EZ ACCESS- benzocaine liquid
Product Max Group 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.

EZ ACCESS - 001

Drug Facts
Active Ingredient

Benzocaine 5%

Purpose

Topical Analgesic

Keep out of reach of children

- **Keep out of reach of children.** If swallowed get medical help or contact a Poison Control Center right away.

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 a consult a doctor.

Directions

- When practical, cleanse the affected area with mild soap and water and rinse thoroughly.
- Gently dry by patting or blotting with a tissue or a soft cloth before the application of this product.
- Apply to the affected area up to 6 times daily.

Other information

Do not use if safety tab is broken or missing

Inactive Ingredients

Methylparaben, PEG-8, Propylene Glycol, Propylparaben, Water

EZ ACCESS product label

Anal Desensitizer

EZ ACCESS

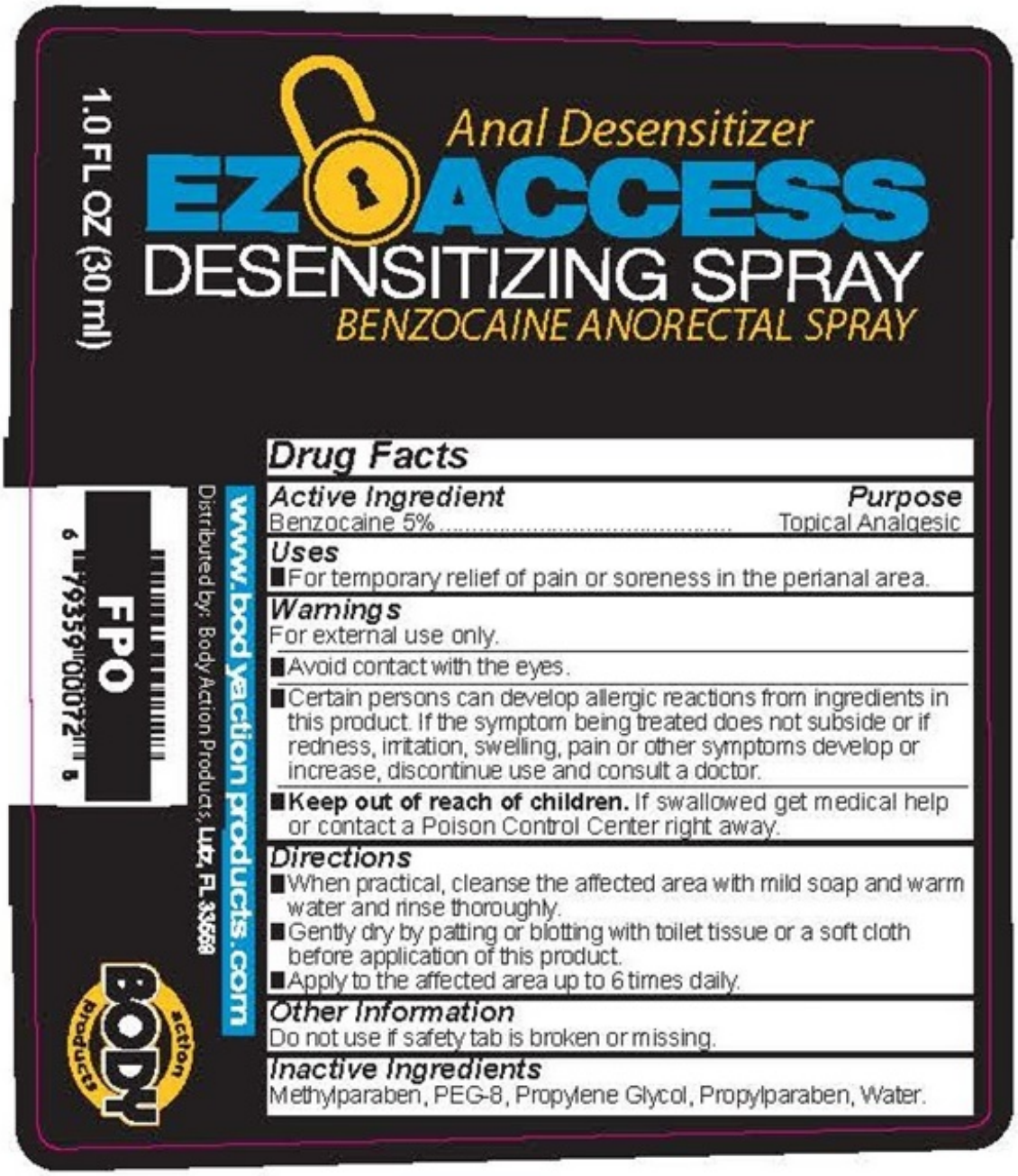
DESENSITIZING SPRAY
BENZOCAINE ANORECTAL SPRAY

1.0 FL OZ (30mL)

www.bodyactionproducts.com

Distributed by: BodyAction Products, Lutz, FL 33559

action BODY products



EZ ACCESS

benzocaine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70742-001
Route of Administration	Topical		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	1417 mg in 30 mL

Inactive Ingredients	
Ingredient Name	**Strength**
METHYLPARABEN (UNII: A2I8C7HI9T)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70742-001-01	30 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/15/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	06/15/2016	

Labeler - Product Max Group Inc (134893911)

Registrant - Product Max Group Inc (134893911)

Revised: 8/2016 Product Max Group Inc