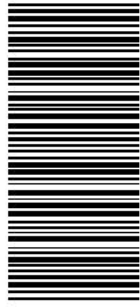


LATANOPROST PF- latanoprost pf solution/ drops
ImprimisRx NJ

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 under refrigeration 2° to 8° C (36° to 46° F)

Bottle Label

<p>Sterile 7.5ml Bottle</p> <div style="background-color: #00aaff; color: white; padding: 5px; font-weight: bold; font-size: 1.2em;">Latanoprost P-F</div> <div style="background-color: yellow; border: 1px solid black; padding: 2px; display: inline-block; margin: 5px 0;">0.005%</div> <p>Ophthalmic Drops Compounded for a licensed professional or patient use by</p> <div style="font-size: 1.5em; font-weight: bold; color: #00aaff;">imprimis</div> <div style="font-size: 2em; font-weight: bold; color: orange; margin-left: 5px;">Rx</div> <p>ImprimisRx NJ 1705 Route 46 West, Suite 6A Ledgewood, NJ 07852 (844) 446-6979</p>	<p>NDC 70261-515-07</p>	<p>Each mL contains: Latanoprost 0.05mg, Sodium Chloride 8mg, Sodium Phosphate Monobasic 1mg, Sodium Phosphate Dibasic 0.7mg, Pluronic L64 0.001ml, Polysorbate 80 NF 0.0005 ml and Sterile Water q.s. Sodium Hydroxide and/or Hydrochloric Acid may have been used to adjust pH.</p> <p>Store under refrigeration 2-8 C (36-46°F). This medicine was compounded for you at the direction of your prescriber. Protect from light. Rx Only - Not for resale</p>
		
		<p>Lot # Use By: Date Compounded: mm/dd/yyyy hh:mm xx</p>

LATANOPROST PF			
latanoprost pf solution/ drops			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70261-515
Route of Administration	OPHTHALMIC		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	LATANOPROST (UNII: 6Z5B6HVF6O) (LATANOPROST - UNII:6Z5B6HVF6O)	LATANOPROST	0.05 mg in 1 mL
Packaging			
#	Item Code	Package Description	Marketing Start Date
		Marketing End Date	
1	NDC:70261-515-07	7.5 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
unapproved drug other		01/01/2018	

Labeler - Imprimis Rx NJ (931390178)

Registrant - Imprimis Rx NJ (931390178)

Establishment

Name	Address	ID/FEI	Business Operations
Imprimis Pharmaceuticals		080431967	manufacture(70261-515)

Revised: 5/2018

Imprimis Rx NJ