Nivolumab





Nivolumab CheckMate 649 Nivolumab CheckMate 649 PRELIMINARY SCORE **FINAL SCORE** CURATIVE **CURATIVE** Overall Survival / Disease-Free Survival / Pathological Complete Response **NON-CURATIVE NON-CURATIVE** Overall Survival **ADJUSTMENTS** Quality of life Progression-Free Survival Non-inferiority (Improved Quality of Life or Reduced Adverse Events) / Response Rate Serious and disabling adverse effects Overall Response Rate / Duration of Response Overall Survival / Disease-Free Survival / Pathological Complete Response Other adjustments INFORMATION Tumour type: Gastrointestinal Cancers Therapeutic Indication: First-line treatment for patients with HER2-negative advanced or metastatic gastric, gastroesophageal junction or oesophageal adenocarcinoma whose tumours express PD-L1 with a CPS ≥5 Experimental Arm: Nivolumab + Fluoropyrimidine and platinum ChT (FOLFOX or CAPOX) Control Arm: ChT (FOLFOX or CAPOX)



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