PROLAYED- lidocaine spray MATRIX MIXOLOGY, INC

Prolayed

Drug Facts

Active Ingredient (in each spray)

Lidocaine 10 mg

Purpose

Male genital desenitizer

Use

Helps in temporarily slowing the onset of ejaculation.

Warnings

For external use only

When using this product

avoid contact with the eyes.

Stop use and ask a doctor if

- this product, used as directed, does not provide relief. Premature ejaculation may be due to a 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 3 or more sprays, not to exceed 10, to head and shaft of penis before intercourse, or use as directed by a doctor.
- Wash product off after intercourse.

Inactive Ingredients

acetic acid, ascorbic acid, chlorhexidine gluconate, cholecalciferol, dimethyl sulfoxide, dipropylene glycol, glucono delta lactone, glycerin, histidine, hydroxyethylcellulose, magnesium stearate, methylparaben, sodium hydroxide, sorbic acid, stearic acid, water

Package Labeling:

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supervision

Directions

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PROLAYED

male genital desensitizer

topical spray

Lidocaine 10 mg in 1 gram

0.5 FLOZ (15ml)

MATRIX MIXOLOGY, INC 11024 BALBOA BLVD STE 50 GRANADA HILLS CA 91354

PROLAYED

lidocaine spray

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79218-001

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient NameBasis of StrengthStrengthLIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)LIDOCAINE10 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
ACETIC ACID (UNII: Q40Q9N063P)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	
CHLORHEXIDINE GLUCONATE (UNII: MOR84MUD8E)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
DIMETHYL SULFOXIDE (UNII: YOW8V9698H)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
GLYCERIN (UNII: PDC6A3C0OX)	
HISTIDINE (UNII: 4QD397987E)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SORBIC ACID (UNII: X045W989B)	

STEARIC ACID (UNII: 4ELV7Z65AP)	
WATER (UNII: 059QF0KO0R)	

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:79218-001- 01	1 in 1 BOX	07/01/2020				
1		15 mL in 1 BOTTLE; Type 0: Not a Combination Product					

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M017	07/01/2020			

Labeler - MATRIX MIXOLOGY, INC (128885636)

Revised: 12/2023 MATRIX MIXOLOGY, INC