

Durvalumab

DUO-E

PRELIMINARY SCORE

CURATIVE



NON-CURATIVE



ADJUSTMENTS

Quality of life



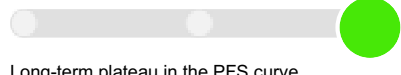
QoL data pending



Serious and disabling adverse effects



Other adjustments



Long-term plateau in the PFS curve



FINAL SCORE

CURATIVE



Overall Survival / Disease-Free Survival / Pathological Complete Response

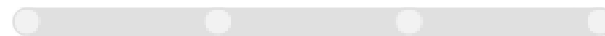
NON-CURATIVE



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: Gynaecological Malignancies

Therapeutic Indication: FDA: Durvalumab is indicated with carboplatin plus paclitaxel followed by single-agent durvalumab for adult patients with primary advanced or recurrent endometrial cancer that is mismatch repair deficient (dMMR). EMA: Durvalumab in combination with carboplatin and paclitaxel is indicated for the first-line treatment of adults with primary advanced or recurrent endometrial cancer who are candidates for systemic therapy followed by maintenance treatment with durvalumab as monotherapy in endometrial cancer that is mismatch repair deficient (dMMR)

Experimental Arm: Durvalumab + Carboplatin and paclitaxel

Control Arm: Carboplatin, paclitaxel, and placebo (arm A)



