GOODSENSE ARTIFICIAL TEARS- polyvinyl alcohol solution
Proficient Rx LP

Drug Facts

Uses

- temporary relieves burning and irritations due to dryness of the eye(s)
- protects against further irritation

Warnings

For external use only. Do not use if solution changes color or becomes cloudy or if you are sensitive to any ingredient in this product.

When using this product

- remove contact lenses before using
- do not touch tip of container to any surface to avoid contamination
- replace cap after each use

Stop use and ask a doctor if

- you feel eye pain
- changes in vision occur
- redness or irritation of the eye gets worse or lasts more than 72 hours

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of the reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Instill 1 or 2 drops in the affected eye(s) as needed.
- Store at 15-25C (59-77F).
- Children under 6 years of age: Ask a doctor

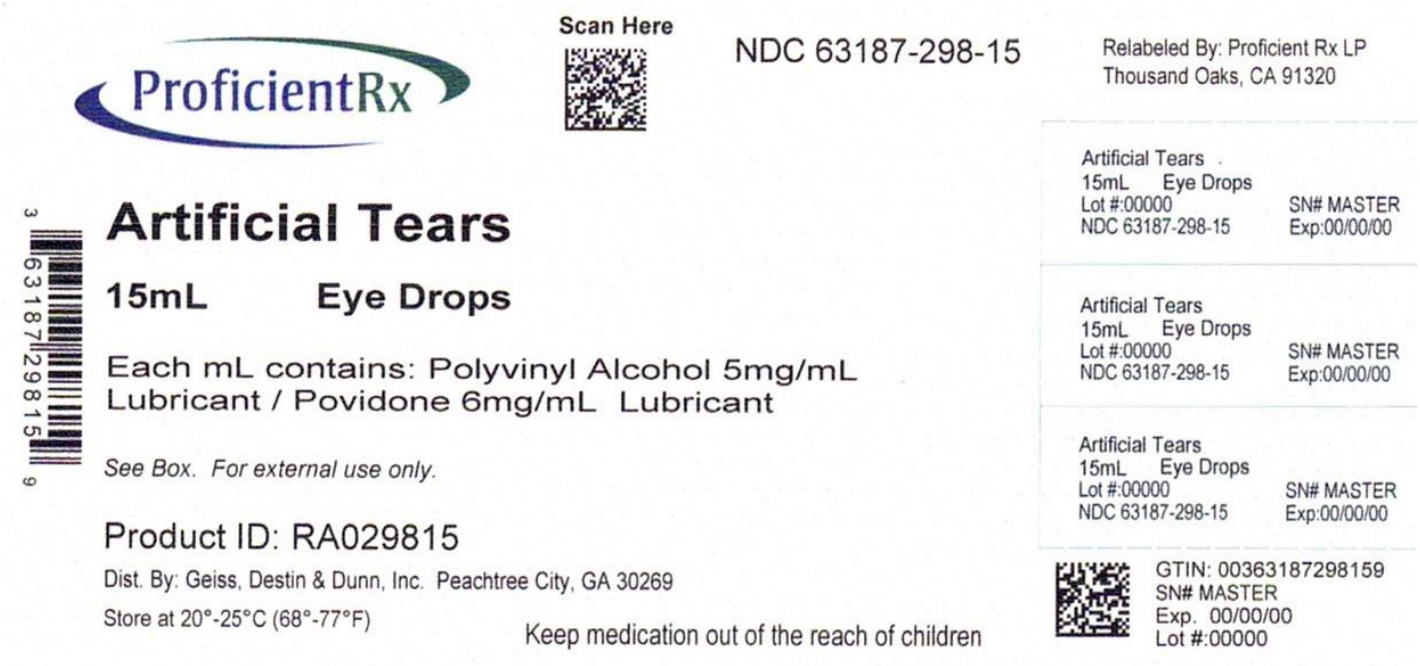
Dosage and Administration

- Instill 1 or 2 drops in the affected eye(s) as needed.

Active Ingredient

Each mL contains: Polyvinyl Alcohol 5mg/ml and Povidone 6mg/ml - lubricant

Inactive Ingredients: Benzalkonium Chloride, Dextrose, Disodium Edetate, Potassium Chloride, Purified Water, Sodium Bicarbonate, Sodium Chloride, Sodium Citrate, Sodium Phosphate (Mono- and Dibasic)



Relabeled By:

Proficient Rx LP

Thousand Oaks, CA 91320

GOODSENSE ARTIFICIAL TEARS

polyvinyl alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63187-298(NDC:11716-0001)
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990) (POLYVINYL ALCOHOL - UNII:532B59J990)	POLYVINYL ALCOHOL, UNSPECIFIED	5 mg in 1 mL
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E)	POVIDONE, UNSPECIFIED	6 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
DEXTROSE, UNSPECIFIED FORM (UNII: IY9XDZ35W2)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
WATER (UNII: 059QF0KO0R)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	

SODIUM PHOSPHATE (UNII: SE337SVY37)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63187-298-15	1 in 1 CARTON	01/01/2019	
1		15 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
OTC Monograph Drug	M018	07/22/2010	

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(63187-298) , RELABEL(63187-298)

Revised: 2/2024 Proficient Rx LP