SPECIALTY QUANTITY LIMIT PROGRAM

SPRYCEL (dasatinib)

I. PROGRAM DESCRIPTION

The standard limit is designed to allow a quantity sufficient for the most common uses of the medication. The recommended dosing parameters for all FDA-approved indications fall within the standard limits. Coverage of an additional quantity may be reviewed on a case-by-case basis upon request.

II. COVERED QUANTITIES

| Medication | Standard Limit | FDA-recommended dosing |
|------------------------------------|---------------------------|---|
| Sprycel (dasatinib) 20 mg tablets | 90 tablets per 30 days | Adult patients: Chronic phase CML: 100 mg orally once daily Accelerated phase CML, myeloid or lymphoid blast phase CML: 140 mg orally once daily Ph+ ALL: 140 mg orally once daily |
| Sprycel (dasatinib) 50 mg tablets | 30 tablets per 30 days | |
| Sprycel (dasatinib) 70 mg tablets | 30 tablets per 30 days | |
| Sprycel (dasatinib) 80 mg tablets | 30 tablets per 30 days | |
| Sprycel (dasatinib) 100 mg tablets | 30 tablets per 30 days | Pediatric patients with chronic phase CML or Ph+ ALL: 40 mg -100 mg based on body weight |
| Sprycel (dasatinib) 140 mg tablets | 30 tablets per 30 days | |

* ALL: acute lymphoblastic leukemia; CML: chronic myeloid leukemia; Ph+: Philadelphia chromosome positive

III. REFERENCES

1. Sprycel [package insert]. Princeton, NJ: Bristol-Myers Squibb Company; December 2018.

Specialty Quantity Limit Sprycel 1722-H P2019

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