



## PRELIMINARY SCORE

## CURATIVE



## NON-CURATIVE



## ADJUSTMENTS

## Quality of life



QoL data pending



## Serious and disabling adverse effects



## Other adjustments



## 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



Overall Response Rate / Duration of Response

Overall Survival / Disease-Free Survival / Pathological Complete Response

## INFORMATION

Tumour type: Thoracic Malignancies

Therapeutic Indication: FDA: Amivantamab in combination with lazertinib for the first-line treatment of locally advanced or metastatic non-small cell lung cancer (NSCLC) with epidermal growth factor receptor (EGFR) exon 19 deletions or exon 21 L858R substitution mutations. EMA: Amivantamab in combination with lazertinib is indicated for the first-line treatment of adult patients with advanced non-small cell lung cancer (NSCLC) with EGFR exon 19 deletions or exon 21 L858R substitution mutations.

Experimental Arm: Amivantamab + lazertinib

Control Arm: Osimertinib

