POPSTAR- delay spray spray
Private Label Productions LLC

PLP-Popstar Delay Spray 14.7 mL

Active Ingredient(s)

Lidocaine 10mg .. Male Genital Desensitizer

Purpose

Male Genital Desensitizer

Use

Helps in temporarily slowing the onset of ejaculation.

Warnings

For external use only.

Do not use if you or your partner are allergic to lidocaine or topical anesthetics.

When using this product

Avoid contact with the eyes
Do not spray on broken, irritated, or sensitive skin.

Stop 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.

Ask a doctor or pharmacist before use

If you have liver problems
If your partner is, or may be pregnant.

Keep out of reach of childern

If swallowed, get medical attention or contact a Poison Control Center immediately.

Directions

Gently shake bottle. Hold upright and press pump until spray dispenses. Apply 3 or more sprays, not to exceed 10, to head and shaft of penis 10 minutes betore intercourse or as directed by a doctor. Wash product off after intercourse.

Other information

Remove safety tab before use.
Store at 68°-77° F (20-25° C).

Inactive ingredients

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Purified Water, Propanediol, Alcohol, Aloe Barbadensis Leaf Juice, Laureth 9, PEG 8, Panthenol, Panax Ginseng Root Extract, Hydroxyethylcellulose, Menthyl Lactate, PPG-26-Buteth-26, PEG-40 Hydrogenated Castor Oil, Caprylyl Glycol, Phenoxyethanol,

Ethylhexylglycerin

Package Label - Principal Display Panel

Lorem ipsum

POPSTAR

delay spray spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77632-151
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	10 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
HYDROXYETHYLCELLULOSE (UNII: T4V6TWG28D)	
MENTHYL LACTATE (UNII: 2BF9E65L7I)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
LAURETH-9 (UNII: 0AWH8BFG9A)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PANAX GINSENG ROOT (UNII: CUQ3A77YXI)	
PEG-40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
PEG-8 (UNII: B697894SGQ)	
PANTHENOL (UNII: WV9CM0O67Z)	
PPG-26-BUTETH-26 (UNII: 2II1K6TZ4P)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77632-151-12	75 in 1 BOX	01/01/2025	
1		14.7 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		
2	NDC:77632-151-11	14.7 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2025	

Labeler - Private Label Productions LLC (046278265)

Revised: 8/2025 Private Label Productions LLC