



# SmartPA Criteria Proposal

Drug/Drug Class:	Corticosteroids, Ophthalmic "Soft" PDL Edit		
First Implementation Date:	July 11, 2019		
Revised Date:	July 7, 2022		
Prepared For:	MO HealthNet		
Prepared By:	MO HealthNet/Conduent		
Criteria Status:	⊠Existing Criteria □Revision of Existing Criteria □New Criteria		

#### **Executive Summary**

Purpose: The MO HealthNet Pharmacy Program will implement a state-specific preferred drug list.

Why Issue Selected: Topical corticosteroids are effective in reducing anterior segment inflammation but are associated with adverse drug reactions including elevation of intraocular pressure and cataract formulation. Newer "soft" ophthalmic corticosteroids have been developed with improved therapeutic indices through retrometabolic drug design. The retrometabolic drug design principles appear to achieve the necessary balance between solubility/lipophilicity, tissue distribution, glucocorticoid receptor binding, and metabolic deactivation to be effective as a topical ophthalmic steroid. The "soft" steroids are safe and effective in treating a wide variety of ocular inflammatory conditions including giant papillary conjunctivitis, seasonal allergic conjunctivitis, and uveitis as well as in the treatment of ocular inflammation and pain following cataract surgery. Clinical studies have confirmed the retrometabolic design of "soft" steroids minimized adverse reactions such as cataract formation and intraocular pressure elevation.

Total program savings for the PDL classes will be regularly reviewed.

## Program-Specific Information:

Preferred Agents	Non-Preferred Agents
Durezol®	Alrex®
Lotemax® Gel/Susp	Difluprednate
·	Eysuvis®
	Inveltys®
	Lotemax® Oint
	Lotemax® SM
	<ul> <li>Loteprednol</li> </ul>

Type of Criteria:	☐ Increased risk of ADE	□ Preferred Drug List		
	☐ Appropriate Indications	☐ Clinical Edit		
Data Sources:	☐ Only Administrative Databases	□ Databases + Prescriber-Supplied		

### **Setting & Population**

- Drug class for review: Corticosteroids, Ophthalmic "Soft"
- Age range: All appropriate MO HealthNet participants

#### **Approval Criteria**

- Failure to achieve desired therapeutic outcomes with trial on 2 or more preferred agents
  - Documented trial period of preferred agents OR
  - Documented ADE/ADR to preferred agents

#### **Denial Criteria**

· Lack of adequate trial on required preferred agents

Therapy will be denied if all approval criteria are not met							
Required Documentation							
Laboratory Results: MedWatch Form:		Progress Notes: Other:					
Disposition of Edit							
Denial: Exception Code "0160" (Preferred Drug List) Rule Type: PDL							

#### **Default Approval Period**

1 year

#### References

- Evidence-Based Medicine and Fiscal Analysis: "Ophthalmic Soft Corticosteroids Therapeutic Class Review", Conduent Business Services, L.L.C., Richmond, VA; January 2022.
- Evidence-Based Medicine Analysis: "Corticosteroids, Ophthalmic Soft Steroids", UMKC-DIC; February 2022.
- USPDI, Micromedex; 2022.
- Facts and Comparisons eAnswers (online); 2022 Clinical Drug Information, LLC.