UP AND UP VAGICAINE- benzocaine, resorcinol cream Target Corporation

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.

Target Corporation Vagicaine Drug Facts

Active ingredients

Benzocaine 20%

Resorcinol 3%

Purpose

External analgesic

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

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 12 years and older	apply a fingertip amount (approximately 1-inch strip) to the
	affected area not more than 3 to 4 times daily. Clean nozzle of
	tube by wiping thoroughly before replacing cap. Keep cap
	tightly closed between uses.
children under 12 years	ask a doctor

Other information

store at 20-25°C (68-77°F)

Inactive ingredients

aloe barbadensis leaf juice, carbomer homopolymer type C, cetyl alcohol, cholecalciferol, corn (zea mays) oil, fragrance, glyceryl monostearate, isopropyl myristate, isopropyl palmitate, isopropyl stearate, lanolin alcohol, methylparaben, mineral oil, PEG-100 stearate, propylene glycol, purified water, retinyl palmitate, sodium sulfite anhydrous, tocopheryl acetate, trisodium HEDTA, trolamine

Questions?

Call 1-888-547-7400

Principal Display Panel

maximum strength

vagicaine

anti-itch cream

benzocaine 20% / resorcinol 3%

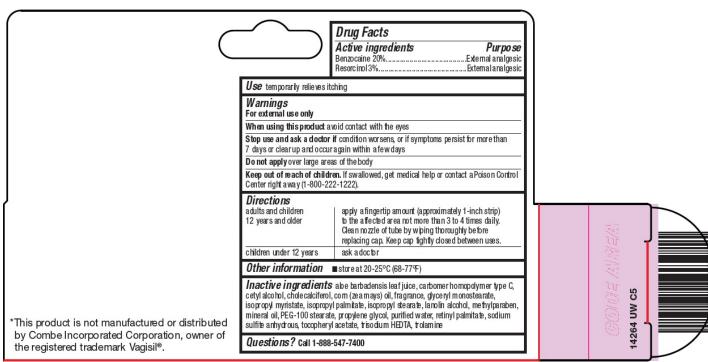
Compare to active ingredients in Vagisil® Maximum Strength

pain relieving cream

instant long-lasting relief from intense itch associated with: antibiotics, discharge, imbalance in pH, irritating vaginal conditions

NET WT 1 OZ (28 g)





UP AND UP VAGICAINE

benzocaine, resorcinol cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-142
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZO CAINE (UNII: U3RS Y48 JW5) (BENZO CAINE - UNII: U3RS Y48 JW5)	BENZOCAINE	20 g in 100 g	
RESORCINOL (UNII: YUL4L094HK) (RESORCINOL - UNII:YUL4L094HK)	RESORCINOL	3 g in 100 g	

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)		
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
CHOLECALCIFEROL (UNII: 1C6V77QF41)		
CORN OIL (UNII: 8470G57WFM)		
GLYCERYL MO NO STEARATE (UNII: 230 O U9 XXE4)		
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)		
ISOPROPYL PALMITATE (UNII: 8 CRQ2TH63M)		
ISOPROPYL STEARATE (UNII: 43253ZW1MZ)		
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
MINERAL OIL (UNII: T5L8T28FGP)		
PEG-100 STEARATE (UNII: YD01N1999R)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
VITAMIN A PALMITATE (UNII: 1D1K0 N0 VVC)		
SODIUM SULFITE (UNII: VTK01UQK3G)		
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)		
TRISO DIUM HEDTA (UNII: K3E0 U7O 8 KI)		
TROLAMINE (UNII: 903K93S3TK)		

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11673-142-64	1 in 1 CARTON	08/25/2009		
1		28 g in 1 TUBE; 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	08/25/2009	

Labeler - Target Corporation (006961700)

Revised: 3/2019 Target Corporation