

REGENER-EYES OPHTHALMIC SOLUTION PROFESSIONAL- glycerin ophthalmic solution solution/ drops
M & M PRODUCTS, LLC

Active Ingredient

Active Ingredient Glycerin 0.5%

Purpose

Purpose: Eye Lubricant

Indications & Usage

Uses:

For the temporary relief of burning and irritation due to dryness of the eye. For use as a lubricant to prevent further irritation or to relieve dryness of the eye. For use as a protectant against further or to relieve dryness of the eye.

Indications:

Regener-Eyes® is formulated to provide temporary relief of burning and irritation due to dryness of the eye. Each product is preservative-free and can be used as a protectant against further irritation or to relieve dryness of the eye.

Warnings

Warnings:

External Use Only. To avoid contamination, do not touch tip of container to any surface. Replace cap after using. If solution changes color or becomes cloudy, do not use.

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 immediately

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Dosage and Administration

Directions:

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

When Using

Store at Room Temperature.

Inactive Ingredient

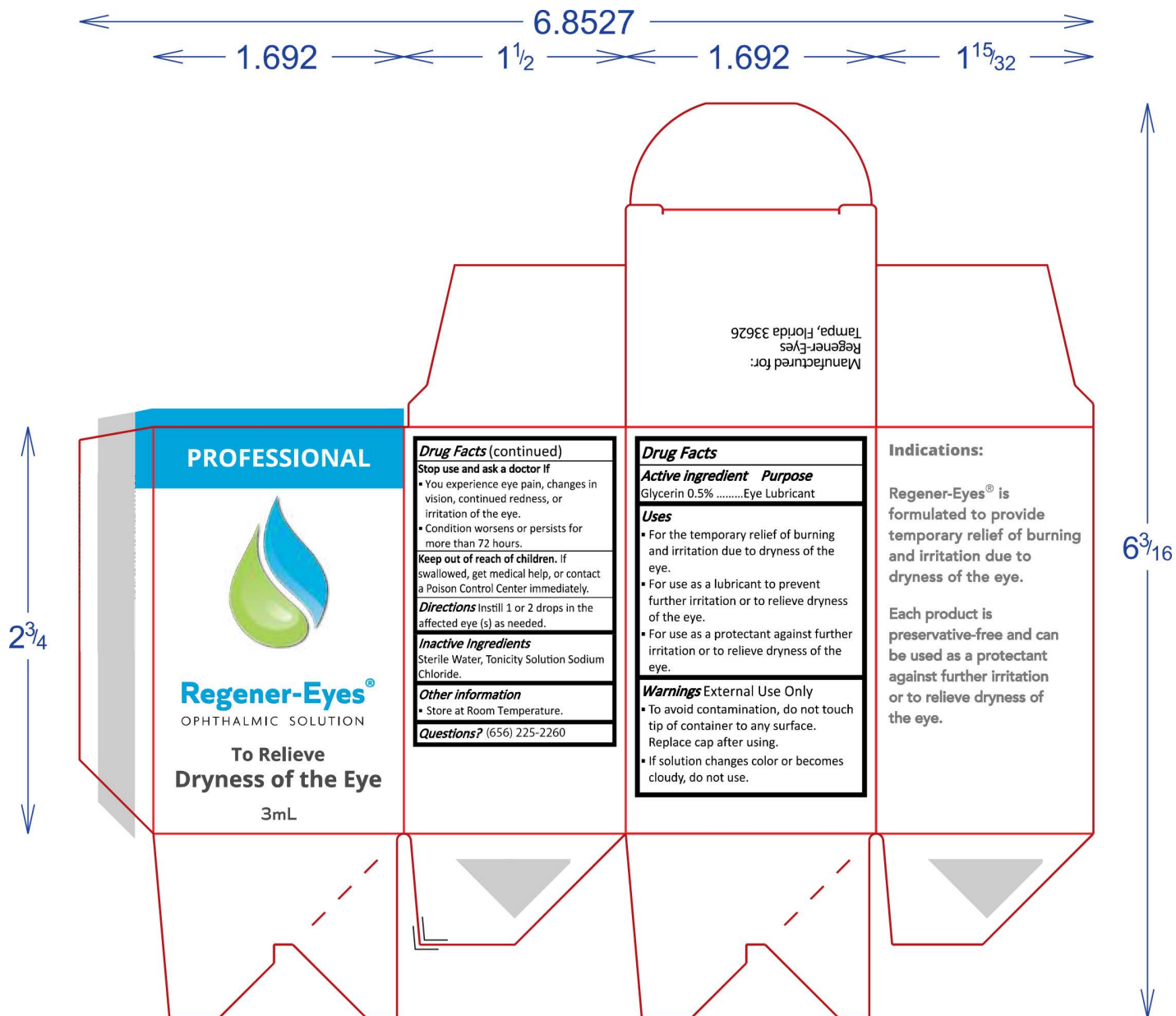
Inactive Ingredients:

Sterile Water, Tonicity Solution Sodium Chloride.

Patient Counseling

Questions?

(656) 225-2260



REGENER-EYES OPHTHALMIC SOLUTION PROFESSIONAL

glycerin ophthalmic solution solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87354-004
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87354-004-01	1 in 1 CARTON	02/01/2026	
1		1 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	02/01/2026	

Labeler - M & M PRODUCTS, LLC (959745410)

Establishment			
Name	Address	ID/FEI	Business Operations
M & M PRODUCTS, LLC		959745410	manufacture(87354-004)