# ANAL GLIDE- benzocaine gel 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.

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anal GLIDE - 092

## Drug Facts Active Ingredient

Benzocaine 5%

## **Purpose**

Topical Analgesic

## Keep out of reach of children

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

#### **Directions**

- 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.
- Apply to the affected area up to 6 times daily.

#### Other Information

Do not use if safety seal is broken or missing.

### **Inactive Ingredients**

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

anal GLIDE EXTRA product label

anal GLIDE EXTRA

**Desensitizer Lubricating Gel** 

#### **PREMIUM**

### **Benzocaine Anorectal Gel**

#### Anal Desensitizer

2 FL OZ (60 ml)

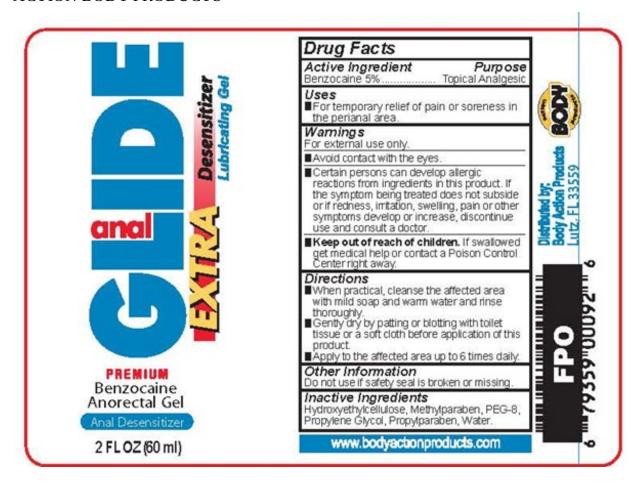
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**Body Action Products** 

Lutz, FL 33559

**ACTION BODY PRODUCTS** 



## ANAL GLIDE

henzocaine gel

benzocaine gel					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (	Source)	NDC:70742-092	
Route of Administration	TOPICAL				
Active Ingredient/Active Moiety					
Ingre	dient Name		Basis of Strengt	h Strength	

BENZOCAINE

2833 mg in 60 mL

**BENZO CAINE** (UNII: U3RS Y48 JW5) (BENZO CAINE - UNII: U3RS Y48 JW5)

Inactive Ingredients			
Ingredient Name	Strength		
HYDROXYETHYL CELLULOSE (140 MPA.S AT 5%) (UNII: 8136 Y38 GY5)			
METHYLPARABEN (UNII: A2I8C7HI9T)			
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
PROPYLPARABEN (UNII: Z8IX2SC1OH)			
WATER (UNII: 059QF0KO0R)			

l	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 ND 01	C:70742-092-	60 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