequent treatment technologies will be taken into account. The availability and cost of biosimilar and generic products should be taken into account. The use of alectinib is conditional on the presence of an ALK gene fusion. The economic modelling should include the costs associated with the diagnostic testing for ALK gene fusions in people with resected NSCLC who would not otherwise have been tested. A sensitivity analysis should be provided without the cost of the diagnostic test. See section 4.8 of the guidance development manual (available here: https://www.nice.org.uk/process/pmg36/chapter/introductionto-health-technology-evaluation). Other Guidance will only be issued in accordance with the considerations marketing authorisation. Where the wording of the therapeutic indication does not include specific treatment combinations, guidance will be issued only in the context of the evidence that has underpinned the marketing authorisation granted by the regulator. **Related NICE** Related technology appraisals: recommendations Nivolumab with chemotherapy for neoadjuvant treatment of resectable non-small-cell lung cancer (2023) NICE technology appraisal guidance 876 Atezolizumab for adjuvant treatment of resected non-smallcell lung cancer (2022) NICE technology appraisal guidance 823 [In CDF]

Osimertinib for adjuvant treatment of EGFR mutation-positive non-small-cell lung cancer after complete tumour resection (2022) TA761 Related technology appraisals in development: Pembrolizumab for adjuvant treatment of resected non-smallcell lung cancer [ID3907] Publication date to be confirmed Pembrolizumab for neoadjuvant and adjuvant treatment of resectable stage 2 to 3B non-small-cell lung cancer [ID5094] Publication date to be confirmed Atezolizumab with chemotherapy for neoadjuvant and adjuvant treatment of resectable non-small-cell lung cancer [ID3894] Publication date to be confirmed Nivolumab for adjuvant treatment of resected non-small-cell lung cancer [ID4053] Publication date to be confirmed Related NICE guidelines: Lung cancer: diagnosis and management (NG122) Related quality standards: Lung cancer in adults (2019) NICE quality standard 17 **Related National** The NHS Long Term Plan, 2019. NHS Long Term Plan **Policy** NHS England (2018/2019) NHS manual for prescribed specialist services (2018/2019) Chapter 105: Specialist cancer services (adults).

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- 6. Schneider JL, Lin JJ, Shaw AT. ALK-positive lung cancer: a moving target. Nat Cancer. Accessed February 2024