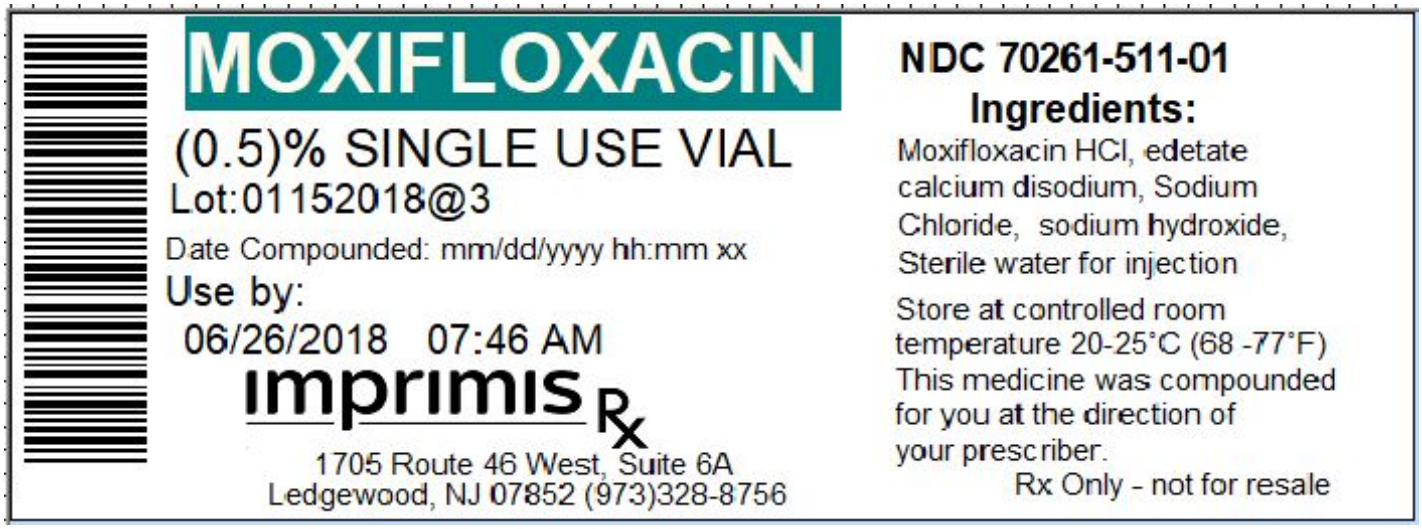
MOXIFLOXACIN PF- moxifloxacin pf injection, solution
ImprimisRx NJ

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)

Vial Label



MOXIFLOXACIN PF
moxifloxacin pf injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:70261-511
Route of Administration	INTRAOCULAR		

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