BODY ACTION PRODUCTS BUTT EZE BENZOCAINE- benzocaine gel
PRODUCT MAX GROUP INC

BODY ACTION PRODUCTS: Butt Eze Benzocaine Gel

Drug Facts

Active Ingredient

Benzocaine 5%

Purpose

Anorectal (Hemorrhoidal) Gel

Uses

- For temporary relief of pain or soreness in the perianal area.

Warnings

For external use only.

- Avoid contact with the eyes.
- Certain persons can develop allergic reactions from ingredients in this product. If the symptom being treated does not subside or if redness, irritation, swelling, pain or other symptoms develop or increase, discontinue use and consult a doctor.

Keep out of reach of children.

If swallowed get medical help or contact a Poison Control Center right away.

Directions

- When practial, 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.
- Apply to the affected area up to 6 times daily.

Other Information

Do not use if safety seal is broken or missing.

Inactive Ingredients

Cannabis Sativa (Hemp) Seed Oil, Hydroxyethylcellulose, Methylparaben, PEG-8, Polysorbate 20, Propylene Glycol, Propylparaben, Water.

Package Labeling:

BODY ACTION PRODUCTS BUTT EZE BENZOCAINE

benzocaine gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70742-286
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	50 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70742-286-00	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/29/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	04/29/2022	

Labeler - PRODUCT MAX GROUP INC (134893911)

Revised: 11/2023

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