LUBELIFE CLIMAX CONTROL DELAY- benzocaine spray
CC Wellness LLC

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.

Benzocaine 7.5%

Male Genital Desensitizer

Uses:
- For temporary male genital desensitization
- Helps to slow the onset of ejaculation

For external use only

Avoid contact with the eyes

Allergy alert: do not use this product if you are allergic (sensitive) to local anesthetics or any of the other listed ingredients

Do not use on broken or inflamed skin

Ask a doctor before use:
- If your partner is pregnant
- If you have or have had liver or kidney problems

Ask a doctor or pharmacist before use if you are already taking prescribed drugs

When using this product
- Do not get into eyes or nostrils
- Do not inhale
- Do not exceed a maximum of 20 sprays in 24 hours
- Always use the minimum amount effective for you
- Allow 5-15 minutes to dry prior to intercourse

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. If you or your partner develop a rash or irritation, such as burning or itching, discontinue use. If symptoms persist, consult a doctor.

Can be used for sexual intercourse and sex play when applied in accordance with usage instructions. Only use in accordance with the instructions, seek medical attention immediately in case of overdose.

Keep out of reach of children

Directions:

- Apply a small amount to head and shaft of penis before intercourse, or use as directed by a doctor
- Wash product off after intercourse

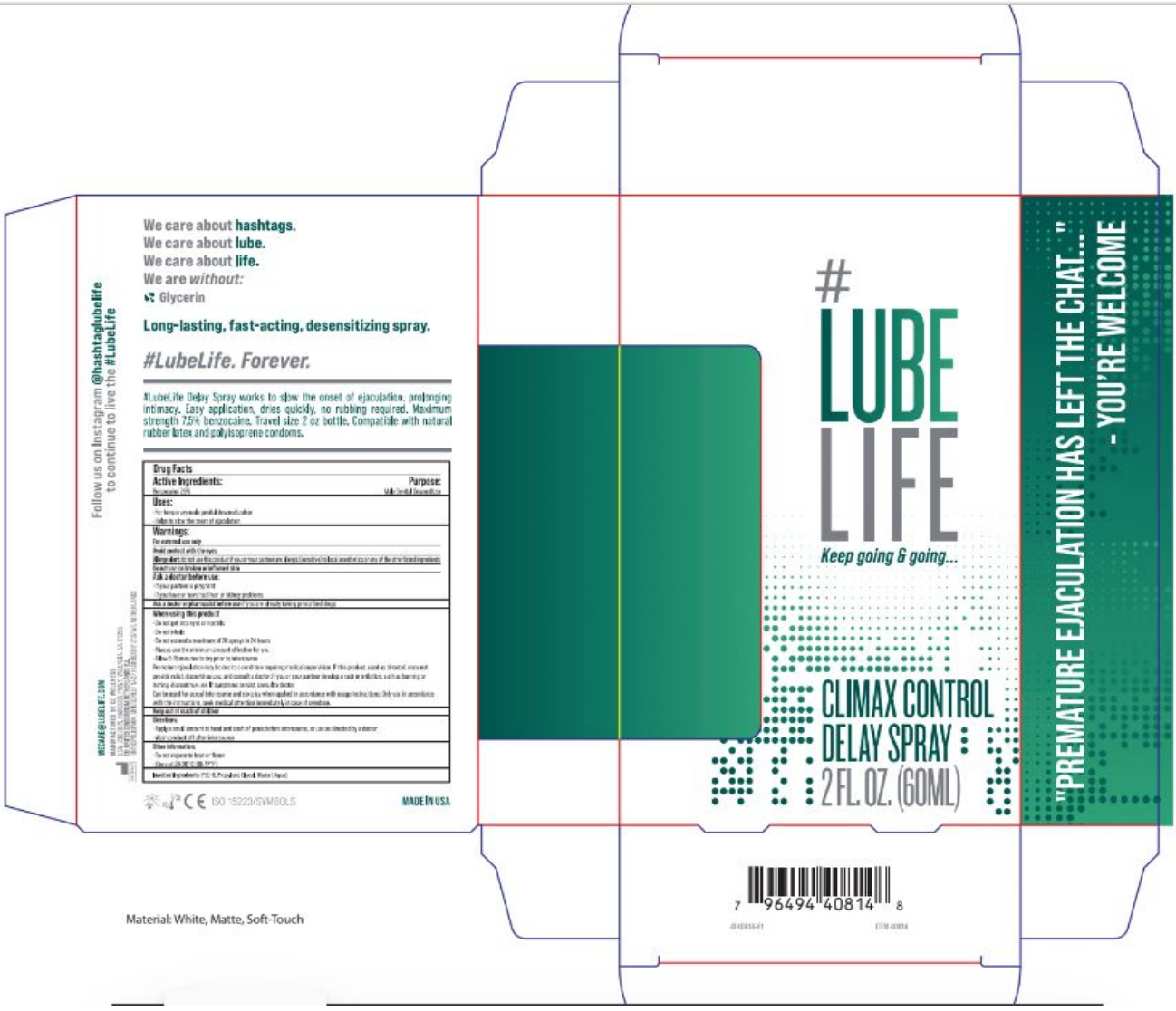
Other information:

- Do not expose to heat or flame
- Store at 20-30°C (68-77°F)

Inactive Ingredients:

PEG-8, Propylene Glycol, Water (Aqua)

Principal Display Panel



LUBELIFE CLIMAX CONTROL DELAY

benzocaine spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71683-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	7.5 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71683-005-01	1 in 1 BOX	01/01/2022	
1	NDC:71683-005-02	60 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	01/01/2022	

Labeler - CC Wellness LLC (067692292)

Establishment

Name	Address	ID/FEI	Business Operations
CC Wellness LLC		067692292	manufacture(71683-005)

Revised: 6/2022 CC Wellness LLC