

**Clinical Policy: Brimonidine Tartrate (Mirvaso)** 

Reference Number: PA.CP.PMN.192

Effective Date: 4.17.19 Revision Log

Last Review Date: 04.19

# **Description**

Brimonidine Tartrate (Mirvaso<sup>®</sup>) is a relatively selective alpha-2 adrenergic agonist topical gel. It may reduce erythema through direct vasoconstriction.

# FDA Approved Indication(s)

Mirvaso is indicated for the topical treatment of persistent (nontransient) facial erythema of rosacea in adults 18 years of age or older.

# Policy/Criteria

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

It is the policy of health plans affiliated with PA Health & Wellness<sup>®</sup> that Mirvaso is **medically necessary** when the following criteria are met:

# I. Initial Approval Criteria

- A. Facial Erythema Associated with Rosacea (must meet all):
  - 1. Diagnosis of persistent facial erythema associated with rosacea;
  - 2. Age  $\geq$  18 years;
  - 3. If papules or pustules are present, a failure of or concomitant treatment with any of the following agents, unless contraindicated or clinically significant adverse effects are experienced: topical metronidazole, oral doxycycline or Finacea;
  - 4. Dose does not exceed 30 mg (1 tube) per month.

**Approval duration:** 12 Months

# **B.** Other diagnoses/indications:

1. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): PA.CP.PMN.53 for Medicaid.

# **II.** Continued Therapy

- A. Facial Erythema Associated with Rosacea (must meet all):
  - 1. Currently receiving medication via PA Health & Wellness benefit and documentation supports positive response to therapy or the Continuity of Care policy (PA.LTSS.PHAR.01) applies;
  - 2. Member is responding positively to therapy;
  - 3. If request is for a dose increase, new dose does not exceed 30 mg (1 tube) per month.

**Approval duration:** 12 Months

# **B. Other diagnoses/indications** (must meet 1 or 2):

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- 1. Currently receiving medication via PA Health & Wellness benefit and documentation supports positive response to therapy or the Continuity of Care policy (PA.LTSS.PHAR.01) applies.
  - Approval duration: Duration of request or 12 months (whichever is less); or
- 2. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): PA.CP.PMN.53 for Medicaid.

# III. Diagnoses/Indications for which coverage is NOT authorized:

**A.** Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policy – PA.CP.PMN.53 for Medicaid or evidence of coverage documents.

### IV. Appendices/General Information

Appendix A: Abbreviation/Acronym Key FDA: Food and Drug Administration

# *Appendix B: Therapeutic Alternatives*

This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may not be a formulary agent for all relevant lines of business and may require prior authorization.

Drug Name	Dosing Regimen	Dose Limit/	
		Maximum Dose	
metronidazole	Apply thin film topically to	No maximum dosage	
(Metrocream <sup>®</sup> 0.75%,	affected area QD for 1% and	information is available.	
Metrogel <sup>®</sup> 1%, Metrolotion <sup>®</sup>	BID for 0.75%		
0.75%)			
Finacea® (15% gel)	Apply in a thin film	No maximum dosage	
(azelaic acid)	topically to the affected area	information is available.	
	BID		
	Reassess if no improvement		
	in 12 weeks.		
doxycycline (Oracea)®	Lesions (papules and	300 mg/day;	
	pustules): 40 mg PO once	40 mg/day for Oracea	
	daily in the morning (1 hour		
	before or 2 hours after a		
	meal)		

Therapeutic alternatives are listed as Brand name® (generic) when the drug is available by brand name only and generic (Brand name®) when the drug is available by both brand and generic.

# Appendix C: Contraindications/Boxed Warnings

# Appendix D: General Information

• Tetracycline agents, including doxycycline and minocycline exhibit anti-inflammatory activities at doses < 50 mg. Anti-inflammatory dose doxycycline does not exert antibiotic selection pressure and thus does not induce antibiotic resistance; its mechanism of action

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in rosacea appears to relate to the anti-inflammatory and biological activities of doxycycline.

V. Dosage and Administration

Indication	Dosing Regimen	<b>Maximum Dose</b>
Facial erythema	Apply a pea-size amount topically QD to	One
associated with	each of the five areas of the face (forehead,	application/day
rosacea	chin, nose, each cheek) avoiding the eyes	
	and lips.	

#### VI. Product Availability

Gel (30 gm tube or pump): 0.33%

### VII. References

- 1. Mirvaso Prescribing Information. Fort Worth, TX: Galderma Laboratories; November 2017. Available at: www.mirvaso.com. Accessed January 31, 2019.
- 2. Fowler J Jr, et al. Efficacy and safety of once-daily topical brimonidine tartrate gel 0.5% for the treatment of moderate to severe facial erythema of rosacea: results of two randomized, double-blind, and vehicle-controlled pivotal studies. J Drugs Dermatol. Jun 2013; 12(6):650-6.
- 3. Micromedex<sup>®</sup> Healthcare Series [database online]. Greenwood Village, Colorado: Thomson Healthcare. Updated periodically. Accessed January 31, 2019.
- 4. Clinical Pharmacology [database online]. Tampa, FL: Gold Standard, Inc.; 2016. Available at: http://www.clinicalpharmacology-ip.com. Accessed January 31, 2019.
- 5. National Rosacea Society. Rosacea treatment algorithms. Available at: <a href="https://www.rosacea.org/physicians/treatmentalgorithms">https://www.rosacea.org/physicians/treatmentalgorithms</a>. January 31, 2019.
- 6. Scaller M, et al. Rosacea treatment update: Recommendations from the global ROSacea Consensus (ROSCO) panel. *Br J Dermatol* 2016. Epub ahead of print. doi: 10.1111/bjd.15173.

Reviews, Revisions, and Approvals		P&T
		Approval Date
Policy created. 2Q 2019 annual review: policy split from PA.CP.PMN.86 Brimonidine (Mirvaso), Oxymetazoline (Rhofade) into individual drug policies; added age limit; references reviewed and updated.	04.17.19	