



Vidaza [azacitidine]
Prior Authorization Request

Your patient's benefit plan requires prior authorization for certain medications. In order to make appropriate medical necessity determinations, your patient's diagnosis and other clinical information is required. Please complete the information requested on the form below and fax this form to Priority Partners, toll-free at 1-866-212-4756 to initiate the review process.

Patient's Name: Date:
Patient's ID: Patient's Date of Birth:
Physician's Name:
Specialty: NPI#:
Physician Office Telephone: Physician Office Fax:

Referring Provider Info: Same as Requesting Provider
Name: NPI#:
Fax: Phone:

Rendering Provider Info: Same as Referring Provider Same as Requesting Provider
Name: NPI#:
Fax: Phone:

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

Patient Weight: kg
Patient Height: cm

Please indicate the place of service for the requested drug:

- Ambulatory Surgical Home Off Campus Outpatient Hospital
On Campus Outpatient Hospital Office

Drug Information:

Strength/Measure Units ml Gm mg ea Un
Directions(sig) Route of administration
Dosing frequency

Criteria Questions:

- A. What is the prescribed drug? Vidaza azacitidine Other
B. What is the ICD-10 code?
1. What is the diagnosis?
Myelodysplastic syndrome (MDS) (If checked, go to 2)

Send completed form to: Priority Partners Fax: 1-866-212-4756

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited.

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- Acute myeloid leukemia (AML) *(If checked, go to 2)*
  - Accelerated phase or blast phase myelofibrosis *(If checked, go to 2)*
  - Blastic plasmacytoid dendritic cell neoplasm (BPDCN) *(If checked, go to 4)*
  - Myelodysplastic syndrome (MDS)/Myeloproliferative Neoplasms (MPN) Overlap Neoplasms (i.e. chronic myelomonocytic leukemia (CMML), juvenile myelomonocytic leukemia (JMML), BCR-ABL negative atypical chronic myeloid leukemia (aCML), MDS/MPN with neutrophilia, unclassifiable MDS/MPN, or MDS/MPN with ring sideroblasts and thrombocytosis) *(If checked, go to 2)*
  - Other, please specify. \_\_\_\_\_ *(If checked, no further questions)*
2. Is the patient currently receiving treatment with the requested medication?
- Yes, *Continue to 3*
  - No, *No Further Questions*
3. Is there evidence of unacceptable toxicity or disease progression while on the current regimen?
- Yes, *No Further Questions*
  - No, *No Further Questions*
4. Is the patient currently receiving treatment with the requested medication?
- Yes, *Continue to 5*
  - No, *Continue to 6*
5. Is there evidence of unacceptable toxicity or disease progression while on the current regimen?
- Yes, *No Further Questions*
  - No, *No Further Questions*
6. Does the patient have relapsed or recurrent disease?
- Yes, *Continue to 8*
  - No, *Continue to 7*
7. Is the requested drug being used for systemic disease with palliative intent?
- Yes, *Continue to 8*
  - No, *Continue to 8*
8. Will the requested medication be used in combination with venetoclax (Venclexta)?
- Yes, *No Further Questions*
  - No, *No Further Questions*

***I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by Priority Partners.***

X \_\_\_\_\_  
**Prescriber or Authorized Signature**

\_\_\_\_\_  
**Date (mm/dd/yy)**

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