LACRIFRESH HYDRAPF- glycerin, povidone solution/ drops AVIZOR, S.A.

LACRIFRESH HYDRAPF

ACTIVE INGREDIENTS	PURPOSES
Glycerin 0.63%	Lubricant
Povidone 0.25%	Lubricant

- For the temporary relief of burning and irritation due to dryness of the eye
- For the temporary relief of discomfort due to minor irritations of the eye or to exposure to wind or sun
- For use as a protectant against further irritation or to relief dryness of the eye

For external use only

Do not use

- If solution changes color or becomes cloudy
- If you are sensitive to any ingredient in this product

When using this product

- Do not touch tip of container to any surface to avoid contamination
- Replace cap after using

Stop use and ask a doctor if

- You experience eye pain, changes in vision, continued redness or irritation of the eye
- 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

Directions

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

Other information

Other information

- Store at 15º-30ºC (59º-86ºF)
- Discard the product within 90 days of first opening

Inactive ingredients

boric acid, calcium chloride, magnesium chloride, potassium chloride, purified water, sodium borate, sodium chloride, sodium hyaluronate

Questions or comments?

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Call 1-800-714-1824

AVIZOR

Lacrifresh

LUBRICANT EYE DROPS

PRESERVATIVE FREE

HydraPF

Dry Eye Relief

Instant Comfort and Hydration

Protects and Lubricates the Eye Surface

Enhanced with

0.63% Glycerin + 0.25% Povidone

Sterile

0.33 FL OZ (10 mL)



LACRIFRESH HYDRAPF

glycerin, povidone solution/ drops				
Product Information				
Product Type	HUMAN OTC DRUG	Item Co	de (Source)	NDC:82808-004
Route of Administration	OPHTHALMIC			
Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
POVIDONE (UNII: FZ989GH94E) (POVIDONE - UNII:FZ989GH94E) POVIDONE 0.25 g in 100 r			0.25 g in 100 mL	

GLYCERIN

Inactive Ingredients			
Ingredient Name	Strength		
SODIUM BORATE (UNII: 91MBZ8H3QO)			
CALCIUM CHLORIDE (UNII: M4I0D6VV5M)			
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)			
BORIC ACID (UNII: R57ZHV85D4)			
SODIUM HYALURONATE (UNII: YSE9PPT4TH)			
POTASSIUM CHLORIDE (UNII: 660YQ98I10)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			

Packaging				
-	tem Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:82808- 004-01	1 in 1 CASE	11/03/2025	
	L	10 mL in 1 BOTTLE, DROPPER; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M018	11/03/2025	

Labeler - AVIZOR, S.A. (465588731)

Registrant - AVIZOR, S.A. (465588731)

Establishment			
Name	Address	ID/FEI	Business Operations
AVIZOR, S.A.		465588731	manufacture(82808-004), label(82808-004), pack(82808-004)

Revised: 10/2025 AVIZOR, S.A.