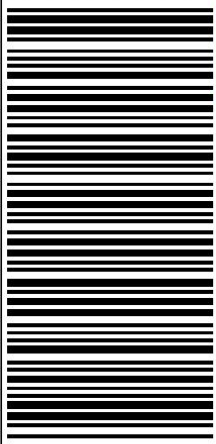


TRIAMCINOLONE-MOXIFLOXACIN PF- triamcinolone-moxifloxacin pf suspension
Imprimis Rx NJ

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)

	Triamcinolone-Moxifloxacin- PF		
	(15/1)mg/mL 0.6ml single dose vial NDC 70261-510-01		
	Lot:	Shake Well	Each mL contains:
	Use by:		Triamcinolone acetate, moxifloxacin, polysorbate 80, edetate calcium disodium, pluronic, sodium hydroxide, Sterile water for injection
imprimis _{Rx}		Store at controlled room temperature 20-25C. Rx only-not for resale	
1705 Route 46 West, Suite 6A Ledgewood, NJ 07852 (973)328-8756			
This medicine was compounded in our pharmacy for use by a licensed professional only			

TRIAMCINOLONE-MOXIFLOXACIN PF				
triamcinolone-moxifloxacin pf suspension				
Product Information				
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:70261-510	
Route of Administration	OPHTHALMIC			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
MOXIFLOXACIN HYDROCHLORIDE (UNII: C53598599T) (MOXIFLOXACIN - UNII:U188XYD42P)		MOXIFLOXACIN	1 mg in 1 mL	
TRIAMCINOLONE ACETONIDE (UNII: F446C597KA) (TRIAMCINOLONE ACETONIDE - UNII:F446C597KA)		TRIAMCINOLONE ACETONIDE	15 mg in 1 mL	
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70261-510-01	0.6 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)

Revised: 2/2018

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