PLEASURE BALM KAMA SUTRA- benzocaine gel Cosmetic Enterprises Ltd

Pleasure Balm

Active Ingredients Purpose

Benzocaine 5.0 % Male genital desensitizer

Use

Helps in temporarily prolonging time until ejaculation

Keep out of reach of children.

Discontinue use and consult a doctor if

This product, used as directed does not provide relief

You or your partner develops a rash or irritation, such as burning or itching

Condition worsens or if symptoms persist for more than 7 days or clear up and occur again within a few days

The following symptoms appear: weakness, confusion, headache, difficulty breathing and/or pale, gray or blue colored skin as these may be signs of methemoglobinemia, a rare disorder which may appear up to 2 hours of use

Warnings

DO NOT use if allergic to Benzocaine or any other ingredient in this product

If swallowed seek medical assistance or contact a poison control or contact a poison control Center right away.

Premature ejaculation may be due to condition requiring medical supervision

Consult a doctor prior to use if partner is pregnant

Avoid contact with eyes: if this happens, rinse thoroughly with water

Known adverse reactions Hypersensitivity/allergy has been known to occur, in such a case, discontinue use.

Directions for use

Adults, 18 and over. Apply a small amount to head and shaft of penis 5-10 minutes before intercourse, or use as directed by a health care practitioner. Use the smallest amount to achieve the desired result. Wipe off any excess gel before intercourse. Wash product off after intercourse. For external Use only.

Inactive Inagredients: Blue 1 (CI 42090), Disodium EDTA, Hydroxyethylcellulose, Mentha Piperita (peppermint oil), PEG-8, Polysorbate 60, Propylene Glycol, Sodium Saccharin, Water (Aqua/Eau), Yellow 5 (CI 19140), Yellow 6 (CI 15985)

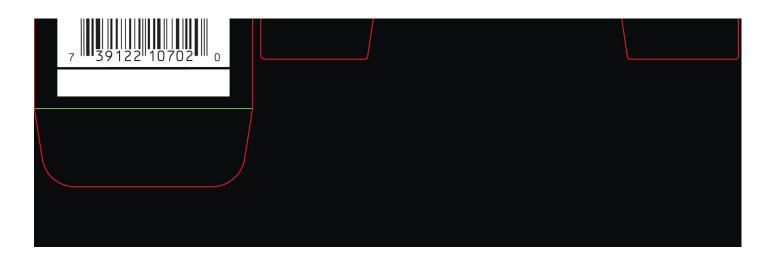
Pleasure Balm

Prolonging Gel for Men

Desensibilisateur Genital Pour hommes

0.4 fl oz 12 ml





PLEASURE BALM KAMA SUTRA

benzocaine gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:56152-1470

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient NameBasis of StrengthStrengthBENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)BENZOCAINE5 g in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
EDETATE DISODIUM (UNII: 7FLD91C86K)		
CETYL HYDROXYETHYLCELLULOSE (350000 MW) (UNII: T7SWE4S2TT)		
MENTHA PIPERITA (UNII: 79M2M2UDA9)		
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)		
POLYSORBATE 60 (UNII: CAL22UVI4M)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
WATER (UNII: 059QF0KO0R)		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		

l	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:56152- 1470-1	12 mL in 1 TUBE; Type 0: Not a Combination Product	12/13/2018		
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Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	09/12/2013		

Labeler - Cosmetic Enterprises Ltd (017701475)

Revised: 10/2023 Cosmetic Enterprises Ltd