# BODY ACTION PRODUCTS ENDLESS LOVE FOR MEN- benzocaine gel PRODUCT MAX GROUP INC

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#### **BODY ACTION PRODUCTS: Endless Love for Men**

### **Drug Facts**

#### **Active Ingredient**

Benzocaine 5%

### Purpose

Male Genital Desensitizer

#### Uses

• Helps in the prevention of premature ejaculation.

#### Warnings

For external use only.

- Avoid contact with the eyes.
- Premature ejaculation may be due to a condition requiring medical supervision. If this
  product, used as directed, does not provide relief, discontinue use and consult a
  doctor.
- If you or your partner develop a rash or irritation, such as burning or itching, discontinue use. If symptoms persist, consult a doctor.

### Keep out of reach of children.

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

#### **Directions**

- Apply a small amount to head and shaft of penis before intercourse.
- Wash product off after intercourse.

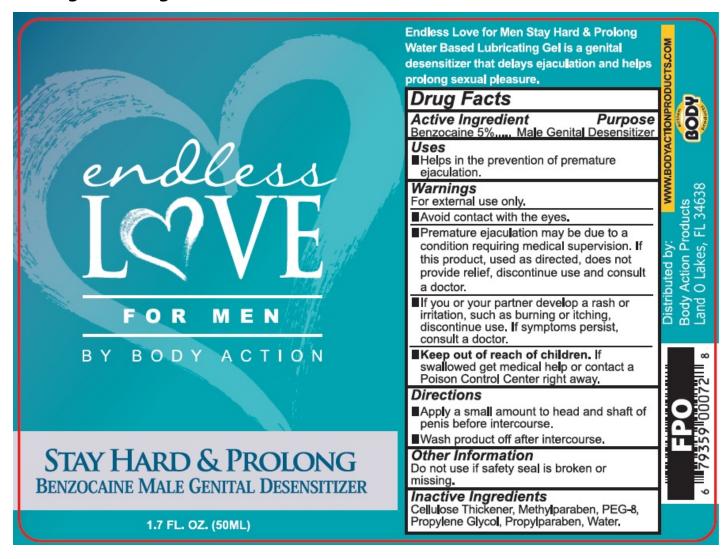
#### Other Information

Do not use if safety seal is broken or missing.

### Inactive Ingredients

Cellulose Thickener, Methylparaben, PEG-8, Propylene Glycol, Propylparaben, Water.

#### **Package Labeling:**



### **BODY ACTION PRODUCTS ENDLESS LOVE FOR MEN**

benzocaine gel

<b>Product Information</b>	Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70742-284	
Route of Administration	TOPICAL			

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

Inactive Ingredients		
Ingredient Name Strength		
METHYLPARABEN (UNII: A2I8C7HI9T)		
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		

PROPYLPARABEN (UNII: Z8I	X2SC1OH)	
WATER (UNII: 059QF0KO0R)		

l	P	Packaging			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:70742-284- 00	50 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/29/2022	

<b>Marketing In</b>	rketing Information		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/29/2022	

## Labeler - PRODUCT MAX GROUP INC (134893911)

Revised: 11/2023 PRODUCT MAX GROUP INC