RAPIDOL HEMORRHOIDAL- glycerin, phenylephrine hcl, pramoxine hcl, white petrolatum cream
Pharmadel LLC

Rapidol Hemorrhoidal Cream (K)

Drug Facts

Active ingredients & Purposes

Active ingredients	*Purposes*
Glycerin 14.4%.....	Protectant
Phenylephrine HCl 0.25%.....	Vasoconstrictor
Pramoxine HCl 1 %.....	Local anesthetic
White petrolatum 15%.....	Protectant

Uses

- for the temporary relief of local itching and discomfort associated with
- hemorrhoid
- anorectal inflammation
- temporarily shrinks hemorrhoidal tissue
- temporarily relieves the symptoms of perianal skin irritation

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 of 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
- do not put this product into the rectum by using fingers or any mechanical device or applicator

Stop use and consult a doctor if

- condition worsens or does not improve within 7 days
- bleeding occurs

If pregnant or breastfeeding,

ask a doctor before use.

KEEP OUT OF THE REACH OF CHILDREN.

In case of accidental ingestion, seek medical help or contact a Poison Control Center immediately.

Directions

- **adults:** when practical, cleanse the affected area with soap and warm water and rinse thoroughly. Gently dry by patting or blotting with toilet tissue or a soft cloth before application of this product

apply externally to the affected area after a bowel movement, no more than 4 times daily

- **children under 12 years of age**: consult a doctor

Other information

- store between 60-77°F (20-25°C)
- do not use if foil seal under cap is broken or missing

Inactive Ingredients

aloe vera leaf, alpha tocopherol (Vit. E), butylated hydroxyanisole, carboxymethylcellulose sodium, cetostearyl alcohol, cetyl alcohol, citric acid monohydrate, edetate disodium, glyceryl monostearate, methylparaben, mineral oil, panthenol, propylparaben, sodium benzoate, steareth-20, xanthan gum, water

Questions?

+1-866-359-3478 (M-F) 9AM to 5 PM EST or www.pharmadel.com

Distributed by/ Distribuido por:

Pharmadel LLC

New Castle, DE 19720

Made in India/ Hecho en India

Product Labeling

RAPIDOL HEMORRHOIDAL

glycerin, phenylephrine hcl, pramoxine hcl, white petrolatum cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55758-448
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	14.4 g in 100 g
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	0.25 g in 100 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	1 g in 100 g
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	15 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PANTHENOL (UNII: WV9CM0O67Z)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
STEARETH-20 (UNII: L0Q8IK9E08)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
MINERAL OIL (UNII: T5L8T28FGP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55758-448-01	1 in 1 CARTON	08/30/2024	
1		30 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	08/30/2024	

Labeler - Pharmadel LLC (030129680)

Revised: 8/2024 Pharmadel LLC