THE PHARMA C COMPANY MEDICATED FOAM WITH WITCH HAZEL- witch hazel liquid
Kleen Test Products Corporation

The Pharma C Company Medicated Foam with Witch Hazel

Drug Facts

Active ingredient

Witch hazel 50%

Purpose

Hemorrhoidal astringent

Uses

- temporarily relieves the local itching and discomfort associated with hemorrhoids
- aids in protecting irritated anorectal areas
- temporarily relieves irritation and burning

Warnings

For external use only.

When using this product

- do not use more than directed unless told to do so by a doctor
- do not put directly in the rectum or vagina by using fingers or mechanical device

Stop use and ask doctor if

- rectal bleeding occurs
- condition worsens or does not improve within 7 days

Keep out of reach of children.

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

Directions

Adults:

- when practical, clean the affected area with mild soap and warm water and rinse thoroughly
- gently dry area by patting or blotting with toilet tissure or a soft cloth
- pump a small amount of foam onto clean toilet paper and apply externally to the affected area
- after application, flush toilet paper and wash hands

- medicated foam may be used up to 6 times daily or after each bowel movement
  Children under 12 years of age: ask a doctor

Other information

Store at a controlled room temperature: 15-30°C (59°-86°F)

Inactive ingredients

aloe barbadensis leaf juice, benzyl alcohol, citric acid, coco-glucoside, ethyl alcohol, glycerin, phenoxyethanol, potassium sorbate, propanediol, purified water, sodium citrate, tocopherol

Questions or comments?

1-844-308-8600

Package Labeling:

THE PHARMA C COMPANY MEDICATED FOAM WITH WITCH HAZEL

witch hazel liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55239-358
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
WITCH HAZEL (UNII: 101I4J0U34) (WITCH HAZEL - UNII:101I4J0U34)	WITCH HAZEL	500 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
COCO GLUCOSIDE (UNII: ICS790225B)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PROPANEDIOL (UNII: 5965N8W85T)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
TOCOPHEROL (UNII: R0ZB2556P8)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55239-358-00	210 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	05/15/2022	

Labeler - Kleen Test Products Corporation (168165814)

Revised: 11/2023 Kleen Test Products Corporation