

Active substances set

Search phrase: ponatinib

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.

Acute lymphoid leukemia

Ponatinib

Ponatinib is indicated in adult patients with Philadelphia chromosome positive acute lymphoblastic leukaemia (Ph+ ALL) who are resistant to dasatinib; who are intolerant to dasatinib and for whom subsequent treatment with imatinib is not clinically appropriate; or who have the T315I mutation.

 **REIMBURSEMENT
WITH RESTRICTIONS**

 **ESMO**

Chronic myeloid leukemia

Ponatinib

Ponatinib is indicated in adult patients with chronic phase, accelerated phase, or blast phase chronic myeloid leukaemia (CML) who are resistant to dasatinib or nilotinib; who are intolerant to dasatinib or nilotinib and for whom subsequent treatment with imatinib is not clinically appropriate; or who have the T315I mutation.

 **FULL
REIMBURSEMENT**

 **ESMO**