

HEB MAXIMUM STRENGTH PAIN RELIEF HEMORRHOIDAL- glycerin, petrolatum, phenylephrine hydrochloride, and pramoxine hydrochloride cream
HEB

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.

HEB Maximum Strength Pain Relief Hemorrhoidal Cream

Drug Facts

<i>Active ingredients</i>	<i>Purposes</i>
Glycerin 14.4%	Protectant
Petrolatum 15%	Protectant
Phenylephrine hydrochloride 0.25%	Vasoconstrictor
Pramoxine hydrochloride 1%	Local Anesthetic

Uses

- temporarily relieves pain, soreness, and burning
- helps relieve local itching and discomfort associated with hemorrhoids
- temporarily shrinks hemorrhoidal tissue
- 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 urinating due to enlargement of the prostate gland

Ask a doctor or pharmacist before use if you are 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 into rectum by using fingers or any mechanical device or applicator

Stop use and ask a physician if

- bleeding occurs
- the condition worsens or does not improve within 7 days
- an allergic reaction develops
- the symptom being treated does not subside or redness, irritation, swelling, pain, or other symptoms develops, persists, or increase

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: 800-222-1222.

Directions

Adults:

- when practical, cleanse affected area by patting or blotting with appropriate cleansing wipe. Gently dry by patting or blotting with tissue or soft cloth before applying cream.
- when first opening tube, remove foil seal
- apply externally or in lower portion of anal canal only
- apply externally to affected area up to 4 times daily, especially at night, in the morning, or after each bowel movement
- for application in lower anal canal; remove cover from dispensing cap. Attach dispensing cap to tube. Lubricate dispensing cap well, then gently insert dispensing cap partway into anus
- thoroughly cleanse dispensing cap after each use, and replace cover
- **Children under 12 years:** ask a doctor

Other information

- store at 15 to 30°C (59 to 86°F).
- close cap tightly after use
- **Tamper Evident:** DO NOT USE IF SEAL ON TUBE IS PUNCTURED OR MISSING.

Inactive ingredients

aloe barbadensis leaf juice, BHA, cetyl alcohol, citric acid, disodium EDTA, glyceryl stearate, laureth-23, methylparaben, mineral oil, panthenol, propyl gallate, propylene glycol, propylparaben, purified water, sodium benzoate, sodium carboxymethylcellulose, steareth-2, steareth-20, stearyl alcohol, tocopherol (vitamin E) acetate, xanthan gum

Questions or comments?

866-323-0107

PRINCIPAL DISPLAY PANEL - 56.7 g Tube Carton

Compare to Preparation H[®] Cream active ingredients*

NDC 37808-992-24

H-E-B[®]

Maximum Strength Pain Relief

Hemorrhoidal Cream

Phenylephrine Hydrochloride / Vasoconstrictor • Pramoxine Hydrochloride / Local Anesthetic

Hemorrhoid Symptom Relief

With Soothing Aloe

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

Includes

Dispensing

Cap

NET WT 2.0 OZ (56.7 g)

Drug Facts (continued)

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Compare to Preparation H® Cream active ingredients*



NDC 37808-992-24

Maximum Strength Pain Relief

Hemorrhoidal Cream

Phenylephrine Hydrochloride / Vasoconstrictor • Pramoxine Hydrochloride / Local Anesthetic

Hemorrhoid Symptom Relief

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Maximum Strength Pain Relief

Hemorrhoidal Cream

Phenylephrine Hydrochloride / Vasoconstrictor
Pramoxine Hydrochloride / Local Anesthetic

Hemorrhoid Symptom Relief

MADE WITH PRIDE
AND CARE FOR
H-E-B®
SAN ANTONIO TX
78204



MADE IN
THE USA



100%
GUARANTEE
promise

If you aren't completely
pleased with this product,
we'll be happy to replace
it or refund your money.
You have our word on it.

8586-1804



Tamper Resistant
do not use if seal on
tube is punctured
or missing.

*This product is not manufactured or distributed by Pfizer, Inc., the distributor of Preparation H®.

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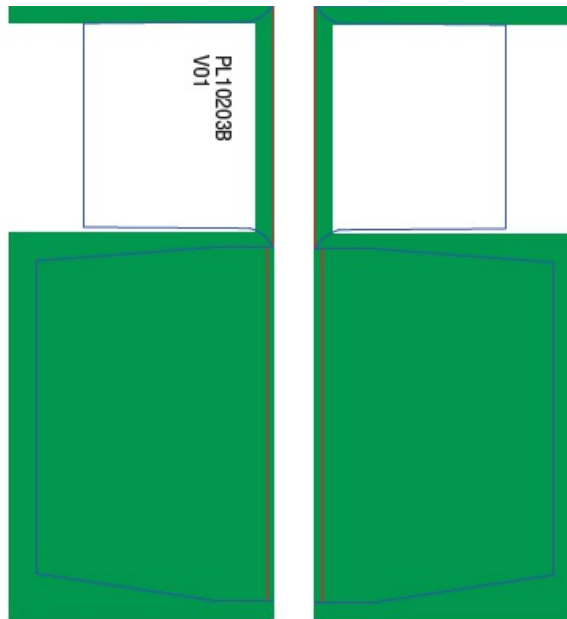
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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-992
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	0.144 mg in 2 g
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	0.15 mg in 2 g
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	0.0025 mg in 2 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	0.1 mg in 2 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
LAURETH-23 (UNII: N72LMW566G)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
PANTHENOL (UNII: WV9CM0O67Z)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
WATER (UNII: 059QF0KO0R)				
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)				
STEARETH-2 (UNII: V56DFE46J5)				
STEARETH-20 (UNII: L0Q8IK9E08)				
STEARYL ALCOHOL (UNII: 2KR89I4HIY)				
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)				
XANTHAN GUM (UNII: TTV12P4NEE)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-992-24	1 in 1 CARTON	10/18/2018	
1		56.7 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 final	part346		10/18/2018	

Labeler - HEB (007924756)

Establishment			
Name	Address	ID/FEI	Business Operations
Natureplex LLC		062808196	MANUFACTURE(37808-992)

Revised: 10/2018

HEB