

**Prescriber Criteria Form**

Iclusig 2026 PA Fax 920-A v1 010126.docx  
Iclusig (ponatinib)  
Coverage Determination

This fax machine is located in a secure location as required by HIPAA regulations.  
Complete/review information, sign and date. Fax signed forms to CVS Caremark at **1-855-633-7673**. Please contact CVS Caremark at **1-866-785-5714** with questions regarding the prior authorization process. When conditions are met, we will authorize the coverage of Iclusig (ponatinib).

Drug Name:  
Iclusig (ponatinib)

**Patient Name:**

**Patient ID:**

**Patient DOB:**

**Patient Phone:**

**Prescriber Name:**

**Prescriber Address:**

**City:**

**State:**

**Zip:**

**Prescriber Phone:**

**Prescriber Fax:**

**Diagnosis:**

**ICD Code(s):**

**Please circle the appropriate answer for each question.**

1	Does the patient have a diagnosis of acute lymphoblastic leukemia (ALL), including patients who have received a hematopoietic stem cell transplant? [If no, then skip to question 3.]	Yes	No
2	Was the diagnosis confirmed by detection of the Philadelphia chromosome or BCR-ABL gene? [No further questions.]	Yes	No
3	Does the patient have a diagnosis of chronic myeloid leukemia (CML), including patients who have received a hematopoietic stem cell transplant? [If no, then skip to question 8.]	Yes	No
4	Does the patient have accelerated or blast phase chronic myeloid leukemia (CML) and no other kinase inhibitor is indicated? [If yes, then no further questions.]	Yes	No
5	Does the patient have chronic phase chronic myeloid leukemia (CML) and has experienced resistance or intolerance to at least 2 prior kinase inhibitors AND at least ONE of those was imatinib, dasatinib, or nilotinib? [If yes, then no further questions.]	Yes	No
6	Is the patient positive for the T315I mutation? [If yes, then no further questions.]	Yes	No

7	Does the patient meet both of the following criteria: A) chronic myeloid leukemia (CML) with no identifiable BCR::ABL1 mutations, B) resistance to primary treatment with imatinib, bosutinib, dasatinib, or nilotinib? [No further questions.]	Yes	No
8	Does the patient have a diagnosis of myeloid and/or lymphoid neoplasms with eosinophilia and fibroblast growth factor receptor 1 (FGFR1) or ABL1 rearrangement? [If no, then skip to question 10.]	Yes	No
9	Is the disease in chronic phase or blast phase? [No further questions.]	Yes	No
10	Does the patient have a diagnosis of gastrointestinal stromal tumors? [If no, then no further questions.]	Yes	No
11	Does the disease meet any of the following: A) residual, B) unresectable, C) recurrent, D) metastatic/tumor rupture? [If no, then no further questions.]	Yes	No
12	Has the disease progressed after use of at least two Food and Drug Administration (FDA)-approved therapies (e.g., imatinib, sunitinib, regorafenib, ripretinib)?	Yes	No

Comments:	

By signing this form, I attest that the information provided is accurate and true as of this date and that the documentation supporting this information is available for review if requested by the health plan.	
<b>Prescriber (or Authorized) Signature:</b> _____ <b>Date:</b> _____	