Active substances set

Search phrase: capivasertib

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 breast cancer

Capivasertib	Capivasertib is indicated in combination with fulvestrant for the treatment of adult patients with oestrogen receptor (ER)-positive, HER2-negative locally advanced or metastatic breast cancer with one or more PIK3CA/AKT1/PTEN-alterations following recurrence or progression on or after an endocrine-based regimen. In pre- or perimenopausal women, Capivasertib plus fulvestrant should be combined with a luteinising hormone releasing hormone (LHRH) agonist. For men, administration of LHRH agonist according to current clinical practice standards should be considered.	REIMBURSEMENT WITH RESTRICTIONS ESMO