

**BODY ACTION PRODUCTS PROLONG LUBRICATING- benzocaine gel
PRODUCT MAX GROUP INC**

BODY ACTION PRODUCTS: Prolong Lubricating Gel

Drug Facts

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 under cap is broken or missing.

Inactive Ingredients

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

Package Labeling:

Drug Facts	
Active Ingredient	Purpose
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www.bodyactionproducts.com

Distributed by:
Body Action Products
Land O Lakes, FL 34638



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**Benzocaine
Male Genital
Desensitizer**

FOR MEN

PROLONG

**Helps Prolong Sexual Pleasure
Lubricating Gel**

NET 2 FL OZ (60ml)

BODY ACTION PRODUCTS PROLONG LUBRICATING				
benzocaine gel				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70742-288	
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
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:70742-288-00	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

PRODUCT MAX GROUP INC