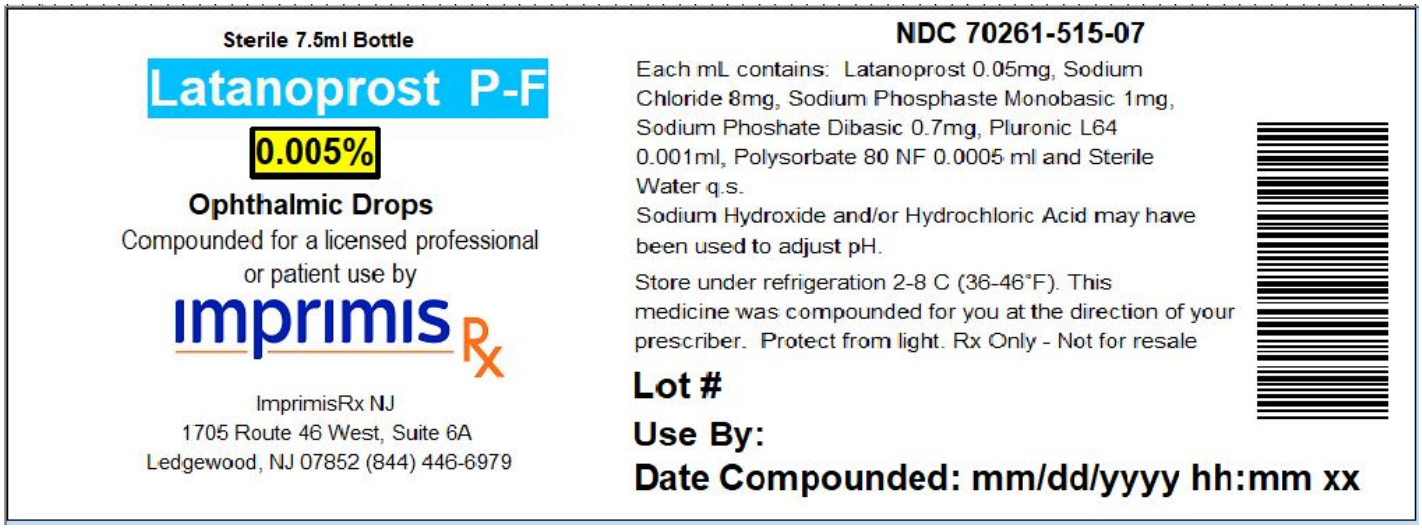
LATANOPROST PF- latanoprost pf solution/ drops
ImprimisRx NJ

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



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 - ImprimisRx NJ (931390178)

Registrant - ImprimisRx NJ (931390178)

Establishment

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

Revised: 5/2018 ImprimisRx NJ