

**Priority Health Medicare prior authorization form**

Fax completed form to: 877.974.4411 toll free, or 616.942.8206

This form applies to:  Medicare Part B  Medicare Part D  
 This request is:  Expedited request  Standard request

Your request will be expedited if you haven't gotten the prescription and Priority Health Medicare determines, or your prescriber tells us, that your life or health may be at risk by waiting.

**Rydapt<sup>TM</sup>** (midostaurin)

**Member**

Last Name: \_\_\_\_\_ First Name: \_\_\_\_\_  
 ID #: \_\_\_\_\_ DOB: \_\_\_\_\_ Gender: \_\_\_\_\_  
 Primary Care Physician: \_\_\_\_\_  
 Requesting Provider: \_\_\_\_\_ Prov. Phone: \_\_\_\_\_ Prov. Fax: \_\_\_\_\_  
 Provider Address: \_\_\_\_\_  
 Provider NPI: \_\_\_\_\_ Contact Name: \_\_\_\_\_  
 Provider Signature: \_\_\_\_\_ Date: \_\_\_\_\_

**Drug information**

New request  Continuation request  
 Drug Product:  Rydapt 25 mg capsule  
 Start date (or date of next dose): \_\_\_\_\_  
 Date of last dose (if applicable): \_\_\_\_\_  
 Dosing frequency: \_\_\_\_\_

**Prior authorization criteria**

The following requirements need to be met before this drug is covered by Priority Health Medicare. These requirements have been approved by the Centers for Medicare and Medicaid Services (CMS), but you may ask us for an exception if you believe one or more of these requirements should be waived.

1. Must be 18 years or older
2. The patient must have one of the following diagnoses and meet any required criteria:
  - Advanced systemic mastocytosis including aggressive systemic mastocytosis (ASM), systemic mastocytosis with associated hematological neoplasm (SM-AHN), or mast cell leukemia (MCL)
  - Acute myeloid leukemia (AML) that meets all of the following:
    - Newly diagnosed
    - FLT3 mutation-positive disease as detected by an FDA-approved test
    - Must use Rydapt in combination with standard cytarabine and daunorubicin induction and cytarabine consolidation chemotherapy

**Additional information**

**Note:** If approved, coverage is provided for 1 year. For AML, Rydapt is limited to 6 cycles.

**Medically accepted indication**

This drug is only covered under Medicare Part D when it is used for a medically accepted indication. A medically accepted indication is a use of the drug that is *either*:

- approved by the Food and Drug Administration. (That is, the Food and Drug Administration has approved the drug for the diagnosis or condition for which it is being prescribed.)
- — or — supported by certain reference books. (These reference books are the American Hospital Formulary Service Drug Information and the DRUGDEX Information System)

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**New request**  
**Priority Health Precertification Documentation**

**A. What condition is this drug being requested for?**

- Acute myeloid leukemia
- Advanced systemic mastocytosis
  - Aggressive systemic mastocytosis (ASM)
  - Systemic mastocytosis with associated hematological neoplasm (SM-AHN)
  - Mast cell leukemia (MCL)
- Other – the patient’s condition is: \_\_\_\_\_

**For Acute Myeloid Leukemia (AML)**

**A. Is the patient newly diagnosed?**

- Yes
- No. *Rationale for use:* \_\_\_\_\_

**B. Does the patient have FLT3 mutation-positive disease as detected by an FDA-approved test?**

- Yes
- No. *Rationale for use:* \_\_\_\_\_

**C. Is Rydapt being used in combination with standard cytarabine and daunorubicin induction and cytarabine consolidation chemotherapy?**

- Yes
- No. *Rationale for use:* \_\_\_\_\_

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**Priority Health Medicare exception request**

**Do you believe one or more of the prior authorization requirements should be waived?**  Yes  No

If yes, you must provide a statement explaining the medical reason why the exception should be approved.

**Would Rydapt likely be the most effective option for this patient?**

- No
- Yes, because: \_\_\_\_\_

**If the patient is currently using Rydapt, would changing the patient’s current regimen likely result in adverse effects for the patient?**

- No
- Yes, because: \_\_\_\_\_