Active substances set

Search phrase: ramucirumab

Below you will find a list of active substances registered by the European Medical Agency (EMA) in the last 15 years, recommended by the European Society of Clinical Oncology (ESMO) and their reimbursement status in the country.

Malignant stomach cancer

Ramucirumab	Ramucirumab in combination with paclitaxel is indicated for the treatment of adult patients with advanced gastric cancer or gastro-oesophageal junction adenocarcinoma with disease progression after prior platinum and fluoropyrimidine chemotherapy. Ramucirumab monotherapy is indicated for the treatment of adult patients with advanced gastric cancer or gastro-oesophageal junction adenocarcinoma with disease progression after prior platinum or fluoropyrimidine chemotherapy, for whom treatment in combination with paclitaxel is not appropriate.	REIMBURSEMENT WITH RESTRICTIONS ESMO

Colon and rectum cancer

Ramucirumab	Ramucirumab, in combination with FOLFIRI (irinotecan, folinic acid, and 5-fluorouracil), is indicated for the treatment of adult patients with metastatic colorectal cancer (mCRC) with disease progression on or after prior therapy with bevacizumab, oxaliplatin and a fluoropyrimidine.	NO REIMBURSEMENT ESMO

Liver cancer

Ramucirumab	Ramucirumab monotherapy is indicated for the treatment of adult patients with advanced or unresectable hepatocellular carcinoma who have a serum alpha fetoprotein (AFP) of ≥ 400 ng/ml and who have been previously treated with sorafenib.	REIMBURSEMENT WITH RESTRICTIONS ESMO

Tracheal, bronchus, and lung cancer

Ramucirumab	Ramucirumab in combination with erlotinib is indicated for the first-line treatment of adult patients with metastatic non-small cell lung cancer with activating epidermal growth factor receptor (EGFR) mutations. Ramucirumab in combination with docetaxel is indicated for the treatment of adult patients with locallyadvanced or metastatic non-small cell lung cancer with disease progression after platinum-based chemotherapy.	NO REIMBURSEMENT ESMO