

Kalbitor (ecallantide)

Effective Date: 10/22/13

Date Developed: 9/3/13 by Albert Reeves MD Last Approval Date: 1/26/16, 1/24/17, 1/23/18, 1/22/19

(Archived 1/22/19)

Kalbitor is a human plasma kallikrein inhibitor aimed at decreasing bradykinin production

Pre-Authorization Criteria:

VCHCP will authorize Kalbitor for the following: acute attacks of hereditary angioedema (HAE) in patients 16 years of age and older.

Dosing: 30 mg subcutaneously; may repeat once within 24 hours

PRECAUTIONS: hypersensitivity; nausea; headache; injection site reactions (pain; pruritis;

edema)

DRUG INTERACTIONS: none known

REFERENCES

- 1. Caballero T, Farkas H, Bouillet L, et al, "International Consensus and Practical Guidelines on the Gynecologic and Obstetric Management of Female Patients With Hereditary Angioedema Caused by C1 Inhibitor Deficiency," *J Allergy Clin Immunol*, 2012, 129(2):308-20.
- 2. Craig T, Riedl M, Dykewicz MS, et al, "When is Prophylaxis for Hereditary Angioedema Necessary?" *Ann Alllergy Ashtma Immunol*, 2009, 102(5):366-72.
- 3. Epstein TG and Bernstein JA. "Current and Emerging Management Options for Hereditary Angioedema in the US," *Drugs*, 2008, 68(18):2561-73.

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Revision Date	Content Revised (Yes/No)	Contributors	Review/Revision Notes
1/24/17	No	Catherine Sanders, MD;	Annual review
		Robert Sterling, MD	
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