Enfortumab

EV-302/KN-A39



Enfortumab EV-302/KN-A39 Enfortumab EV-302/KN-A39 PRELIMINARY SCORE **SCORE CURATIVE CURATIVE** Overall Survival / Disease-Free Survival / Pathological Complete Response **NON-CURATIVE NON-CURATIVE** Overall Survival **ADJUSTMENTS** Quality of life Progression-Free Survival QoL data pending 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: Genitourinary Cancers Therapeutic Indication: FDA: Enfortumab vedotin in combination with pembrolizumab for patients with locally advanced or metastatic urothelial cancer. EMA: Pembrolizumab, in combination with enfortumab vedotin, is indicated for the first-line treatment of unresectable or metastatic urothelial carcinoma in adults. Experimental Arm: Enfortumab + Pembrolizumab Control Arm: Platinum-based ChT



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