**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**

**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 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)

www.bodyactionproducts.com

Distributed by:

Body Action Products

Lutz, FL 33559

ACTION BODY PRODUCTS


# ANAL GLIDE

benzocaine gel

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:70742-092 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **BENZOCAINE** (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5) | BENZOCAINE | 2833 mg in 60 mL |

| Inactive Ingredients | |
|---|---|
| Ingredient Name | Strength |
| **HYDROXYETHYL CELLULOSE (140 MPA.S AT 5%)** (UNII: 8136Y38GY5) | |
| **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:70742-092-01 | 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