INTIMATE WIPE- benzocaine liquid
Veru Inc.

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.

Drug Facts

Active ingredient

Benzocaine USP 4%

Purpose

Male Genital Desensitizer

Use

Helps in temporarily prolonging time until 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 supervision.
- You or your partner develop a rash or irritation, such as burning or itching
- Symptoms persist.

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

Inactive ingredients

ethyl alcohol (SDA 40B), propylene glycol, purified water

Principal Display Panel – Box Label

PLAYBOY

INTIMATE WIPES

MALE GENITAL DESENSITIZER

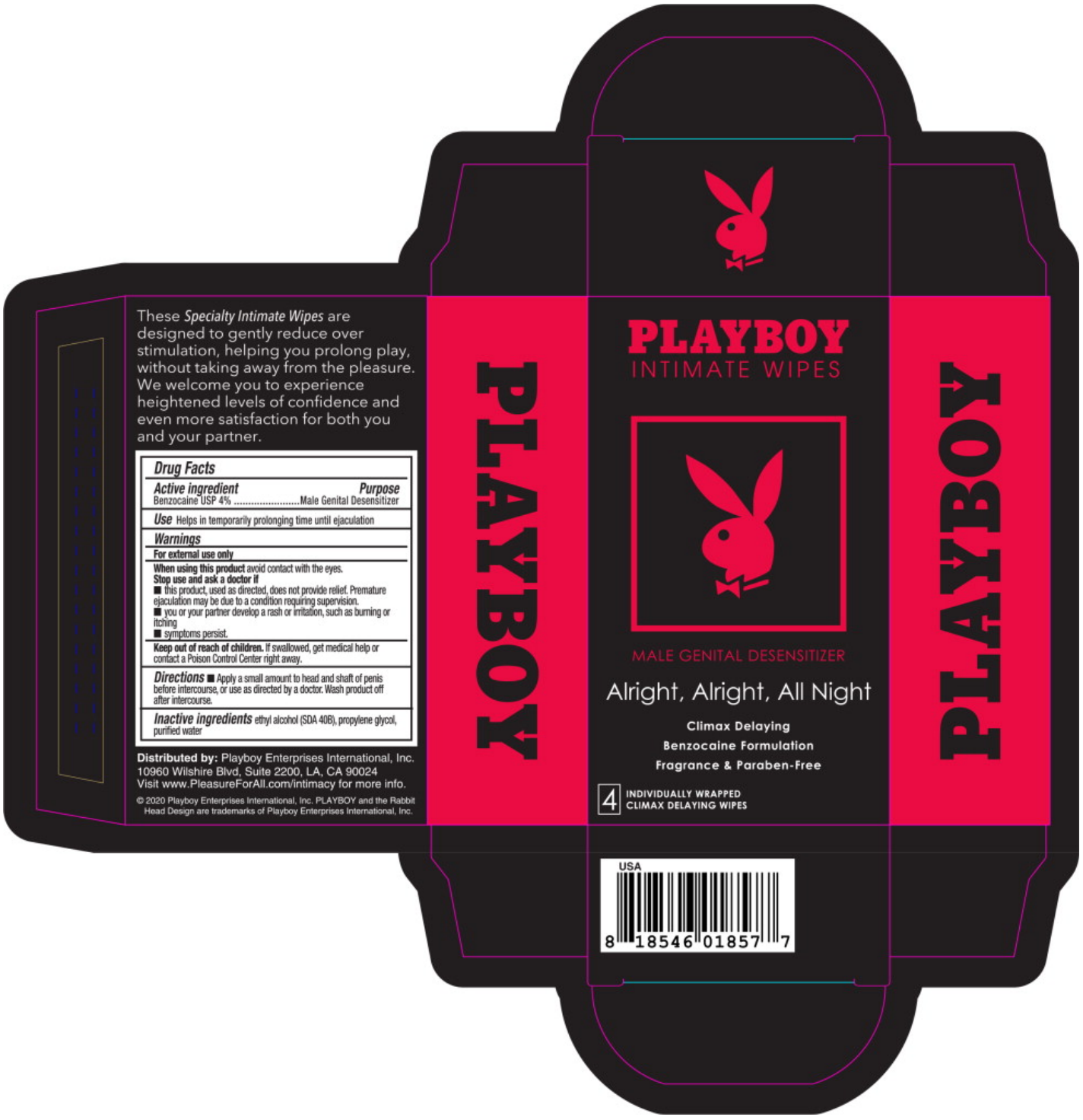
Alright, Alright, All Night

Climax Delaying

Benzocaine Formulation

Fragrance & Paraben-Free

4 INDIVIDUALLY WRAPPED
CLIMAX DELAYING WIPES

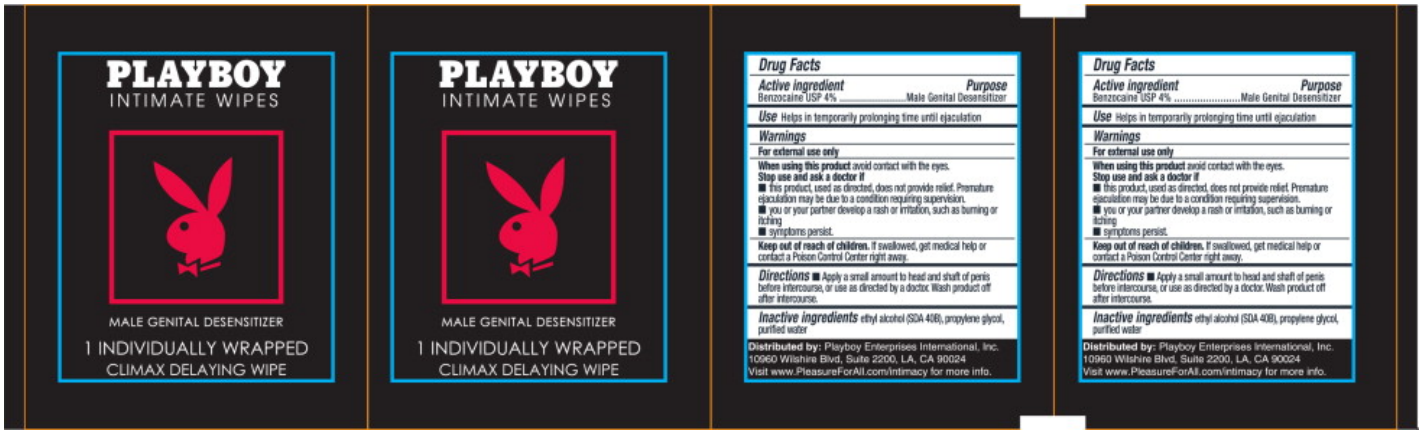


Principal Display Panel – Packet Label

PLAYBOY

INTIMATE WIPES

MALE GENITAL DESENSITIZER
1 INDIVIDUALLY WRAPPED
CLIMAX DELAYING WIPE



INTIMATE WIPE

benzocaine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69681-434
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
benzocaine (UNII: U3RSY48JW5) (benzocaine - UNII:U3RSY48JW5)	benzocaine	4 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
alcohol (UNII: 3K9958V90M)	
water (UNII: 059QF0KO0R)	
propylene glycol (UNII: 6DC9Q167V3)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69681-434-04	4 in 1 BOX	06/08/2020	
1		1.2 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part348	06/08/2020	

Labeler - Veru Inc. (055300578)

Registrant - Safetec of America, Inc. (874965262)

Revised: 6/2020 Veru Inc.