DORZOLAMIDE PF- dorzolamide pf solution/ drops Imprimis Rx NJ

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Store at 20° to 25° C (68° to 77° F)

Bottle Label



Ophthalmic Drops

Compounded for a licensed professional or patient use by



ImprimisRx NJ 1705 Route 46 West, Suite 6A Ledgewood, NJ 07852 (844)446-6979

NDC 70261-516-10

Each mL contains: Dorzolamide 20mg, Sodium Chloride 4.8 mg, Sodium citrate 1.5mg, Pluronic L64 0.001ml and Sterile Water q.s.

Sodium Hydroxide and/or Hydrochloric Acid may have been used to adjust pH.

Store at room temperature 20-25°C(68-77°F). This medicine was compounded for you at the direction of your prescriber. Protect from light. Rx Only - Not for resale

Lot#

Use By:

Date Compounded: mm/dd/yyyy hh:mm xx



DORZOLAMIDE PF

dorzolamide pf solution/ drops

Product Information

Product TypeHUMAN PRESCRIPTION DRUGItem Code (Source)NDC:70261-516

Route of Administration OPHTHALMIC

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

DORZOLAMIDE HYDROCHLORIDE (UNII: QZO5366EW7) (DORZOLAMIDE - UNII:9JDX055TW1)

DORZOLAMIDE | 20 mg in 1 mL

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# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:70261-516- 10	10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	0 1/0 1/20 18	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		0 1/0 1/20 18	

Labeler - Imprimis Rx NJ (931390178)

Revised: 5/2018 ImprimisRx NJ