HEMORRHOIDAL PADS- witch hazel patch
Meijer Distribution Inc.

Drug Facts

Active Ingredient

Witch Hazel 50%

Purpose

Hemorrhoidal Astringent

Uses

Temporarily relieves these external symptoms associated with hemorrhoids:

- itching
- burning
- irritation

Warnings

For external rectal use only.

When using this product

- do not exceed the recommended daily dosage unless directed by a doctor.
- do not insert into rectum or vagina using fingers or mechanical device.

Stop use and ask a 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 at 1-800-222-1222.

Directions

As hemorrhoidal treatment for adults:

- 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 applying.
- gently apply to the affected area by patting and then discard.
- can be used up to six times daily or after each bowel movement.

Children under 12 years of age

- consult a doctor.

Other Information

Store at controlled room temperature: 15º-30ºC (59º-86ºF)

Inactive Ingredients

Water, Sodium citrate, Citric acid, Glycerin, Phenoxyethanol, Potassium sorbate

Do not use this product if seal on jar is broken or missing.

* This product is not manufactured or distributed by Blistex, Inc., owner of the registered trademark TUCKS®.

Principal Display Panel

NDC 41250-868-10

meijer

Compare to Tucks

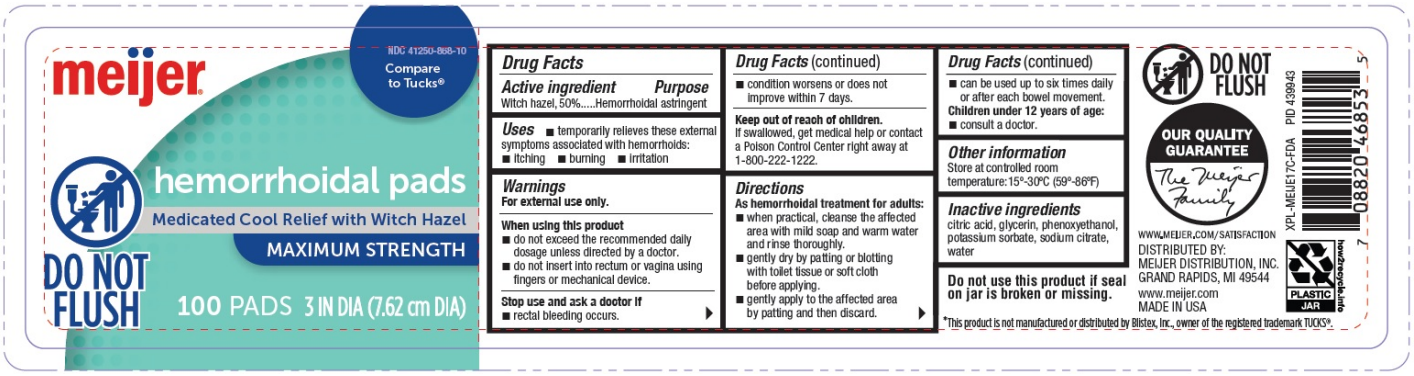
hemorrhoidal pads

Medicated Cool Relief with Witch Hazel

MAXIMUM STRENGTH

100 PADS 3 IN DIA (7.62 cm DIA)

DO NOT FLUSH



HEMORRHOIDAL PADS

witch hazel patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41250-868
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 1000 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-868-10	100 in 1 BOX	10/02/2019	
1		500 mg in 1 PATCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	06/28/2018	

Labeler - Meijer Distribution Inc. (006959555)

Registrant - Meijer Distribution Inc. (006959555)

Establishment

Name	Address	ID/FEI	Business Operations
U. S. Nonwovens Corp		080453184	manufacture(41250-868)

Revised: 1/2024 Meijer Distribution Inc.