PRELIMINARY SCORE

CURATIVE

NON-CURATIVE

ORR

ADJUSTMENTS

Quality of life

Not qualified for an ESMO-MCBS credit

Serious and disabling adverse effects

Other adjustments

Repotrectinib TRIDENT-1

SCORE

CURATIVE

Overall Survival / Disease-Free Survival / Pathological Complete Response

NON-CURATIVE

Overall Survival

Progression-Free Survival

Non-inferiority (Improved Quality of Life or Reduced Adverse Events) / Response Rate

3

Overall Response Rate / Duration of Response

Overall Survival / Disease-Free Survival / Pathological Complete Response

INFORMATION

Tumour type: Thoracic Malignancies
Therapeutic Indication: FDA: Repotrectinib used for locally advanced or metastatic ROS1-positive non-small cell lung cancer. FDA approval that includes patients with ROS1-positive NSCLC who have previously received a ROS1 tyrosine kinase inhibitor (TKI). EMA: Repotrectinib as monotherapy is indicated for the treatment of adult patients with ROS1-positive advanced non-small cell lung cancer (NSCLC) who have previously received a ROS1 tyrosine kinase inhibitor (TKI).
Experimental Arm: Repotrectinib
Control Arm: Single arm