FEMININE ANTI-ITCH CREME- benzocaine, resorcinol cream Quality Choice (CHAIN DRUG MARKETING ASSOCIATION)

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.

Quality Choice Maximum Strength Feminine Anti-itch Cream

Drug Facts Active ingredients

Benzocaine 20%

Resorcinol 3%

Purpose

External analgesic

Keep out of reach of children

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Use

temporarily relieves itching

Warnings

For external use only

When using this product

avoid contact with the eyes

Stop use and ask a doctor if

condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days

Do not apply Directions

Do not apply over large area of the body.

adults and children 12 years and older apply a fingertip amount (approximately 1-inch strip) to the affected area not more than 3 to 4 times daily

children under 12 years ask a doctor

Other Information

store at 20° -25° C (68° -77° F)

Inactive Ingredients

aloe barbadensis leaf extract, carbomer, cetyl alcohol, cholecalciferol, glyceryl monostearate, isopropyl myristate, isopropyl palmitate, isopropyl sterate, lanolin, methylparaben, mineral iol, PEG-100 stearate, propylene glycol, purified water, retinyl palmitate, sodium hydroxide, sodium sulfite, tocopheryl acetate, trisodium HEDTA

Questions or comments?

800-935-2362

Quality Choice Maximum Strength Feminine Anti-itch Cream





in the Vaginal Area

Feminine Anti-Itch Creme

Relief from Intense Burning and Itching

Plus Aloe

20% Benzocaine 3% Resorcinol External Analgesic



Made in India M 92-89064 02/23

Model 89-201 /23 WC\$10052

This product is not manufactured or distributed by Combe Incorporated, owner of the registered trademark. Yagis? Maximum Strength Meditated Anti-Inch Creme.*

Drug Facts

Active ingredients Purposes Berzocaine 20%External analgesic Resorcinol 3%External analgesic

Use temporarily relieves itching

Warnings

For external use only

When using this product avoid contact with eyes

Stop use and ask a doctor if condition worsens, or if symptoms persist for more than 7 days or clear upand occur again within a few days.

Do not apply overlarge areas of the body

Keep out of reach of children. If swallowed, getmedical help or contact a Poison Control Center right away (1-800-222-1222)

times daily

ask a doctor

Direc frons adults and children 12 years

and older

apply a tinger to amount (approximately 1-inch strip) to the affected area not more than 3 to 4

children under 12 years

Other Information

Inactive ingredients also barbadensis leaf extract, carbomer, cetyl alcohol, cholecalciterol, glyceryl m onostearate, isopropyl m yristate, isopropyl palmitafe, isopropyl stearafe, ianolin, methylparaben, mineral oil, FEG-100 stearafe, propylene glycol, purified water, refinyl palmitate, sodium hydroxide, sodium sultite, to copheryl acetate, trisodium HEDTA

Questions or comments? 1-900-935-2362



1 **OZ** (NetWt283g)

TOPICAL

FEMININE ANTI-ITCH CREME

benzocaine, resorcinol cream

Route of Administration

Product Information	oduct Information		
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-949

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength

BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	5.67 g in 28 g
RESORCINOL (UNII: YUL4L094HK) (RESORCINOL - UNII:YUL4L094HK)	RESORCINOL	0.85 g in 28 g

Inactive Ingredients	
Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
ISOPROPYL STEARATE (UNII: 43253ZW1MZ)	
LANOLIN (UNII: 7EV65EAW6H)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM SULFITE (UNII: VTK01UQK3G)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TRISODIUM HEDTA (UNII: K3E0U7O8KI)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-949- 01	28 g in 1 TUBE; Type 0: Not a Combination Product	02/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/01/2020	
	part348	02/01/2020	

Labeler - Quality Choice (CHAIN DRUG MARKETING ASSOCIATION) (011920774)

Revised: 2/2023 Quality Choice (CHAIN DRUG MARKETING ASSOCIATION)