# WALGREENS- hemorrhoidal pain relief ointment Walgreens

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# Walgreens ® Hemorrhoidal Pain Relief Ointment

#### **ACTIVE INGREDIENTS**

Mineral oil 14%

Petrolatum 74.9%

Phenylephrine HCI 0.25%

### **PURPOSES**

**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 and/or intrarectal 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 health 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 with top end of cap
- 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)

#### INACTIVE INGREDIENTS

benzoic acid, butylated hydroxyanisole, corn oil, glycerin, lanolin, lanolin alcohols, methylparaben, mineral oil, paraffin, propylparaben, purified water, thymus vulgaris (thyme) flower/leaf oil, tocopherols excipient, white wax

## **QUESTIONS OR COMMENTS?**

1-800-925-4733

## 28 g Carton Label





# 57 g Carton Label

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"GLUE AREA"



# **WALGREENS**

hemorrhoidal pain relief ointment

<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-6503
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	2.5 mg in 1 g		
MINERAL OIL (UNII: T5L8T28FGP) (MINERAL OIL - UNII:T5L8T28FGP)	MINERAL OIL	140 mg in 1 g		
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	749 mg in 1 g		

Inactive Ingredients			
Ingredient Name	Strength		
BENZOIC ACID (UNII: 85KN0B0MIM)			
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)			
CORN OIL (UNII: 8470G57WFM)			
GLYCERIN (UNII: PDC6A3C0OX)			
LANOLIN (UNII: 7EV65EAW6H)			
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)			
METHYLPARABEN (UNII: A2I8C7HI9T)			
PARAFFIN (UNII: 1900E3H2ZE)			
PROPYLPARABEN (UNII: Z8IX2SC1OH)			
WATER (UNII: 059QF0KO0R)			
THYME (UNII: CW6570BU4N)			
WHITE WAX (UNII: 7G1J5DA97F)			
TOCOPHEROL (UNII: ROZB2556P8)			

Color yellow (smooth yellow ointment)  Shape Flavor  Size Imprint Code	Product Characteristics		
Flavor Imprint Code	Color	yellow (smooth yellow ointment)	Score
	Shape		Size
	Flavor		Imprint Code
Contains	Contains		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-6503- 01	1 in 1 CARTON	03/01/2004	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:0363-6503- 02	1 in 1 CARTON	03/01/2004	
2		57 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing In	formation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	03/01/2004	

# Labeler - Walgreens (008965063)

# Registrant - Unipack LLC (116015769)

Establishmen	t		
Name	Address	ID/FEI	Business Operations
Unipack LLC		009248480	manufacture(0363-6503)

Revised: 12/2024 Walgreens