

**BODY ACTION PRODUCTS CLIMAX CONTROL GEL- benzocaine gel
PRODUCT MAX GROUP INC**

BODY ACTION PRODUCTS: Climax Control Gel

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 under cap is broken or missing.

Inactive Ingredients

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

Package Labeling:

Drug Facts		  www.bodyactionproducts.com Distributed by: Body Action Products Land O Lakes, FL 34638		 Climax Control Gel Helps in Temporarily Prolonging the Time Until Ejaculation Benzocaine Male Genital Desensitizer 2 fl. oz. (60 mL)
Active Ingredient	Purpose			
Benzocaine 5% 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.				
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BODY ACTION PRODUCTS CLIMAX CONTROL GEL

benzocaine gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70742-282
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

Marketing Start Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70742-282-00	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	M017	04/29/2022		

Labeler - PRODUCT MAX GROUP INC (134893911)

Revised: 11/2023

PRODUCT MAX GROUP INC