CLIMAX CONTROL BENZOCAINE- benzocaine gel
Natureplex LLC

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.

Climax Control Gel
Benzocaine 7.5%

Drug Facts

Active ingredient

Benzocaine 7.5%

Purpose

Male genital desensitizer

Uses

- helps in temporarily prolonging time until ejaculation

Warnings

For external use only.

When using this product

- avoid contact with eyes

Stop use and ask a doctor if

- this product, used as directed, does not provide relief. Premature ejaculation may be due to conditions requiring medical supervision
- you or your partner develop a rash or irritation, such as burning or itching

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away: 800-222-1222.

Directions

- apply a small amount to head and shaft of penis before intercourse, or use as directed by a physician
- wash product off after intercourse

Other information

- store at 15 to 30°C (59 to 86°F)
- do not use if seal on tube is punctured or missing
- contents filled by weight, not volume

Inactive ingredients

hydroxypropylcellulose, PEG-8, propylene glycol

Questions or comments?

866-323-0107 or www.natureplex.com

PRINCIPAL DISPLAY PANEL - 28 g Tube Carton

Helps
Prolong
Sexual
Pleasure

MAXIMUM STRENGTH
DESENSITIZER
CLIMAX CONTROL GEL

NET WT. 1 Oz. (28g)

benzocaine gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67234-038
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Benzocaine (UNII: U3RSY48JW5) (Benzocaine - UNII:U3RSY48JW5)	Benzocaine	0.075 g in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:67234-038-01	1 in 1 CARTON	01/25/2016	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part333B	01/25/2016	

Labeler - Natureplex LLC (062808196)

Establishment

Name	Address	ID/FEI	Business Operations
Natureplex LLC		062808196	MANUFACTURE(67234-038)

Revised: 10/2017 Natureplex LLC