

TRIAMCINOLONE-MOXIFLOXACIN PF- triamcinolone-moxifloxacin pf injection, suspension

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

NDC 71384-510-01

Triamcinolone-Moxifloxacin PF (15/1) mg/ml Injection

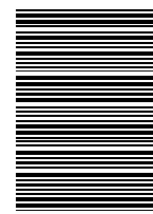
Shake Well

Volume: 0.6ml/vial

Quantity: 20

Lot: XXXXXX

imprimis ^{Rx}®



Date Compounded:

DDMMYYYY

Expires on:

DDMMYYYY

Imprimis NJOF, LLC.
1705 Route 46 West, Unit 6B
Ledgewood, NJ (844)446-6979

Store at 20 to 25°C

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

Rev. 2

**In case of adverse event contact:
www.fda.gov/medwatch or (800) FDA1088**

TRIAMCINOLONE-MOXIFLOXACIN PF

triamcinolone-moxifloxacin pf injection, suspension

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:71384-510
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRIAMCINOLONE ACETONIDE (UNII: F446C597KA) (TRIAMCINOLONE ACETONIDE - UNII:F446C597KA)	TRIAMCINOLONE ACETONIDE	15 mg in 1 mL
MOXIFLOXACIN HYDROCHLORIDE MONOHYDRATE (UNII: B8956S8609) (MOXIFLOXACIN - UNII:U188XYD42P)	MOXIFLOXACIN HYDROCHLORIDE MONOHYDRATE	1 mg in 1 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71384-510-01	20 in 1 BOX	01/05/2018	

1	0.6 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product		
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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)

Revised: 2/2020

Imprimis NJOF, LLC