

Dacogen [decitabine]

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. If you have questions regarding the prior authorization please contact Priority Partners at 888-819-1043 Option 4.

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 Pro	ovider
Name:	NPI#:
Fax:	Phone:
Rendering Provider Info: □ Same as Referring Prov	
Name:	NPI#:
Fax:	Phone:
	nits in accordance with FDA-approved labeling, r evidence-based practice guidelines.
Patient Weight:kg	
Patient Height:cm	
Please indicate the place of service for the requested dr. Ambulatory Surgical On Campus Outpatient Hospital Office	rug: ☐ Off Campus Outpatient Hospital
Drug Information:	
Strength/Measure	<i>Units</i> □ ml □ Gm □ mg □ ea □ Un
Directions(sig)	
Dosing frequency	
Dosing frequency	_
Criteria Questions:	
A. What drug is being prescribed? □ Dacogen □ decitabine HCPCS code J0894 (manufacturer no decitabine HCPCS code J0893 (Sun Pharma) □ Other	ot otherwise specified below)
B. What is the ICD-10 code?	

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. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. JHHC Dacogen [decitabine] SGM 2288-A – 07/2023.

1. What is the patient's diagnosis?	
☐ Myelodysplastic syndrome (MDS) (<i>If checked</i> ,	go to 2)
☐ Acute myeloid leukemia (AML) (If checked, go	o to 2)
☐ Accelerated phase or blast phase myelofibrosis	s (If checked, go to 2)
myelomonocytic leukemia [CMML], BCR-ABL i MDS/MPN with neutrophilia, unclassifiable MDS (<i>If checked, go to 9</i>)	neoplasm (MDS/MPN) overlap neoplasms (i.e. chronic negative atypical chronic myeloid leukemia [aCML], S/MPN, MDS/MPN with ring sideroblasts and thrombocytosis)
☐ Other, please specify.	(If checked, no further questions)
2. Is the patient currently receiving treatment with ☐ Yes, Continue to 3 ☐ No, No Further Questions	n the requested medication?
3. Is there evidence of unacceptable toxicity or dis ☐ Yes, No Further Questions ☐ No, No Further Questions	sease progression on the current regimen?
4. Is the patient currently receiving treatment with ☐ Yes, Continue to 5 ☐ No, Continue to 6	n the requested medication?
5. Is there evidence of unacceptable toxicity or dis ☐ Yes, No Further Questions ☐ No, No Further Questions	sease progression on the current regimen?
 6. Does the patient have relapsed or refractory dis ☐ Yes, Continue to 8 ☐ No, Continue to 7 	ease?
7. Is the requested drug being used for systemic d ☐ Yes, <i>Continue to 8</i> ☐ No, <i>Continue to 8</i>	isease with palliative intent?
8. Will the requested medication be used in comba ☐ Yes, No Further Questions ☐ No, No Further Questions	ination with venetoclax (Venclexta)?
9. Is the patient currently receiving treatment with ☐ Yes, Continue to 10 ☐ No, No Further Questions	n the requested medication?
10. Is there evidence of unacceptable toxicity or d ☐ Yes, No Further Questions ☐ No, No Further Questions	lisease progression on the current regimen?

I attest that this information is accurate and true, and that docume information is available for review if requested by Priority Partner	
X	Date (mandal) m
Prescriber or Authorized Signature	Date (mm/dd/yy)

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