# **U**NOVARTIS

### TAF MEK Lung cancer - BRAF and mechanism of action - HCP

### Prescribing information

Image



Image



## **BRAF** mutation and mechanism of action for TAFINLAR® (dabrafenib) + MEKINIST® (trametinib)

TAFINLAR in combination with MEKINIST is indicated in adult patients with advanced nonsmall cell lung cancer (NSCLC) with a *BRAF* V600 mutation.<sup>1,2</sup>

For the full safety profile, please refer to the Summary of Product Characteristics (SmPC) for <u>TAFINLAR</u> and <u>MEKINIST</u>.

Adverse event reporting: Details of how to report adverse events are available at

the bottom of the page. Please refer to the respective SmPC for all licensed indications.

# **BRAF** represents a novel therapeutic target for the treatment of advanced NSCLC<sup>3</sup>

### Testing for BRAF in the UK

In England, *BRAF* testing can now be done as part of the next-generation sequencing (NGS) panel. For more information, go to the national genomic test directory for cancer in England.<sup>4</sup>

Visit the national genomic test directory for cancer in England



Molecular characterisation of NSCLC marked a turning point in the treatment of lung tumours harbouring kinase alterations suitable for targeted drug inhibition.<sup>3</sup>

## The role of BRAF mutations in NSCLC

Approximately 1.5-3.5% of patients with NSCLC have a BRAF mutation.<sup>3</sup>

*BRAF* mutations lead to constitutive activation of the mitogen-activated protein kinase (MAPK) pathway.<sup>5</sup>

*BRAF* V600E is the most common variant, occurring in roughly 50% of *BRAF*-mutated tumours.<sup>6</sup>

NSCLC with *BRAF* V600E mutations has histological features suggestive of an aggressive tumour.<sup>3</sup>

In patients with NSCLC who received platinum-based chemotherapy, outcomes were less favourable among those with a *BRAF* V600E-mutated tumour vs those with *BRAF* wild-type (WT) tumours.<sup>5</sup>

#### Image



Adapted from Hirsch FR, et al. 2017.<sup>7</sup>

# **TAFINLAR + MEKINIST** target two distinct points on the MAPK pathway to provide concomitant inhibition<sup>1,2</sup>

## Mechanism of action for TAFINLAR + MEKINIST

TAFINLAR + MEKINIST target two different kinases in the MAPK pathway (*BRAF* and MEK1/2, respectively) to block the signalling that leads to abnormal cell division and growth.<sup>1,2</sup>

Combination therapy with TAFINLAR + MEKINIST results in prolonged inhibition of tumour growth compared with either drug alone in *BRAF* V600E-mutant tumours, both *in vitro* and *in vivo*.<sup>1,2</sup>

### Schematic of the MAPK signalling pathway showing where TAFINLAR and



ALK, anaplastic lymphoma kinase; *BRAF*, v-raf murine sarcoma viral oncogene homolog B1; *BRAF* V600, mutation of the *BRAF* gene at valine (V) 600; *BRAF* V600E, mutation of the *BRAF* gene at valine (V) 600 to glutamate (E); EGFR, epidermal growth factor receptor; ERK, extracellular signal-regulated kinases; HER2, human epidermal growth factor receptor-2; KRAS, Kirsten rat sarcoma viral oncogene homolog; MAPK, mitogen-activated protein kinase; MEK, mekinist; MEK1/2, mitogen-activated protein kinase 1/2; MET, mesenchymal epithelial transition factor receptor; NGS, next-generation sequencing; NSCLC, non-small cell lung cancer; NTRK1, neurotrophic receptor tyrosine kinase 1; PIK3CA, phosphatidylinositol-4,5-bisphosphate 3-kinase catalytic subunit alpha; RAS, rat sarcoma; RET, ret proto-oncogene; ROS1, proto-oncogene 1, receptor tyrosine kinase; SmPC, summary of product characteristics; WT, wild-type.

#### References

- 1. TAFINLAR (dabrafenib) Summary of Product Characteristics.
- 2. MEKINIST (trametinib) Summary of Product Characteristics.
- 3. Leonetti A, et al. Cancer Treat Rev 2018;66:82-94.
- 4. NHS England. National Genomic Test Directory for cancer. Available at: <u>https://www.england.nhs.uk/publication/national-genomic-test-directories/</u> [Accessed April 2025].

- 5. Planchard D, et al. *Lancet Oncol* 2016;17:984-993.
- 6. O'Leary C, et al. Transl Lung Cancer Res 2019;8:1119-1124.
- 7. Hirsch FR, et al. Lancet 2017;389:299-311.

×

Efficacy

Efficacy

See more details

Hide details

×

Safety profile

Safety profile

See more details

Hide details

×

Dosing and administration

Dosing and administration

See more details

Hide details

UK | April 2025 | FA-11392355

Adverse events should be reported. Reporting forms and information can be found at <u>www.mhra.gov.uk/yellowcard</u>. Adverse events should also be reported to Novartis online through the pharmacovigilance intake (PVI) tool at <u>www.novartis.com/report</u>, or alternatively email <u>medinfo.uk@novartis.com</u> or call 01276 698370.

#### Source URL:

https://www.pro.novartis.com/uk-en/medicines/oncology/tafinlar-mekinist/lung-cancer/brafmoa