

<b>Clinical Policy Title:</b>	propranolol HCl Oral Solution
<b>Policy Number:</b>	RxA.389
<b>Drug(s) Applied:</b>	Hemangeol®
<b>Original Policy Date:</b>	03/06/2020
<b>Last Review Date:</b>	09/14/2020
<b>Line of Business Policy Applies to:</b>	All lines of business

## Background

Propranolol HCl oral solution (Hemangeol®) is a beta-adrenergic blocker. Hemangeol oral solution is indicated for the treatment of proliferating infantile hemangioma requiring systemic therapy.

## Dosing Information

Drug Name	Indication	Dosing Regimen	Maximum Dose
Propranolol HCl Oral Solution (Hemangeol®)	Proliferating infantile hemangioma	Starting dose is 0.15 mL/kg (0.6 mg/kg) PO twice daily . After 1 week, increase to 0.3 mL/kg (1.1 mg/kg) twice daily. After 2 weeks, increase to a maintenance dose of 0.4 mL/kg (1.7 mg/kg) twice daily	Depends on weight

## Dosage Forms

- Oral solution: 4.28 mg/mL

## Clinical Policy

Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

### I. Initial Approval Criteria

#### A. Proliferating Infantile Hemangioma (must meet all):

1. Diagnosis of proliferating infantile hemangioma;
2. Age 5 weeks of age or older;
3. Weight is 2 kg or more;

#### Approval duration

**Commercial:** 6 months

**Medicaid:** 6 months

**HIM:** 6 months

This clinical policy has been developed to authorize, modify, or determine coverage for individuals with similar conditions. Specific care and treatment may vary depending on individual need and benefits covered by the plan. This policy is not intended to dictate to providers how to practice medicine, nor does it constitute a contract or guarantee regarding payment or results. This document may contain prescription brand name drugs that are trademarks of pharmaceutical manufacturers that are not affiliated with RxAdvance.

## II. Continued Therapy Approval

### A. Proliferating Infantile Hemangioma (must meet all):

1. Member is currently receiving medication that has been authorized by RxAdvance or the member has met initial approval criteria listed in this policy;
2. Member is responding positively to therapy;
3. Member meets one of the following (a or b):
  - a. Member has not received  $\geq 12$  months of consecutive therapy;
  - b. Documentation supports recurrence of hemangioma.

#### Approval duration

**Commercial:** 6 months

**Medicaid:** 6 months

**HIM:** 6 months

## III. Appendices

### APPENDIX A: Abbreviation/Acronym Key

FDA: Food and Drug Administration

HCl: hydrochloride

IH: infantile hemangioma

### APPENDIX B: Therapeutic Alternatives

Not applicable

### APPENDIX C: Contraindications/Boxed Warnings

- Contraindication(s):
  - Asthma or history of bronchospasm;
  - Pheochromocytoma;
  - Premature infants with corrected age  $<5$  weeks;
  - Infants weighing less than 2 kg;
  - Known hypersensitivity to propranolol or excipients;
  - Bradycardia ( $<80$  beats per minute), greater than first degree heart block, decompensated heart failure;
  - Blood pressure less than 50/30mmHg;
- Boxed warning(s):
  - None

### APPENDIX D: General Information

Management of IH

- IHs are the most common tumors of childhood. While they often involute after proliferation, there are some that rapidly develop complications, resulting in pain, functional impairment, or permanent disfiguration. For such complicated cases of IH, propranolol is a first-line medical therapy.
- Although the most dramatic improvement using propranolol for IH occurs within 3 to 4 months of initiation of therapy, the optimal treatment duration has not been established:
  - The FDA recommends the maintenance dose be maintained for 6 months. This is likely based on the clinical trial for approval which evaluated patients after 6 months of treatment.

- The American Academy of Pediatrics indicates that many continue therapy until patients reach an age when IH would normally begin to regress without treatment- often until at least 8 to 12 months of age, which, in most studies, equated to 3 to 12 months of therapy.
- While Hemangeol is effective, rebound growth has been observed in 6% to 25% of children. In the Hemangeol clinical trial, 10% of patients deemed successes after 6months of therapy later required re-treatment for recurrence.

**References**

1. Hemangeol Prescribing Information. Parsippany, NJ: Pierre Fabre Pharmaceuticals, Inc; April 2020. Available at: <https://www.hemangeol.com/hcp/wp-content/themes/responsive-child/docs/HEMANGEOL-Notice-USA-May-12-2020.pdf> . Accessed July 14, 2020.
2. Darrow DH, Greene AK, Mancini AJ, et al. American Academy of Pediatrics clinical report (guidance for the clinician in rendering pediatric care): diagnosis and management of infantile hemangioma. Pediatrics. 2015; 136(4): e1060-e1104.
3. Hemangeol. Lexi-Drug. Lexicomp. Wolters Kluwer Health, Inc. Riverwoods, IL. Accessed with subscription at: <http://online.lexi.com>. Accessed July 14, 2020.
4. Krowchuk DP, Frieden IJ, Mancini AJ, et al: Clinical practice guideline for the management of infantile hemangiomas. Pediatrics 2019; 143(1):e20183475.

Review/Revision History	Review/Revised Date	P&T Approval Date
Policy established.	01/2020	03/06/2020
Policy was reviewed: <ol style="list-style-type: none"> <li>1) Policy description table updated</li> <li>2) Dosing Information updated</li> <li>3) Continuation therapy criteria II.A.1. rephrased to “Member is currently receiving medication that has been authorized by RxAdvance or the member has met initial approval criteria listed in this policy”</li> <li>4) Appendix C, contraindications updated</li> <li>5) Initial therapy and continued therapy approval duration was updated</li> <li>6) References were updated</li> </ol>	07/14/2020	09/14/2020