REDNESS AND DRY EYE RELIEF- glycerin, naphazoline hydrochloride solution
KC Pharmaceuticals, Inc

KC (MFG) for Walmart (PLD) NDC: 79903-362-01
Equate Redness & Dry Eye Relief

Active ingredients
Glycerin 0.25%
Naphazoline HCl 0.012%

Active ingredients Purposes
Glycerin 0.25%.....Lubricant
Naphazoline HCl 0.012%.....Redness reliever

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

Inactive Ingredients

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

Directions:
instill 1 to 2 drops Ianthe affected eye(s) up to 4 times daily

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

REDNESS AND DRY EYE RELIEF

glycerin, naphazoline hydrochloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55651-620
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.25 g in 100 mL

NAPHAZOLINE HYDROCHLORIDE (UNII: MZ1131787D) (NAPHAZOLINE - UNII:H231GF11BV)	NAPHAZOLINE HYDROCHLORIDE	0.012 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
WATER (UNII: 059QF0KO0R)	
BORIC ACID (UNII: R57ZHV85D4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55651-620-01	1 in 1 BOX	12/26/2025	
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	12/26/2025	

Labeler - KC Pharmaceuticals, Inc (174450460)

Revised: 12/2025 KC Pharmaceuticals, Inc