Pembrolizumab

KEYNOTE-054



Pembrolizumab KEYNOTE-054 Pembrolizumab KEYNOTE-054 PRFLIMINARY SCORE **SCORE** CURATIVE **CURATIVE** DFS Overall Survival / Disease-Free Survival / Pathological Complete Response **NON-CURATIVE NON-CURATIVE** Overall Survival **ADJUSTMENTS** Quality of life Progression-Free Survival Not qualified for an ESMO-MCBS credit 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: Skin Cancers Therapeutic Indication: EMA: Pembrolizumab for adjuvant treatment of adults and adolescents aged 12 years and older with stage IIB, IIC or III melanoma and who have undergone complete resection.. FDA: Pembrolizumab for the adjuvant treatment of patients with melanoma with involvement of lymph node(s) following complete resection. Experimental Arm: Pembrolizumab Control Arm: Placebo



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