

## SPECIALTY GUIDELINE MANAGEMENT

### VIDAZA (azacitidine) azacitidine (generic)

#### POLICY

##### I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

###### A. FDA-Approved Indications

Myelodysplastic syndromes (MDS): azacitidine/Vidaza is indicated for treatment of patients with the following French-American-British (FAB) myelodysplastic syndrome subtypes: refractory anemia (RA) or refractory anemia with ringed sideroblasts (if accompanied by neutropenia or thrombocytopenia or requiring transfusions), refractory anemia with excess blasts (RAEB), refractory anemia with excess blasts in transformation (RAEB-T), and chronic myelomonocytic leukemia (CMML).

###### B. Compendial Uses

1. Acute myeloid leukemia (AML)
2. Accelerated phase or blast phase myelofibrosis
3. Blastic plasmacytoid dendritic cell neoplasm (BPDCN)
4. Myelodysplastic syndrome (MDS)/Myeloproliferative Neoplasms (MPN) Overlap Neoplasms

All other indications are considered experimental/investigational and not medically necessary.

##### II. CRITERIA FOR INITIAL APPROVAL

###### A. **Myelodysplastic syndromes (MDS)**

Authorization of 12 months may be granted for the treatment of MDS.

###### B. **Acute myeloid leukemia (AML)**

Authorization of 12 months may be granted for the treatment of AML.

###### C. **Accelerated phase or blast phase myelofibrosis**

Authorization of 12 months may be granted for the treatment of accelerated phase or blast phase myelofibrosis.

###### D. **Blastic plasmacytoid dendritic cell neoplasm (BPDCN)**

Authorization of 12 months may be granted for the treatment of relapsed or refractory disease in combination with venetoclax.

###### E. **Myelodysplastic syndrome (MDS)/Myeloproliferative Neoplasms (MPN) Overlap Neoplasms**

Authorization of 12 months may be granted for the treatment of MDS/MPN overlap neoplasms (i.e. chronic myelomonocytic leukemia (CMML), BCR-ABL negative atypical chronic myeloid leukemia (aCML), unclassifiable MDS/MPN, or MDS/MPN with ring sideroblasts and thrombocytosis).

<b>Reference number</b>
2280-A

### III. CONTINUATION OF THERAPY

Authorization of 12 months may be granted for continued treatment in members requesting reauthorization for an indication listed in Section II when there is no evidence of unacceptable toxicity or disease progression while on the current regimen.

### IV. REFERENCES

1. Vidaza [package insert]. Summit, NJ: Celgene Corporation; March 2020.
2. Azacitidine injection [package insert]. Parsippany, NJ: Actavis Pharma Inc.; July 2020.
3. National Comprehensive Cancer Network. The NCCN Drugs & Biologics Compendium. <http://www.nccn.org>. Accessed July 15, 2020.