

HIGHER CONTROL MALE CLIMAX CONTROL LUBRICATING- benzocaine gel
Product Max Group Inc

HIGHER CONTROL Male Climax Control Lubricating

□ 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.

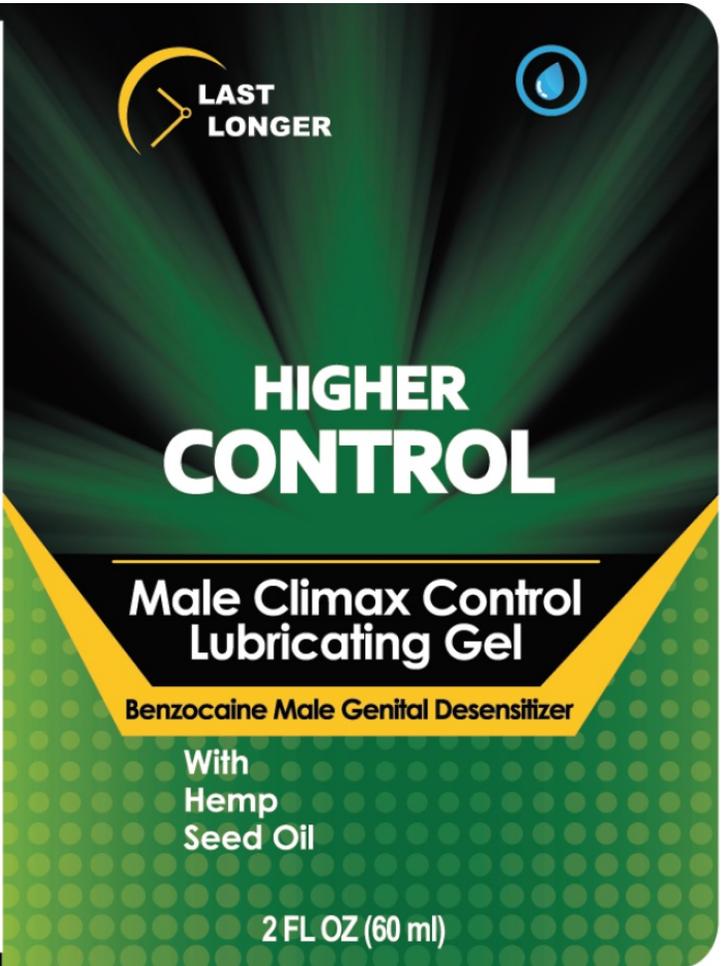
Other Information

Do not use if safety seal on bottle is broken or missing.

Inactive Ingredients

Cannabis Sativa (Hemp) Seed Oil, Hydroxyethylcellulose, Methylparaben, PEG-8, Polysorbate 20, Propylene Glycol, Propylparaben, Water.

Package Labeling:

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HIGHER CONTROL MALE CLIMAX CONTROL LUBRICATING			
benzocaine gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70742-180
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
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)			
METHYLPARABEN (UNII: A2I8C7HI9T)			
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)			
POLYSORBATE 20 (UNII: 7T1F30V5YH)			

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70742-180-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2018	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/01/2018	

Labeler - Product Max Group Inc (134893911)

Revised: 11/2023

Product Max Group Inc