MOXIFLOXACIN PF- moxifloxacin pf injection, solution
Imprimis NJOF, LLC

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.

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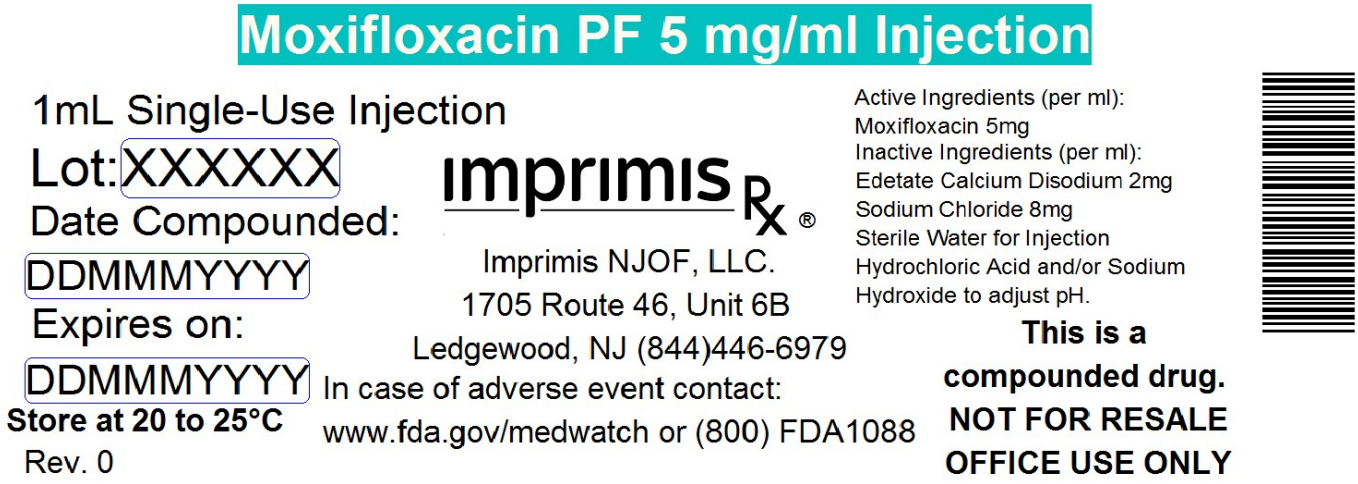
20 ^{0} to 25 ^{0} C (*68 ^{0} to 77 ^{0} F)

Vial Label



MOXIFLOXACIN PF
moxifloxacin pf injection, solution

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:71384-511
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MOXIFLOXACIN HYDROCHLORIDE MONOHYDRATE (UNII: B8956S8609) (MOXIFLOXACIN - UNII:U188XYD42P)	MOXIFLOXACIN	5 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71384-511-01	20 in 1 BOX	01/05/2018	
1		1 mL in 1 VIAL, SINGLE-DOSE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/05/2018	

Labeler - Imprimis NJOF, LLC (080431967)

Registrant - Imprimis NJOF, LLC (080431967)

Establishment

Name	Address	ID/FEI	Business Operations
Imprimis NJOF, LLC		080431967	manufacture(71384-511)

Revised: 3/2020 Imprimis NJOF, LLC