
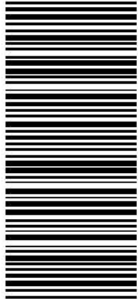


PREDNISOLONE ACETATE PF- prednisolone acetate pf suspension/ drops
ImprimisRx NJ

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

Sterile 5 ml Bottle		NDC 70261-501-05	
Prednisolone Acetate P-F1%			
Shake Well Before Use			
Ophthalmic Drops			
Compounded for a licensed professional or patient use by		Each mL contains: Prednisolone Acetate 10mg, Sodium Chloride, Polysorbate 80, Poloxamer 407, Edetate Disodium, Sodium Citrate, Glycerin and Sterile Water q.s. Sodium Hydroxide may have been used to adjust pH.	
		Store at controlled 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	
ImprimisRx NJ 1705 Route 46 West, Suite 6A Ledgewood, NJ 07852 (844) 446-6979		Lot # Use By:	
			

PREDNISOLONE ACETATE PF

prednisolone acetate pf suspension/ drops

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70261-501
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PREDNISOLONE ACETATE (UNII: 8B2807733D) (PREDNISOLONE - UNII:9PHQ9Y1OLM)	PREDNISOLONE ACETATE	10 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70261-501-05	5 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	01/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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unapproved drug other		01/01/2018	
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Labeler - Imprimis Rx NJ (931390178)

Revised: 5/2018

ImprimisRx NJ