THERA PLUS MAXIMUM STRENGTH FEMININE ANTI-ITCH- benzocaine, resorcinol cream FOURSTAR GROUP USA, INC.

Thera Plus Maximum Strength Feminine Anti-Itch Cream

Drug Facts

Active ingredients

Benzocaine 20% Resorcinol 3%

Purpose

External Analgesic

Uses

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 persists for more than 7 days or clear up and occur again within a few days.

Do not applyover large areas of the body.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

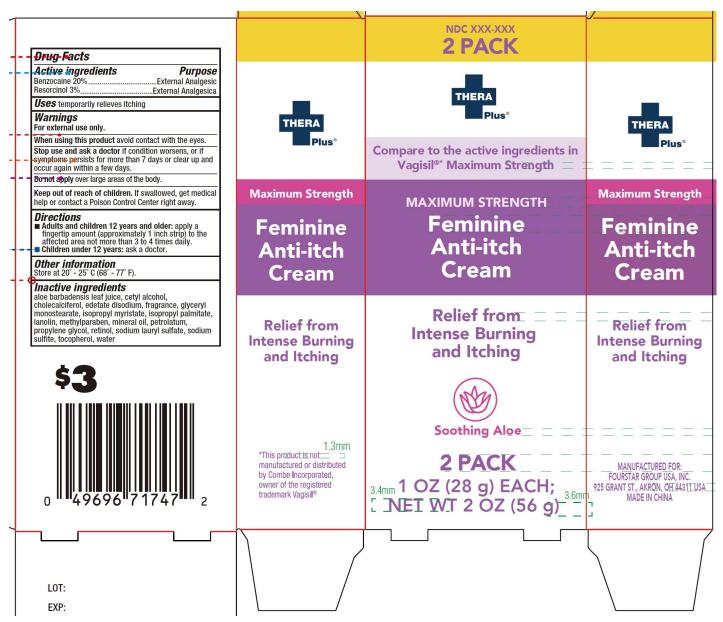
- Adults and children 12 years and older: apply a fingertip amount (approximately 1 inch strip) to the affected area not more than 3 to 4 times daily.
- Children under 12 years:ask a doctor.

Other information

Inactive ingredientsa

aloe barbadensis leaf juice, cetyl alcohol, cholecalciferol, edetate disodium, fragrance, glyceryl monostearate, isopropyl myristate, isopropyl palmitate, lanolin, methylparaben, mineral oil, petrolatum, propylene glycol, retinol, sodium lauryl sulfate, sodium sulfite, tocopherol, water

Package Labeling:



THERA PLUS MAXIMUM STRENGTH FEMININE ANTI-ITCH benzocaine, resorcinol cream Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:80684-142

Route	Ot A	dmir	nistra	ation

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Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	200 mg in 1 g	
RESORCINOL (UNII: YUL4L094HK) (RESORCINOL - UNII:YUL4L094HK)	RESORCINOL	30 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
CHOLECALCIFEROL (UNII: 1C6V77QF41)		
EDETATE DISODIUM (UNII: 7FLD91C86K)		
GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)		
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)		
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)		
LANOLIN (UNII: 7EV65EAW6H)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
MINERAL OIL (UNII: T5L8T28FGP)		
PETROLATUM (UNII: 4T6H12BN9U)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
RETINOL (UNII: G2SH0XKK91)		
SODIUM LAURYL SULFATE (UNII: 368GB5141J)		
SODIUM SULFITE (UNII: VTK01UQK3G)		
TOCOPHEROL (UNII: R0ZB2556P8)		
WATER (UNII: 059QF0KO0R)		

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:80684-142- 00	28 g in 1 TUBE; Type 0: Not a Combination Product	02/26/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/26/2025	

Labeler - FOURSTAR GROUP USA, INC. (140099503)