

**XPECT INDUSTRIAL EYE RELIEF- glycerin, naphazoline hcl solution/ drops**  
**Cintas Corporation**

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**Xpect Industrial Eye Relief 15mL (PLD)**

**Active ingredients**

Glycerin 0.25%

Naphazoline HCl 0.012%

**Purposes**

Lubricant

Redness reliever

**Uses**

- relieves redness of the eye due to minor eye irritations
- for use as a protectant against further irritation or to relieve dryness of the eye
- for the temporary relief of burning and irritation due to dryness of the eye

**Warnings**

For external use only

**Do not use**

if solution changes color or becomes cloudy

**Ask a doctor before use if you have**

narrow angle glaucoma

**When using this product**

- to avoid contamination, do not touch tip of container to any surface
- replace cap after using
- overuse may produce increased redness of the eye
- pupils may become enlarged temporarily

**Stop use and ask a doctor if**

- you experience eye pain
- you experience changes in vision
- you experience continued redness or irritation of the eye
- the condition worsens or persists for more than 72 hours

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

**Directions**

instill 1 to 2 drops in the affected eye(s) up to 4 times daily

**Other information**

store at room temperature

**Inactive ingredients**

benzalkonium chloride, boric acid, edetate disodium, purified water, sodium borate

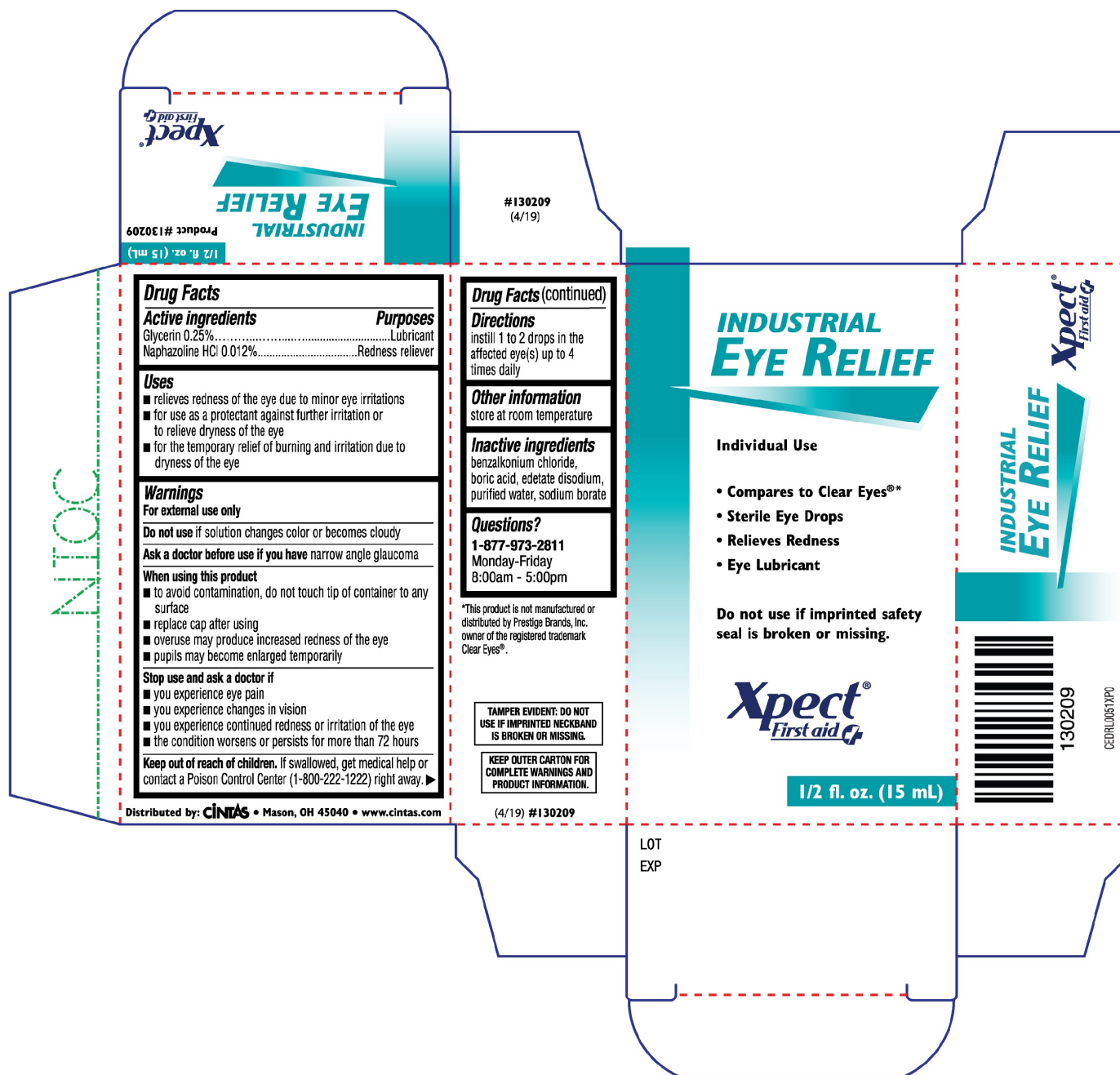
**Questions?**

1-877-973-2811

Monday-Friday

8:00am-5:00pm

**Xpect Industrial Eye Relief 15mL**



## XPECT INDUSTRIAL EYE RELIEF

glycerin, naphazoline hcl solution/ drops

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:42961-255
Route of Administration	OPHTHALMIC		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII: H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.012 g in 100 mL

GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)			GLYCERIN	0.25 g in 100 mL
Inactive Ingredients				
Ingredient Name				Strength
BORIC ACID (UNII: R57ZHV85D4)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
SODIUM BORATE (UNII: 91MBZ8H3QO)				
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)				
WATER (UNII: 059QF0KO0R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:42961-255-01	1 in 1 BOX	09/18/2022	
1		15 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
OTC Monograph Drug		M018	03/01/2022	

**Labeler** - Cintas Corporation (056481716)

**Registrant** - KC Pharmaceuticals, Inc. (174450460)

## Establishment

Name	Address	ID/FEI	Business Operations
KC Pharmaceuticals, Inc.		174450460	manufacture(42961-255) , pack(42961-255) , label(42961-255)

Revised: 12/2023

Cintas Corporation