

MANDELAY MALE GENITAL DESENSITIZER- benzocaine gel
Majestic Drug Co., 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.

MANDELAY®

Drug Facts

Active Ingredient

Benzocaine 7.5%

Purpose

Male genital desensitizer

Uses

- Helps in temporarily prolonging the 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 condition 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.

Directions

- Apply a small amount to head and shaft of penis 2-3 minutes before intercourse, or use as directed by a physician.
- Wash product off after intercourse.

Inactive Ingredients

Carbowax, Carbomer 974P.

PRINCIPAL DISPLAY PANEL - 28.35 g Tube Box

MANDELAY®

Climax Control Gel

NET WT 1.0 OZ. (28.35g)

Climax Control Gel

Make It Last!

MANDELAY®

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Manufactured for
Majestic Drug Co., Inc.
P.O.Box 490
South Fallsburg, NY 12779 USA
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www.mandelay.com



MANDELAY®
Climax Control Gel

MANDELAY®
Benzocaine Male
Genital Desensitizer
formulated
to delay ejaculation.

MANDELAY®
Maximum strength
to help prolong
sexual pleasure.

Compatible with natural
rubber latex, polyisoprene,
and polyurethane condoms

NET WT 1.0 OZ. (28.35g)





MANDELAY MALE GENITAL DESENSITIZER

benzocaine gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10705-077
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Benzocaine (UNII: U3RSY48JW5) (Benzocaine - UNII:U3RSY48JW5)	Benzocaine	2.13 g in 28.35 g

Inactive Ingredients

Ingredient Name	Strength
Carbomer Homopolymer Type B (Allyl Pentaerythritol Crosslinked) (UNII: HHT01ZNK31)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10705-077-01	1 in 1 BOX	10/31/1995	
1	NDC:10705-077-28	28.35 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	part348	10/31/1995	

Labeler - Majestic Drug Co., INC. (001496777)