Louisiana Medicaid
Oncology Agents – Oral – Hematologic

The *Louisiana Uniform Prescription Drug Prior Authorization Form* should be utilized to request prior authorization for non-preferred oral hematologic oncology agents.

Additional Point-of-Sale edits may apply.

By submitting the authorization request, the prescriber attests to the conditions available HERE. ~~*These agents may have **Black Box Warnings** and/or may be subject to **Risk Evaluation and Mitigation Strategy (REMS)** under FDA safety regulations. Please refer to individual prescribing information for details.*~~

Approval Criteria for Initiation and Continuation of Therapy

~~**Approval Criteria for Initial and Reauthorization Requests**~~

- ~~For lenalidomide capsule (generic for Revlimid®) – there has been a treatment failure or intolerable side effect with or contraindication to brand Revlimid®; **OR**~~
- There is no preferred alternative that is the exact same chemical entity, formulation, strength, etc.; **AND**
- Previous use of a preferred product - **ONE** of the following is required:
  - The recipient has had a *treatment failure* with at least one preferred product; **OR**
  - The recipient has had an *intolerable side effect* to at least one preferred product; **OR**
  - The recipient has *documented contraindication(s)* to all of the preferred products that are appropriate to use for the condition being treated; **OR**
  - There is *no preferred product that is appropriate* to use for the condition being treated; **OR**
  - The prescriber states that the recipient is currently using the requested medication; **AND**
- The requested medication has been prescribed for an approved diagnosis (if applicable).~~; **AND**~~

- ~~By submitting the authorization request, the prescriber attests to the following:~~
  - ~~The prescribing information for the requested medication has been thoroughly reviewed, including any Black Box Warning, Risk Evaluation and Mitigation Strategy (REMS), contraindications, minimum age requirements, recommended dosing, and prior treatment requirements; **AND**~~
  - ~~All laboratory testing and clinical monitoring recommended in the prescribing information have been completed as of the date of the request and will be repeated as recommended; **AND**~~
  - ~~The recipient has no concomitant drug therapies or disease states that limit the use of the requested medication and will not be receiving the requested medication in combination with any other medication that is contraindicated or not recommended per FDA labeling.~~

Duration of approval for initiation and continuation of therapy~~Duration of initial and reauthorization approval~~:
Up to 12 months based upon patient-specific factors and the condition being treated.

References

Clinical Pharmacology [database online]. Tampa, FL: Gold Standard, Inc.; https://www.clinicalkey.com/pharmacology/

DiPiro JT, Talbert RL, Yee GC, Matzke GR, Wells BG, Posey L. eds. Pharmacotherapy: A Pathophysiologic Approach, 10e New York, NY: McGraw-Hill; https://accesspharmacy.mhmedical.com/book.aspx?bookid=1861

Revision / Date	Implementation Date
Single PDL Implementation	May 2019
Reviewed current criteria and no changes made / September 2019	January 2020
Added specific wording for use of Gleevec®, separated "Oncology Agents" into individual therapeutic class documents / November 2019	January 2020
Removed wording requiring use of brand Gleevec® and removed reference, removed references for Pomalyst® and Revlimid®, formatting changes / July 2020	July 2020
Formatting changes / September 2021	January 2022
Added wording for Revlimid® / November 2022	January 2023
Removed specific wording for the use of Revlimid®, formatting changes / April 2024	July 2024