HEMORRHOID PAIN RELIEF- lidocaine cream Telebrands Corp

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.

Hemorrhoid Pain Relief

Active ingredient

Lidocaine 5%

Purpose

Lidocaine 5%.....Local Anesthetic

Uses

For the temporary relief of pain, burning, itching and discomfort associated with hemorrhoids.

Warnings

For external use only

Allergy alert: If prone to allergic reaction from aspirin or salicylates, consult a doctor before use.

When using this product

- do not exceed the recommended daily dosage unless directed by a doctor.
- use only as directed. Read and follow all directions and warnings on this label.
- do not put this product into the rectum by using finger or any mechanical device or applicator.

Stop use and ask a doctor if

- if condition worsens or does not improve within 7 days
- in case of bleeding
- symptom being treated does not subside or if redness, irritation, swelling, pain, or other symptoms develop or increase. Certain persons can develop allergic reactions to ingredients in this product.

If pregnant or breast-feeding, ask a health professional before use.

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

Directions

- **Adults:** When practical, cleanse the affected area with mild 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 to affected area.
- Apply to the affected area up to 6 times a day
- Children under 12 years of age: consult a doctor.

Other information

Do not use if seal is broken or missing

Inactive ingredients

Water/Aqua/Eau, Ethylhexyl Stearate, Butylene Glycol, Dimethicone, Stearic Acid, Caprylic/Capric Triglyceride, Cannabis Sativa (Hemp) Seed Oil, Glyceryl Stearate, PEG-100 Stearate, Cetearyl Alcohol, Helianthus Annuus (Sunflower) Seed Oil, Curcuma Longa (Turmeric) Root Extract, Allantoin, Glycerin, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Phenoxyethanol, Caprylyl Glycol, Ethylhexylglycerin, Hexylene Glycol, Sodium Hydroxide, Disodium EDTA

Questions?

Call (855) 877-4503 (M-F, 9am-5pm EST)

Hempvana

NDC 73287-028-01

Maximum OTC Strength

Hemorrhoid Pain Relief

Topical Anesthetic Cream

5% Lidocaine

+ Hemp Seed Oil For Moisturization

Net Wt. 3oz (85g)

100% Fragrance Free





HEMORRHOID PAIN RELIEF

lidocaine cream

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC	73287-028
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis o	f Strength	Strength

Inactive Ingredients	
Ingredient Name	Strength
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
TURMERIC (UNII: 856YO1Z64F)	
ALLANTOIN (UNII: 344S277G0Z)	
GLYCERIN (UNII: PDC6A3C0OX)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
PEG-100 STEARATE (UNII: YD01N1999R)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
ETHYLHEXYL STEARATE (UNII: EG3PA2K3K5)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
WATER (UNII: 059QF0KO0R)	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:73287-028- 01	1 in 1 CARTON	08/29/2023		
1		85 g in 1 JAR; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	M015	08/29/2023		

Labeler - Telebrands Corp (177266558)

Establishment				
Name	Address	ID/FEI	Business Operations	
Neutraderm, Inc.		146224444	manufacture(73287-028)	

Revised: 8/2023 Telebrands Corp