MAGIC STAMINA- benzocaine liquid
Product Max Group Inc

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.

Magic STAMINA - 088

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 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 tab is broken or missing.

Inactive Ingredients

Methylparaben, PEG-8, Propylene Glycol, Propylparaben, Water

Magic STAMINA product label

Benzocaine Male genital Desensitizer

Magic STAMINA®

Climax contral spray

1.0 FL. OZ. (30 mL)

Distributed By: Body Action Products, Lutz FL 33559

www.bodyactionproducts.com

MAGIC STAMINA

benzocaine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70742-088
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	1417 mg in 30 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:70742-088-01	30 mL in 1 PACKAGE; Type 0: Not a Combination Product	06/15/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	06/15/2016	

Labeler - Product Max Group Inc (134893911)

Registrant - Product Max Group Inc (134893911)

Revised: 6/2016 Product Max Group Inc