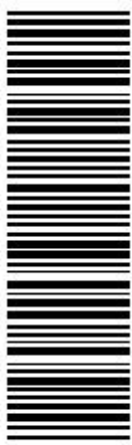



MOXIFLOXACIN PF- moxifloxacin pf injection, solution
Imprimis Rx NJ

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)

Vial Label

	MOXIFLOXACIN	NDC 70261-511-01
	(0.5)% SINGLE USE VIAL	Ingredients:
	Lot:01152018@3	Moxifloxacin HCl, edetate calcium disodium, Sodium Chloride, sodium hydroxide, Sterile water for injection
	Date Compounded: mm/dd/yyyy hh:mm xx	Store at controlled room temperature 20-25°C (68 -77°F)
	Use by: 06/26/2018 07:46 AM	This medicine was compounded for you at the direction of your prescriber.
 1705 Route 46 West, Suite 6A Ledgewood, NJ 07852 (973)328-8756	Rx Only - not for resale	

MOXIFLOXACIN PF				
moxifloxacin pf injection, solution				
Product Information				
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70261-511	
Route of Administration	INTRAOCCULAR			
Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
MOXIFLOXACIN HYDROCHLORIDE MONOHYDRATE (UNII: B8956S8609) (MOXIFLOXACIN - UNII:U188XYD42P)	MOXIFLOXACIN	5 mg in 1 mL		
Inactive Ingredients				
Ingredient Name	Strength			
SODIUM CHLORIDE (UNII: 451W47IQ8X)	8 mg in 1 mL			
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70261-511-01	1 mL in 1 VIAL, GLASS; 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(70261-511)

Revised: 5/2018

ImprimisRx NJ