

**0.25 % PHENYLEPHRINE HCL HEMORRHOIDAL- mineral oil, petrolatum, phenylephrine hydrochloride ointment**  
**Universal Distribution Center LLC**

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**0.25 % PHENYLEPHRINE HCl HEMORRHOIDAL OINTMENT**

***Drug Facts***

***Active ingredient***

Mineral Oil 14%  
Petrolatum 74.9%  
Phenylephrine HCl 0.25%

***Purpose***

Protectant  
Protectant  
Vasoconstrictor

***Uses***

helps relieve the local itching and discomfort associated with hemorrhoids • temporarily shrinks hemorrhoidal tissue and relieves burning • temporarily provides a coating for relief of anorectal discomforts • temporarily protects the inflamed, irritated anorectal surface to help make bowel movements less painful

***Warnings***

**For external use only.**

**Ask a doctor before use if you have** • heart disease • high blood pressure • thyroid disease • diabetes • difficulty in urination due to enlargement of the prostate gland

**Ask a doctor or pharmacist before use if you are** presently taking a prescription drug for high blood pressure or depression

**When using this product** do not exceed the recommended daily dosage unless directed by a doctor

**Stop use and ask a doctor if** • bleeding occurs • condition worsens or does not improve within 7 days • introduction of applicator into the rectum causes additional pain

**If pregnant or breast-feeding,** ask a healthcare professional before use

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

## Directions

• adults: when practical, cleanse the affected area by patting or blotting with an appropriate cleansing wipe. Gently dry by patting or blotting with a tissue or a soft cloth before applying ointment • when first opening the tube, puncture foil seal secured on tube • apply to the affected area up to 4 times daily, especially at night, in the morning or after each bowel movement • intrarectal use: remove cover from applicator, attach applicator to tube, lubricate applicator well and gently insert applicator into the rectum • thoroughly cleanse applicator after each use and replace cover • also apply ointment to external area • regular use provides continual therapy for relief of symptoms • children under 12 years of age: ask a doctor.

## Other information

• store at 20°–25°C (68°–77°F) • do not use if tube seal under cap is broken

## Inactive ingredients

beeswax, ceteareth-20, cetostearyl alcohol, glycerin

## FAST RELIEF OINTMENT

- ✓ Relieves burning
- ✓ Soothes pain
- ✓ itching & swelling

### Mfd for and Distributed by:

Universal Distribution Center LLC  
330 Applegarth Road,  
Monroe Township, NJ 08831  
[www.universaldc.com](http://www.universaldc.com)

MADE IN CHINA

## Packaging

### Outer Package Label



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Petrolatum 74.9%	Protectant
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ITEM#: 56088



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**0.25%  
PHENYLEPHRINE  
HCl**

**HEMORRHOIDAL  
OINTMENT**

Mineral Oil  
Petrolatum



- Relieves burning
- Soothes pain
- itching & swelling.

NET WT 2oz (57g)



**0.25%  
PHENYLEPHRINE HCl**

**HEMORRHOIDAL  
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HEMORRHOIDAL  
OINTMENT  
0.25% PHENYLEPHRINE HCl



## Inner Package Label





**0.25%  
PHENYLEPHRINE HCl**

**HEMORRHOIDAL  
OINTMENT**

Mineral Oil  
Petrolatum



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- ✓ Soothes pain
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mineral oil, petrolatum, phenylephrine hydrochloride ointment

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:52000-438
<b>Route of Administration</b>	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
<b>MINERAL OIL</b> (UNII: T5L8T28FGP) (MINERAL OIL - UNII:T5L8T28FGP)	MINERAL OIL	14 g in 100 g
<b>PETROLATUM</b> (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	74.9 g in 100 g
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	0.25 g in 100 g

Inactive Ingredients	
Ingredient Name	Strength
<b>YELLOW WAX</b> (UNII: 2ZA36H0S2V)	
<b>POLYOXYL 20 CETOSTEARYL ETHER</b> (UNII: YRC528SWUY)	
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-438-02	1 in 1 BOX	01/19/2026	
1		57 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	01/19/2026	

**Labeler** - Universal Distribution Center LLC (019180459)