Reference number(s)
1686-H

SPECIALTY QUANTITY LIMIT PROGRAM

ZELBORAF (vemurafenib)

I. PROGRAM DESCRIPTION

The standard limit is designed to allow a quantity sufficient for the most common uses of the medication. If the member's plan allows a quantity limit exception review for the requested medication, coverage of an additional quantity may be provided up to the exception limit with prior authorization.

II. COVERED QUANTITIES

Medication	Standard Limit	Exception Limit*	FDA-recommended dosing
Zelboraf (vemurafenib) 240 mg tablet	240 per 30 days	300 per 30 days	• 960 mg (four tablets) every 12 hours • 1200 mg (five tablets) every 12 hours if co-administered with a strong CYP3A4 inducer

*Coverage up to the exception limits may be provided with prior authorization via the Specialty Post Limit Quantity Exception Criteria for approval.

III. REFERENCES

1. Zelboraf [package insert]. South San Francisco, CA: Genentech USA, Inc.; November 2017.

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