

Division: Pharmacy Policy	Subject: State of Florida's Agency for Health Care Administration's
	Prior Authorization Criteria
Original Development Date:	December 2, 2014
Original Effective Date:	
Revision Date:	

HEMANGEOL® (propranolol oral solution)

LENGTH OF AUTHORIZATION:

Initial Review: 6 months

Re-treatment if necessary: 6 months

INITIAL REVIEW CRITERIA:

- 1. Infant has a diagnosis of proliferating infantile hemangioma
- 2. Infant's age is in the range of 5 weeks (adjusted gestational age) to 5 months
- 3. Infant weighs a minimum of 2 kilograms
- 4. Infant has <u>none</u> of the contraindications as listed below:
 - Contraindications:
 - Known hypersensitivity to propranolol or excipients
 - Asthma or history of bronchospasm
 - o Bradycardia (< 80 beats per minute)
 - Greater than first degree heart block
 - Decompensated heart failure
 - Blood pressure < 50/30 mmHg
 - Pheochromocytoma

RE-TREATMENT REVIEW CRITERIA:

1. Patient had initial successful treatment with Hemangeol for 6 months resulting in complete or nearly complete resolution of the target hemangioma but has experienced a recurrence.

DOSING & ADMINISTRATION:

- Starting dose is 0.6 mg/kg twice daily
- After one week, increase the dose to 1.1 mg/kg twice daily
- After two weeks of treatment, increase the dose to 1.7 mg/kg twice daily and maintain for 6 months
- Heart rate and blood pressure should be monitored for two hours after initiation of drug and with all dose increases
- Twice daily doses must be administered a minimum of nine hours apart
- To minimize the risk of hypoglycemia, administer during or right after a feeding
- Readjust the dose periodically as the child's weight increases