

Technology Guidance

Alectinib

for adjuvant treatment of resectable, ALK mutation-positive non-small-cell lung cancer

Technology Guidance from the MOH Drug Advisory Committee

Guidance Recommendations

The Ministry of Health's Drug Advisory Committee has not recommended alectinib for inclusion on the MOH List of Subsidised Drugs for adjuvant treatment of resectable, anaplastic lymphoma kinase mutation-positive non-small-cell lung cancer. The decision was due to the unacceptable pricing and price-volume agreement proposed by the company.

Clinical indication, subsidy class and MediShield Life claim limit for alectinib are provided in the Annex.

Updated: 16 September 2025



ANNEX

Recommendations by the MOH Drug Advisory Committee

Drug preparation	Approved clinical indication	Subsidy class	MediShield Life claim limit per month (implementation date)
Alectinib 150 mg	Adjuvant treatment as	Not recommended	\$2,000
capsule	monotherapy after tumour resection in patients with stage II to IIIA or stage IIIB (T3N2), ALK mutation-positive NSCLC, according to the 8th edition of the AJCC/UICC cancer staging system. Treatment should be continued until disease recurrence, unacceptable toxicity, or for a maximum of 2 years.	for subsidy	(1 Nov 2025)

Abbreviations: ALK, anaplastic lymphoma kinase; AJCC/UICC: American Joint Committee on Cancer/International Union Against Cancer; NSCLC, non-small-cell lung cancer



VERSION HISTORY

Guidance on alectinib for adjuvant treatment of resectable, ALK mutation-positive nonsmall-cell lung cancer

This Version History is provided to track any updates or changes to the guidance following the first publication date. It is not part of the guidance.

1. Publication of guidance

Date of Publication 2 Jun 2025

2. Guidance updated to include alectinib on the Cancer Drug List

Date of Publication 16 Sep 2025

Agency for Care Effectiveness - ACE in Agency for Care Effectiveness (ACE)

About the Agency

The Agency for Care Effectiveness (ACE) was established by the Ministry of Health (Singapore) to drive better decision-making in healthcare through health technology assessment (HTA), clinical guidance, and education.

As the national HTA agency, ACE conducts evaluations to inform government funding decisions for treatments, diagnostic tests and vaccines, and produces guidance for public hospitals and institutions in Singapore.

The guidance is not, and should not be regarded as, a substitute for professional or medical advice. Please seek the advice of a qualified healthcare professional about any medical condition. The responsibility for making decisions appropriate to the circumstances of the individual patient remains with the healthcare professional.

Find out more about ACE at www.ace-hta.gov.sg/about

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