CLIMAX SHIELD DELAY- benzocaine liquid
PRODUCT MAX GROUP INC

Climax Shield Delay Spray

Drug Facts

Active Ingredient

Benzocaine 7.5%

Purpose

Male Genital Desensitizer

Uses

- Helps in the prevention of premature ejaculation.

Warnings

For external use only.

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.

Avoid contact with the eyes.

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 the head and shaft of the penis before intercourse, or use as directed by a doctor. Wash product off after intercourse.

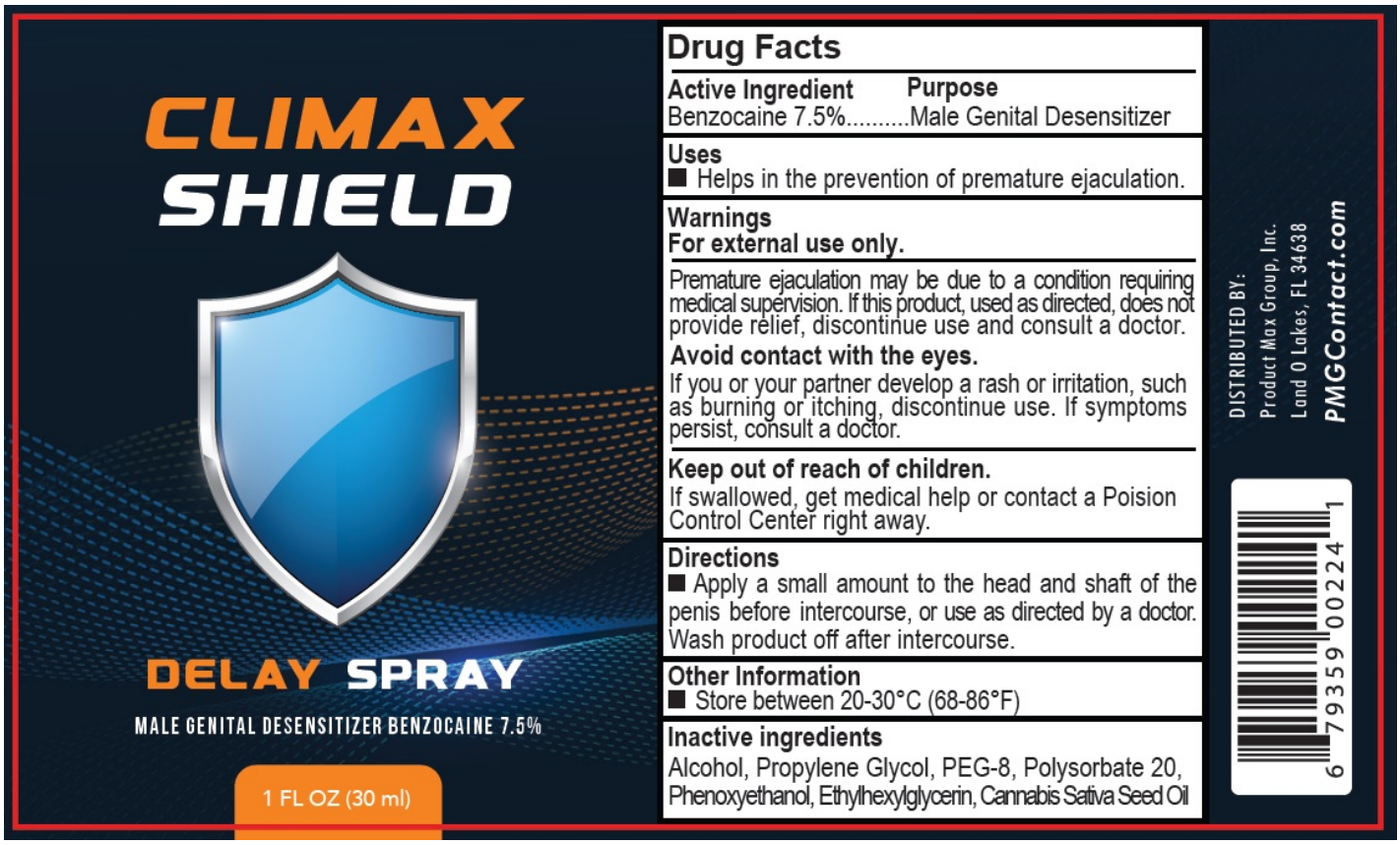
Other Information

- Store between 20-30°C (68-86°F)

Inactive ingredients

Alcohol, Propylene Glycol, PEG-8, Polysorbate 20, Phenoxyethanol, Ethylhexylglycerin, Cannabis Sativa Seed Oil

Package Labeling:



CLIMAX SHIELD DELAY

benzocaine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70742-598
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	75 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70742-598-00	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/28/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/28/2025	

Labeler - PRODUCT MAX GROUP INC (134893911)

Registrant - Pure Source, LLC (080354456)

Revised: 5/2025

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