

**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 ° to 25 ° C (\*68 ° to 77 ° F)

**Vial 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** <sup>Rx</sup>®

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**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:71384-511
<b>Route of Administration</b>	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