

# Chronic Myelogenous Leukemia (CML) Pathways

Patient Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Member Number: \_\_\_\_\_ Treatment Start Date: \_\_\_\_\_

ICD-10 Code: \_\_\_\_\_ Pathology: \_\_\_\_\_

Stage:  New diagnosis or  Relapse

Line of Treatment:  First Line  Second Line  Third Line  Third Line +

ECOG Performance Status:  0  1  2  3  4

## Biomarkers:

CML Phase:  Chronic Phase  Accelerated Phase  Lymphoid Blast Phase  Myeloid Blast Phase  Not Reported

Imatinib resistant or intolerant:  Yes  No

Philadelphia chromosome:  Positive  Negative

T315I:  Positive  Negative

Mutation:  V299L  T315I

## First line of therapy (1<sup>st</sup> line)

Dasatinib\* (Sprycel) for intermediate or high risk disease

Imatinib (Gleevec)

Nilotinib\* (Tasigna) for intermediate or high risk disease

## Second line of therapy (2<sup>nd</sup> line) | Following treatment failure, suboptimal response<sup>†</sup>, or intolerance to first line therapy

Bosutinib (Bosulif)

Dasatinib (Sprycel)

Nilotinib (Tasigna)

Ponatinib<sup>‡</sup> (Iclusig)

## Third line of therapy (3<sup>rd</sup> line)

Ponatinib (Iclusig)

\* For patients with intermediate or high risk disease based on Sokal or Hasford Score:

- Sokal: Intermediate Risk=0.8-1.2; High Risk>1.2
- Hasford: Intermediate Risk=781-1480; High Risk>1480

† Defined as lack of complete hematologic response or BCR-ABL1 transcripts > 10% (IS) or lack of partial cytogenetic response on bone marrow cytogenetics.

‡ Pathway option for second line therapy only after failure, suboptimal response, or intolerance of a second generation TKI has been used in the first line setting, or T315I mutation has been identified.

**Note:** Pathway lists are solely for the purpose of eligibility for enhanced reimbursement and are independent of specific health plan medical policy coverage criteria. Health plan medical policy/clinical guidelines should be consulted to determine whether proposed services will be covered.

