

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.

Store at 20° to 25° C (68° to 77° F)

Package Label

Moxifloxacin PF 5 mg/ml Injection

1mL Single-Use Injection

Lot: XXXXXX

Date Compounded:

DDMMYYYY

Expires on:

DDMMYYYY

Store at 20 to 25°C

Rev. 0

imprimis ^{Rx}®

Imprimis NJOF, LLC.

1705 Route 46, Unit 6B

Ledgewood, NJ (844)446-6979

In case of adverse event contact:

www.fda.gov/medwatch or (800) FDA1088

Active Ingredients (per ml):
Moxifloxacin 5mg
Inactive Ingredients (per ml):
Edetate Calcium Disodium 2mg
Sodium Chloride 8mg
Sterile Water for Injection
Hydrochloric Acid and/or Sodium Hydroxide to adjust pH.



**This is a
compounded drug.
NOT FOR RESALE
OFFICE USE ONLY**

MOXIFLOXACIN PF

moxifloxacin pf injection, solution

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:71384-509
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:71384-509-01	20 in 1 BOX	02/10/2020	
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		02/10/2020	

Labeler - Imprimis NJOF, LLC (080431967)

Registrant - Imprimis NJOF, LLC (080431967)

Revised: 3/2020

Imprimis NJOF, LLC