PREBOOST- benzocaine liquid
The Female Health Company d/b/a Veru Healthcare

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.

PREBOOST®

Drug Facts

Active Ingredient

Benzocaine USP 4%

Purpose

Male Genital Desensitizer

Use

Helps in temporarily prolonging time until ejaculation

Warnings

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

Inactives

Purified water, Ethyl Alcohol (SDA 40B), Propylene Glycol

PRINCIPAL DISPLAY PANEL - 10 Wipes Packet Box

Clinically Proven to Make Sex Last Longer

PREBOOST®
KEEP IT GOING

10 Single-Use Wipes
Male desensitizing wipes for
prolonging the time until ejaculation

H Fisch MD

Created by **Dr. Harry Fisch**, Named
"One of America's Best Doctors"

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Easy-to-use • FDA Compliant

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veru HEALTHCARE

Manufactured for
The Female Health Company
d/b/a Veru Healthcare
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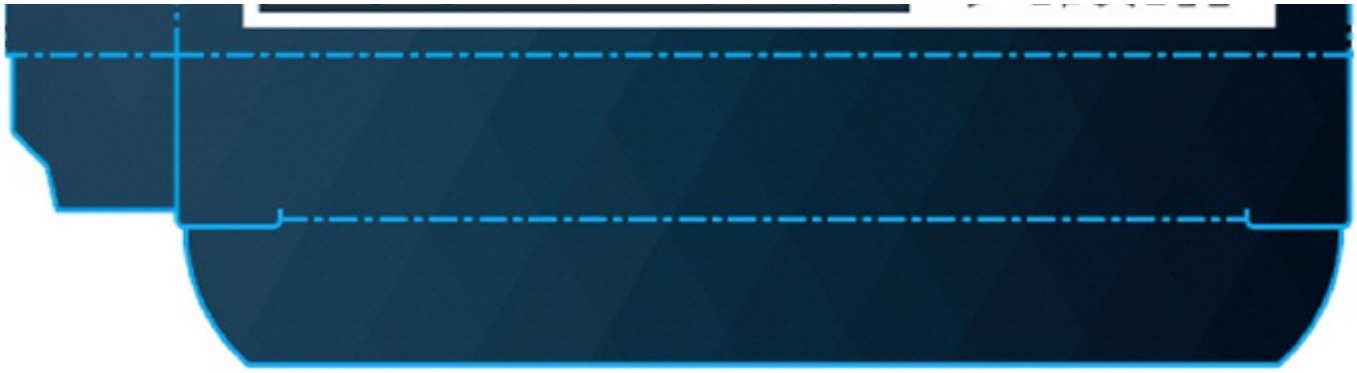

PREBOOST

benzocaine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69681-432
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:69681-432-10	10 in 1 BOX	04/05/2017	
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 not final	part348	03/30/2015	

Labeler - The Female Health Company d/b/a Veru Healthcare (055300578)

Establishment

Name	Address	ID/FEI	Business Operations

Safetec of America, Inc.		874965262	MANUFACTURE(69681-432)