Reference number(s)
3056-H

SPECIALTY QUANTITY LIMIT PROGRAM

RUCONEST (C1 esterase inhibitor [recombinant])

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 the treatment of hereditary angioedema 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
Ruconest 2100 unit vial for reconstitution	60 vials in 90 days	For patients weighing less than 84 kg- 50 units per kg For patients weighing 84 kg or more- 4200 units (2 vials) Do not exceed 4200 units per dose. No more than two doses should be administered within a 24 hour period

III. REFERENCES

1. Ruconest [package insert]. Bridgewater, NJ: Pharming Healthcare, Inc.; March 2018.

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