

**MAXIMUM OTC STRENGTH FEMININE ANTI-ITCH CREME- benzocaine,
resorcinol cream**

Quality Choice (CHAIN DRUG MARKETING ASSOCIATION)

Quality Choice Maximum OTC Strength Feminine Anti-itch Creme

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: do not use; consult a doctor

Other Information

store at 20° -25° C (68° -77° F)

Inactive Ingredients

aloe vera leaf, alpha-tocopherol, butylated hydroxytoluene, carbomer, cetyl alcohol
cholecalciferol, glyceryl monostearate,

isopropyl myristate, isopropyl palmitate, isopropyl stearate, mineral oil, PEG 100
stearate, PEG 4000, phenoxy ethanol, propylene glycol, sodium hydroxide, sodium lauryl
sulfate, trisodium edta, vitamin A palmitate, water

Questions or comments?

1-866-326-1313

Quality Choice Maximum OTC Strength Feminine Anti-itch Creme



Maximum OTC Strength
**Feminine
Anti-Itch Creme**



NDC 83324-019-01

Compare to the Active Ingredients in
Yagisil[®] Maximum Strength
Anti-Itch Creme

Helps Relieve Intense Burning and Itching
in the Vaginal Area

Maximum
OTC Strength
**Feminine
Anti-Itch
Creme**

**Relief from
Intense Burning
and Itching**

Plus Also

2.0% Benzocaine
3% Resorcinol
External Analgesic

1 OZ (Net Wt 28.3g)



Distributed by COMA, Inc
Nora, MI 48276
www.coma.com
Distributor: 800-935-2382
Made in India Model 89-201
92-89084 11/23 VC810072



LOT: 510072MMYY

EXP: YYYY-MM-DD

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Drug Facts

Active ingredients Purposes
Benzocaine 2.0% 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 up and occur again within a few days

Do not apply over large areas of the body

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

Directions

adults and children 2 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 2 years do not use; consult a doctor

Other information

store at 20°-25°C (68°-77°F)

Inactive ingredients

aloe vera leaf, alpha tocopherol, butylated hydroxytoluene, carbomer, cetyl alcohol, cholecalciferol, glyceryl mono stearate, isopropyl myristate, isopropyl stearate, mineral oil, PEG 100 stearate, PEG 4000, phenoxy ethanol, propylene glycol, sodium hydroxide, sodium lauryl sulfate, trisodium edta, vitamin a palmitate, water

Questions or comments?

1-866-326-1313

*This product is not manufactured or distributed by Combe Incorporated, owner of the registered trademark Yagisil[®]

MAXIMUM OTC STRENGTH FEMININE ANTI-ITCH CREME

benzocaine, resorcinol cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-019
Route of Administration	TOPICAL		

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: YUL4LO94HK) (RESORCINOL - UNII:YUL4LO94HK)	RESORCINOL	0.85 g in 28 g

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CHOLECALCIFEROL (UNII: 1C6V77QF41)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
ISOPROPYL STEARATE (UNII: 43253ZW1MZ)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
POLYETHYLENE GLYCOL 4000 (UNII: 4R4HFI6D95)	
MINERAL OIL (UNII: T5L8T28FGP)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	
EDETATE TRISODIUM (UNII: 420IP921MB)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-019-01	28.3 g in 1 TUBE; Type 0: Not a Combination Product	01/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/01/2024	

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

Revised: 4/2024

Quality Choice (CHAIN DRUG MARKETING ASSOCIATION)