# PREFERRED PLUS HEMORRHOID- glycerin, petrolatum, phenylephrine hydrochloride, pramoxine hydrochloride cream Kinray, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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#### **Drug Facts**

# **Active ingredients**

Glycerin 14.4%

Petrolatum 15%

Phenylephrine HCl 0.25%

Pramoxine HCl1 %

## **Purposes**

**Protectant** 

**Protectant** 

Vasoconstrictor

Local anesthetic

#### Uses

- helps relieve the local itching and discomfort associated with hemorrhoids
- temporarily shrinks hemorrhoidal tissue
- temporarily relieves burning
- aids in protecting irritated anorectal areas

#### **Warnings**

#### For external use only

# Ask a doctor before use if you have

- heart disease
- high blood pressure
- diabetes
- thyroid disease
- trouble urinating due to an enlarged 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
- do not put this product into the rectum by using fingers or any mechanical device or applicator

#### Stop use and ask a doctor if

bleeding occur

• condition gets worse or does not Improve within 7 days

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 and gently dry the affected area by patting or blotting with an appropriate cleansing tissue or soft cloth before applying gel
- when first opening the tube, puncture foil seal with top end of cap
- apply externally to the affected area up to 4 times daily, especially at night, in the morning or after each bowel movement
- Children under 12 years of age: ask a doctor

#### Other information

- store at 20°-25°C (68°-77°F)
- for lot number and expiration date, see crimp of tube or see box

#### **Inactive ingredients**

• cetearyl alcohol, cetyl alcohol, monosterate, mineral oil, peg 40 castor oil, propylene glycol, purified water, stearyl alcohol, tocopheryl acetate (Vitamin E)

## **Principal Display Panel - 1 oz Carton Label**

TAMPER-EVIDENT: Do not use if printed "safety sealed" overwrap is broken or missing.

NDC 61715-092-01

Preferred Plus Pharmacy®

**HEMORRHOIDAL** 

**CREAM** *Maximum Strength Pain Relief* 

\*Compare to the active ingredients in PREPARATION H® Cooling Gel

#### **Smooth Cream Formula**

- Rapid Soothing Pain Relief from Painful Burning, Itching and Discomfort
- Shrinks Swollen Hemorrhoidal Tissue
- Protects Irritated Tissue
- Relieves External Discomfort

#### SATISFACTION GUARANTEED

**Preferred Plus Pharmacy**®

**NET WT 1 OZ (28 g)** 



# PREFERRED PLUS HEMORRHOID

glycerin, petrolatum, phenylephrine hydrochloride, pramoxine hydrochloride cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:61715-092
Route of Administration	RECTAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Glycerin (UNII: PDC6A3C0OX) (Glycerin - UNII:PDC6A3C0OX)	Glycerin	144 mg in 1 L	
Petrolatum (UNII: 4T6H12BN9U) (Petrolatum - UNII:4T6H12BN9U)	Petro latum	150 mg in 1 L	
Phenylephrine Hydrochloride (UNII: 04JA59TNSJ) (Phenylephrine - UNII:1WS297W6MV)	Phenylephrine Hydrochloride	2.5 mg in 1 L	
Pramoxine Hydrochloride (UNII: 88AYB867L5) (Pramoxine - UNII:068X84E056)	Pramoxine Hydrochloride	10 mg in 1 L	

Inactive Ingredients		
Ingredient Name	Strength	
Aloe Vera Leaf (UNII: ZY81Z83H0X)		
Cetostearyl Alcohol (UNII: 2DMT128M1S)		
Cetyl Alcohol (UNII: 936JST6JCN)		
Glyceryl Monostearate (UNII: 230 O U9 XXE4)		
Mineral Oil (UNII: T5L8T28FGP)		

Peg-40 Castor Oil (UNII: 4ERD2076EF)	
Water (UNII: 059QF0KO0R)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Stearyl Alcohol (UNII: 2KR89I4H1Y)	
.ALPHATocopherol Acetate (UNII: 9E8X80D2L0)	

P	Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date	
1	NDC:61715-092-01	1 in 1 CARTON	03/28/2014		
1		0.028 L 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 final	part346	03/28/2014	

# **Labeler -** Kinray, Inc. (012574513)

Establishment				
Name	Address	ID/FEI	Business Operations	
PJ Noyes Company		018927392	MANUFACTURE(61715-092)	

Revised: 1/2018 Kinray, Inc.