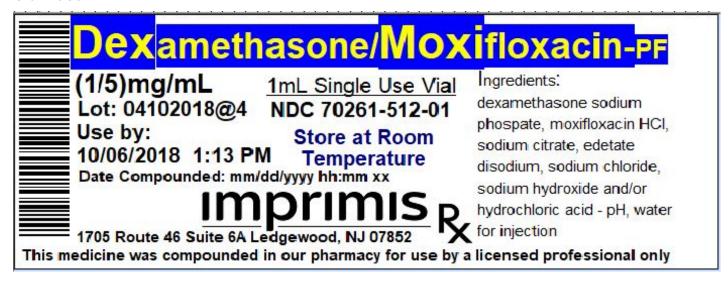
DEX-MOXI PF- dexamethas one - moxifloxacin pf injection, solution Imprimis Rx NJ

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)

Vial Label



DEX-MOXI PF

dexamethasone - moxifloxacin pf injection, solution

| Product Information | Product Information | | | | |
|-------------------------|-------------------------|--------------------|---------------|--|--|
| Product Type | HUMAN PRESCRIPTION DRUG | Item Code (Source) | NDC:70261-512 | | |
| Route of Administration | INTRAOCULAR | | | | |

| Active Ingredient/Active Moiety | | | | |
|--|----------------------------|-----------------|--|--|
| Ingredient Name | Basis of Strength | Strength | | |
| DEXAMETHASO NE SO DIUM PHO SPHATE (UNII: AI9376 Y64P) (DEXAMETHASONE - UNII:7S5I7G3JQL) | DEXAMETHASONE PHOSPHATE | 1 mg in 1 mL | | |
| MOXIFLOXACIN HYDRO CHLORIDE 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:70261-512- 01 | 1 mL in 1 VIAL, SINGLE-USE; Type 0: Not a Combination Product | 0 1/0 1/20 18 | | |
| | | | | |

| Marketing Information | | | | | |
|-----------------------|--|----------------------|--------------------|--|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | | |
| unapproved drug other | | 0 1/0 1/20 18 | | | |
| | | | | | |

Labeler - Imprimis Rx NJ (931390178)

Revised: 5/2018 ImprimisRx NJ