

Policies Repository



Policy Title Olopatadine Hydrochloride (Pataday™) 0.2 Percent

Policy Number FS.CLIN.6

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This Pharmacy Policy document describes the status of pharmaceutical information and/or technology at the time the document was developed. Since that time, new information relating to drug efficacy, interactions, contraindications, dosage, administration routes, safety or FDA approval may have changed. If the Medical/Pharmacy Reviewer is aware of any new information on the subject of this document, please provide it promptly to the Medical/Pharmacy Policy Department. This information may include new FDA approved indications, withdrawals or other FDA alerts. This type of information is relevant not only when considering whether this Policy should be updated, but also when applying it to current requests for coverage.

Members are advised to use participating pharmacies in order to receive the highest level of benefits.

Policy **Olopatadine hydrochloride (Pataday™) 0.2 percent** is indicated for the temporary prevention of itchy eyes due to allergic conjunctivitis.

The use of olopatadine hydrochloride (Pataday™) 0.2 percent requires prior authorization (ie, clinical pharmacy and/or Medical Director review).

Policy Description

Olopatadine hydrochloride (Pataday™) 0.2 percent is an inhibitor of histamine release from the mast cell and a relatively selective histamine H1-antagonist that inhibits the in vivo and in vitro type 1 immediate hypersensitivity reaction. It is devoid of effects on alpha-adrenergic, dopamine, muscarinic type 1 and 2, and serotonin receptors. **Olopatadine hydrochloride (Pataday™) 0.2 percent** contains the same active ingredient as Patanol and has not been shown to have superior efficacy to Patanol.

Policy Guideline Inclusion

Olopatadine hydrochloride (Pataday™) 0.2 percent is approved when **all** of the following inclusion criteria are met:

- Documentation of allergic conjunctivitis
- Documentation of trial and failure or contraindication to all of the following agents:
 - Olopatadine hydrochloride ophthalmic solution (Patanol®)
 - Azelastine hydrochloride ophthalmic solution (Optivar®)

Policy Guideline Exclusion

Olopatadine hydrochloride (Pataday™) 0.2 percent is denied when **any** of the following exclusion criteria are present:

- No documentation of allergic conjunctivitis
- No documentation of trial and failure or contraindication to all of the following agents:
 - Olopatadine hydrochloride ophthalmic solution (Patanol®)
 - Azelastine hydrochloride ophthalmic solution (Optivar®)

Policy List of Applicable Drugs

Brand Name	Generic Name
Pataday	Olopatadine Hydrochloride

Dosing and Administration

Refer to the specific manufacturer's prescribing information for administration and dosage details for each specific agent.

Policy References

Epocrates. Pataday™ (Olopatadine Hydrochloride Ophthalmic Solution) 0.2 Percent. [Epocrates Web site]. 2006. Available at: <https://online.epocrates.com/noFrame/#container> (via subscription only). Accessed November 18, 2008.

Pataday™ (Olopatadine Hydrochloride Ophthalmic Solution) 0.2 Percent [package insert]. Fort Worth, TX: Alcon; 2008.

Slonim CB, Bell BG, Favetta JR, Bielory L, Siegel CS. Allergy: understanding the role of inflammation. The benefits of multi-action topical allergy remedies deserve to be more widely known, both in ophthalmology and in general medicine. Review of Ophthalmology. [Review of Ophthalmology Web site]. Available at: http://www.revophth.com/index.asp?page=1_13397.htm. Accessed July 28, 2009.

US Food and Drug Administration (FDA). Center for Drug Evaluation and Research. Application No. 21-545. Medical Review. Pataday™ (Olopatadine Hydrochloride Ophthalmic Solution) 0.2 Percent. [FDA web site]. 12/21/04. Available at: http://www.fda.gov/cder/foi/nda/2004/021545s000_MedR.pdf. Accessed November 18, 2008. [Link no longer active]

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