STRENZA PLUS- moxifloxacin 0.5%, bromfenac 0.09%, prednisolone acetate 1%
Brisk Pharmaceuticals

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.

STRENZA PLUS

Rx Only

NDC 73614-732-03

For Use in Eyes Only

STRENZA Plus ™

Each Pack Contains:

Moxifloxacin Ophthalmic Solution, USP 0.5% - One 3 ml Bottle

Bromfenac Ophthalmic Solution 0.09% - One 1.7 ml Bottle

Prednisolone Acetate Ophthalmic Suspension, USP 1% - One 5 ml Bottle

Brisk Pharmaceuticals

Packaged by:

Unit Dose Solutions Inc.,

Morrisville, NC 27560

Distributed by:

Brisk pharmaceuticals,

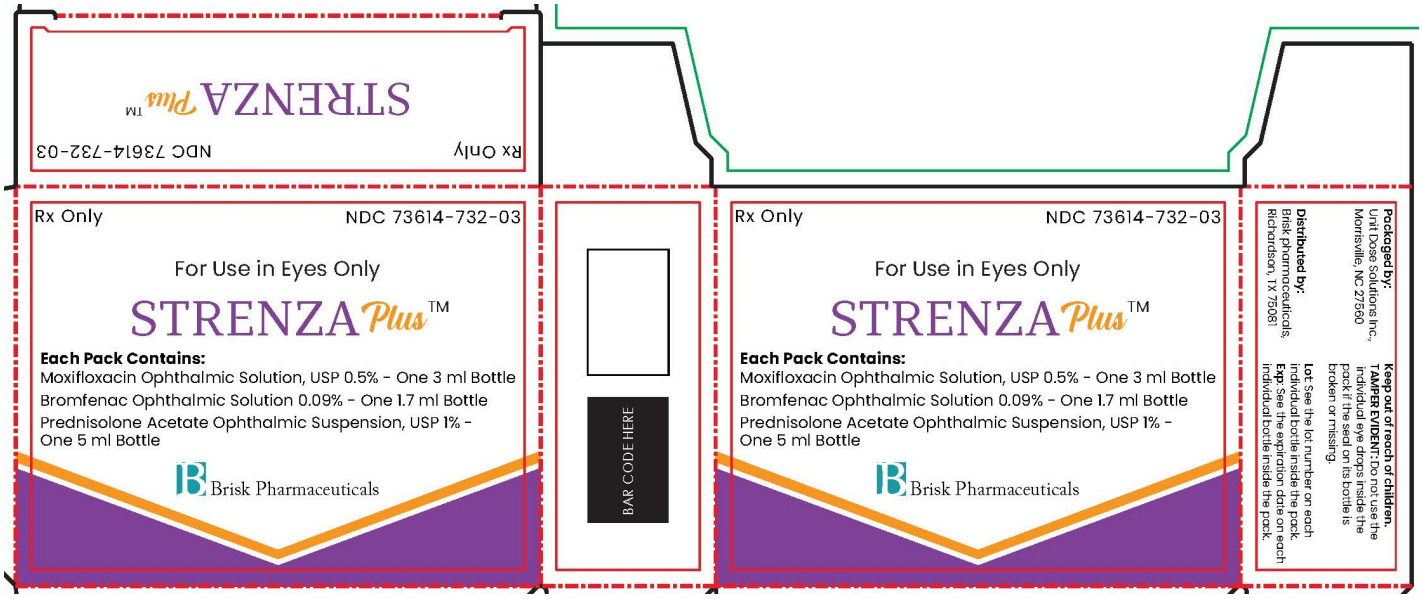
Richardson, TX 75081

Keep out of reach of children.

TAMPER EVIDENT: Do not use the individual eye drops inside the pack if the seal on its bottle is broken or missing.

Lot: See the lot number on each individual bottle inside the pack.

Exp: See the expiration date on each individual bottle inside the pack.



STRENZA PLUS

moxifloxacin 0.5%, bromfenac 0.09%, prednisolone acetate 1% kit

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:73614-732

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73614-732-03	1 in 1 CARTON; Type 1: Convenience Kit of Co-Package	02/10/2025	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	3 mL
Part 2	1 BOTTLE, PLASTIC	5 mL
Part 3	1 BOTTLE, DROPPER	1.7 mL

Part 1 of 3

MOXIFLOXACIN

moxifloxacin solution/ drops

Product Information	
Item Code (Source)	NDC:68180-422
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

Inactive Ingredients	
Ingredient Name	**Strength**
BORIC ACID (UNII: R57ZHV85D4)	
WATER (UNII: 059QF0KO0R)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Product Characteristics			
Color	yellow (Yellow Colored Transparent)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1		3 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA202867	07/01/2017	

Part 2 of 3

PREDNISOLONE ACETATE

prednisolone acetate suspension/ drops

Product Information	
Item Code (Source)	NDC:61314-637
Route of Administration	OPHTHALMIC

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PREDNISOLONE ACETATE (UNII: 8B2807733D) (PREDNISOLONE - UNII:9PHQ9Y1OLM)	PREDNISOLONE ACETATE	10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1		5 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA authorized generic	NDA017469	12/15/1994	

Part 3 of 3

BROMFENAC

bromfenac solution/ drops

Product Information	
Item Code (Source)	NDC:72266-142
Route of Administration	OPHTHALMIC

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BROMFENAC SODIUM (UNII: 8ECV571Y37) (BROMFENAC - UNII:864P0921DW)	BROMFENAC	0.9 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM SULFITE (UNII: VTK01UQK3G)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	0.05 mg in 1 mL
BORIC ACID (UNII: R57ZHV85D4)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72266-142-01	1 in 1 CARTON		
1		1.7 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211029	07/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		02/10/2025	

Labeler - Brisk Pharmaceuticals (117250794)

Revised: 2/2025 Brisk Pharmaceuticals