MINT RX ENDURANCE WIPES- benzocaine liquid
PHARMAMEDRX

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.

MINT RX ENDURANCE WIPES
BENZOCAINE USP 5%

Active ingredient

Benzocaine 5%

Purpose

Male genital desensitizer

Keep out of reach of children

If swallowed, get medical help or contact a poison control center right away

Use

Helps in temporarily prolonging time until ejaculation

Warnings

For external use ony

When using this product avoid contact with 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 medical supervision

You or your partner develop a rash or irritation such as burning or itching

Symptoms persist

Directions

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

Inactive ingredients

Purified water, ethyl alcohol, glycerin, Vitamin E, phenoxyethanol

MINT RX ENDURANCE WIPES
BENZOCAINE USP 5%

MINT RX ENDURANCE WIPES

benzocaine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77691-433
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	5 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
TOCOPHEROL (UNII: R0ZB2556P8)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:77691-433-08	8 in 1 PACKAGE	05/20/2020	
1	NDC:77691-433-03	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	05/20/2020	

Labeler - PHARMAMEDRX (063814205)

Registrant - PHARMAMEDRX (063814205)

Establishment

Name	Address	ID/FEI	Business Operations
PHARMAMEDRX		063814205	manufacture(77691-433)

Revised: 5/2020 PHARMAMEDRX