GOOD SENSE MEDICATED- witch hazel cloth
Geiss, Destin & Dunn, Inc

Drug Facts

Active ingredient

Witch hazel 50%

Purpose

Astringent

Uses

For temporary relief of local discomfort, burning and irritation associated with hemorrhoids

Warnings

For external use only. Avoid contact with eyes.

Stop use and ask a doctor if

- condition worsens or does not improve within 7 days
- in case of bleeding

Do Not Use

Do not put this product into rectum using fingers or any mechanical device or applicator

Keep out of reach of children.

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

Directions

- When practical, cleanse the affected area with mild soap and warm water and rinse thoroughly
- Gently dry by patting or blotting with toilet tissue or soft cloth before each application of this product
- Gently apply to the affected area by patting and discard
- Apply as needed
- **For use on children under 12 years of age:** consult a doctor

Other information

Store at room temperature: 15 º C - 30 º C (59 - 86 º F)

Inactive ingredients

Alcohol, Aloe, Citric Acid, Decyl Glucoside, Glycerin, Phenoxyethanol, Potassium Sorbate, Propylene Glycol, Sodium Benzoate, Water.

Principal Display Panel

GOOD SENSE

Pre-moistened

N 50804-047-60

Medicated Pads

Hemorrhoidal/Vaginal Pads with Witch Hazel

- Instantly relieves hemorrhoidal burning and itching
- Witch hazel cools, soothes and protects

Compare to the active ingredient in Tucks®* Medicated Pads*

100% SATISFACTION GURANTEED

*This product is not manufactured or distributed by McNEIL-PPC, Inc., owner of the registered trademark Tucks®.

GOOD SENSE MEDICATED

witch hazel cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50804-047
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

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
ALOE (UNII: V5VD430YW9)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50804-047-60	100 in 1 JAR; Type 0: Not a Combination Product	12/16/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	12/16/2015	

Labeler - Geiss, Destin & Dunn, Inc (076059836)

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