TUCKS MEDICATED COOLING PADS- witch hazel solution
Blistex Inc

Active Ingredient

Witch hazel (50% w/w)

Purpose

Astringent

Uses

temporarily relieves the local itching and discomfort associated with hemorrhoids

aids in protecting irritated ahorectal 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 by using fingers or any mechanical device or applicator

Stop use and ask doctor if

rectal bleeding occurs

condition worsens or does not improve within 7 days

Keep out of the 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 by patting or blotting with toilet tissue or a soft cloth before applying

apply externally to the affected area up to 6 times daily or after each bowel movement

Children under 12 years of age: ask a doctor

Other information

DO NOT FLUSH-after application, discard pad in trash and wash hands

for use as a moist compress -- if necessary, first cleanse the area and place wipe in contact with irritated area for a soothing and cooling effect. Leave in place for up to 15 minutes and repeat as needed but not to exceed directions for use

Inactive ingredients

Water

Glycerin

Sodium Citrate, Unspecified form

Citric Acid Monohydrate

Ethylhexylglycerin

Phenoxyethanol

Edetate Disodium

Potassium Sorbate

label

TUCKS MEDICATED COOLING PADS

witch hazel solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10157-2103
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
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10157-2103-0	1 in 1 CARTON	08/30/2018	
1		100 in 1 JAR		
1		1 mL in 1 APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	08/30/2018	

Labeler - Blistex Inc (005126354)

Registrant - Accupac LLC (061595175)

Establishment

Name	Address	ID/FEI	Business Operations
Blistex Inc		005126354	manufacture(10157-2103)

Revised: 8/2024 Blistex Inc