Active substances set

Search phrase: ibrutinib

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.

non-Hodgkin lymphoma

Ibrutinib	Ibrutinib as a single agent is indicated for the treatment of adult patients with relapsed or refractory mantle cell lymphoma (MCL).	REIMBURSEMENT WITH RESTRICTIONS ESMO

Multiple myeloma and plasma cell neoplasms

Ibrutinib	Ibrutinib as a single agent is indicated for the treatment of adult patients with Waldenström's macroglobulinaemia (WM) who have received at least one prior therapy, or in first line treatment for patients unsuitable for chemo-immunotherapy. Ibrutinib in combination with rituximab is indicated for the treatment of adult patients with WM.	REIMBURSEMENT WITH RESTRICTIONS ESMO

Chronic lymphoid leukemia

Ibrutinib	Ibrutinib as a single agent or in combination with rituximab or obinutuzumab or venetoclax is indicated for the treatment of adult patients with previously untreated chronic lymphocytic leukaemia (CLL). Ibrutinib as a single agent or in combination with bendamustine and rituximab (BR) is indicated for the treatment of adult patients with CLL who have received at least one prior therapy.	REIMBURSEMENT WITH RESTRICTIONS ESMO