

**LUBRICANT EYE DROPS SOLUTION- polyethylene glycol 400 and propylene glycol solution**  
**AvPAK**

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**Lubricant Eye Drops Solution**

<b><i>Active Ingredients</i></b>	<b><i>Purpose</i></b>
Polyethylene Glycol 400 0.4%	Lubricant
Propylene Glycol 0.3%	Lubricant

***Uses***

- For the temporary relief of burning, irritation and discomfort due to dryness of the eye and exposure to wind or sun.
- May be used as a protectant against further irritation.

***Warnings***

**For use in the eyes only**

**Do not use**

- If solution changes color or becomes cloudy.

**When using this product**

- avoid contamination, do not touch tip of container to any surface
- replace cap after use

**Stop use and ask a doctor if**

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

**Keep this and all drugs out of the reach of children.**

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

***Directions***

- shake well before use
- instill 1 or 2 drops in the affected eye(s) as needed

***Other Information***

- store at room temperature 15°- 30°C (59°-86°F).
- do not use if imprinted seal on cap is torn, broken or missing

- discard 90 days after opening
- retain outer carton for full product information

**Inactive Ingredients:**

Benzalkonium chloride (preservative), boric acid, calcium chloride, hypromellose, magnesium chloride, potassium chloride, purified water, sodium chloride, zinc chloride. May contain hydrochloric acid and/or sodium hydroxide to adjust pH.

**Questions?**

1-855-361-3993

Distributed by:

AvKARE

Pulaski, TN 38478

www.avkare.com

Rev. 12/2022 AV 12/2022

**PRINCIPAL DISPLAY PANEL**

**AVKARE**  
NDC 50268-126-15  
**Lubricant Eye Drops Solution**  
Polyethylene Glycol 400 0.4% /  
Propylene Glycol 0.3%  
(Eye Lubricants)  
**Lubricant Eye Drops**  
**STERILE**  
0.5 FL OZ (15 mL)

**DO NOT USE IF TAMPER EVIDENT SEAL IS TORN, BROKEN OR MISSING**

**Drug Facts**  
**Active ingredient**—**Purpose**  
Polyethylene Glycol 400 0.4%.....Eye Lubricant  
Propylene Glycol 0.3%.....Eye Lubricant  
**Use:** • For the temporary relief of burning, irritation and discomfort due to dryness of the eye or exposure to wind or sun. May be used as a protectant against further irritation.  
**Warnings:** For use in the eyes only  
**Keep this and all drugs out of the reach of children.**  
**Directions:** Instill 1 or 2 drops in the affected eye(s) as needed.  
**Other information: DO NOT USE IF IMPRINTED SEAL ON CAP IS TORN, BROKEN OR MISSING.**  
• Retain outer carton for full product information.  
**Inactive ingredients:** See outer carton.  
**Questions? 1-855-361-3993**

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Rev. 12/2022 AV 12/2022

<b>LUBRICANT EYE DROPS SOLUTION</b>			
polyethylene glycol 400 and propylene glycol solution			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50268-126
<b>Route of Administration</b>	OPHTHALMIC		
<b>Active Ingredient/Active Moiety</b>			

Ingredient Name	Basis of Strength	Strength
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ) (POLYETHYLENE GLYCOL 400 - UNII:B697894SGQ)	POLYETHYLENE GLYCOL 400	4 mg in 1 mL
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3) (PROPYLENE GLYCOL - UNII:6DC9Q167V3)	PROPYLENE GLYCOL	3 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>BORIC ACID</b> (UNII: R57ZHV85D4)	
<b>CALCIUM CHLORIDE</b> (UNII: M4I0D6VV5M)	
<b>MAGNESIUM CHLORIDE</b> (UNII: 02F3473H9O)	
<b>POTASSIUM CHLORIDE</b> (UNII: 660YQ98I10)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>ZINC CHLORIDE</b> (UNII: 86Q357L16B)	
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50268-126-15	1 in 1 CARTON	02/25/2003	
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	02/25/2003	

**Labeler - AvPAK (832926666)**

Revised: 1/2026

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