

MEDICATED WIPES- 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: 15°-30°C (59°-86°F)

Inactive Ingredients

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

Principal Display Panel

NDC 41250-988-48

meijer

Compare to Preparation H® active ingredient*

medicated wipes

Hemorrhoidal Wipes with Witch Hazel

MAXIMUM STRENGTH

Provides Temporary Relief of Itching, Burning and Irritation

48 WIPES

6 IN x 5 IN (15cm x 12cm)

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NDC 41250-988-48

Compare to
Preparation H®
active ingredient*



medicated wipes

Hemorrhoidal Wipes with Witch Hazel

MAXIMUM STRENGTH

**DO NOT
FLUSH**

Provides Temporary Relief of
Itching, Burning and Irritation

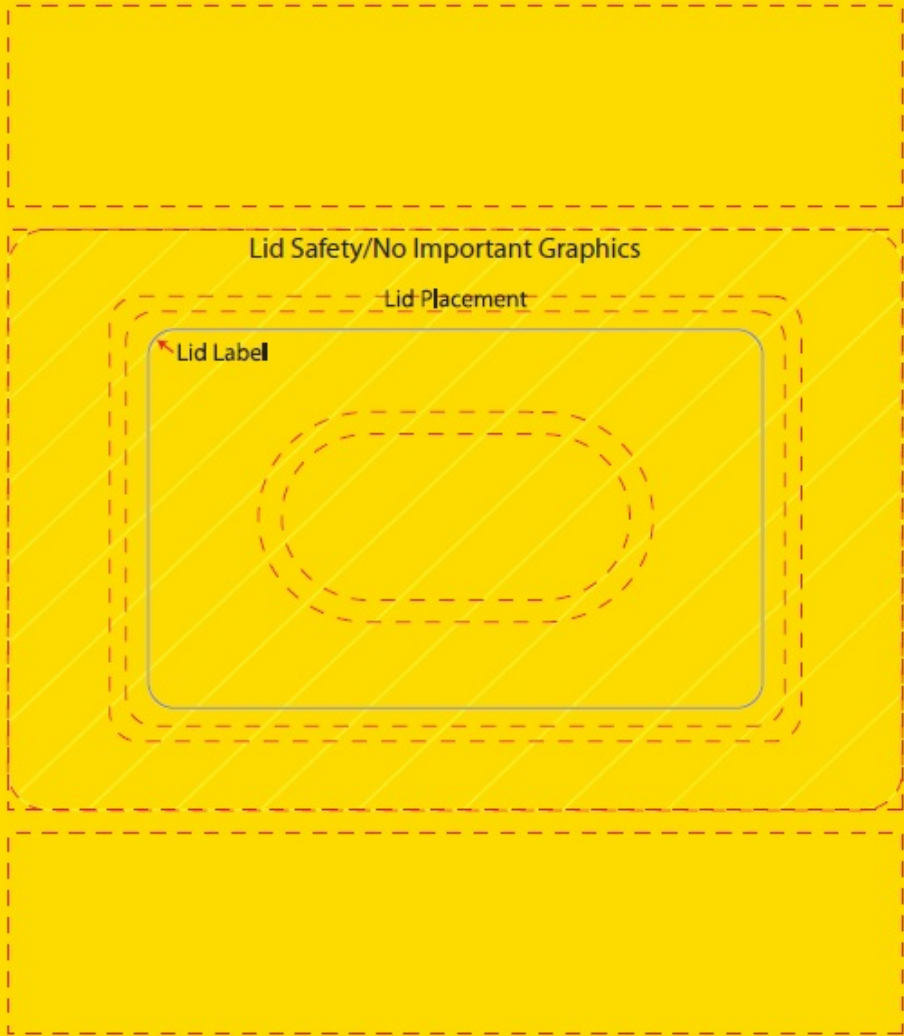
**6 IN x 5 IN
48 WIPES (15 cm x 12 cm)**

Drug Facts (continued)
Directions (continued)
 ■ 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.

Drug Facts (continued)
Other information
 Store at controlled room temperature: 15°-30°C (59°-86°F)
Inactive ingredients
 citric acid, glycerin, phenoxyethanol, potassium sorbate, sodium citrate, water
 DISTRIBUTED BY: MEIJER DISTRIBUTION, INC.
 GRAND RAPIDS, MI 49544



© This product is not manufactured or distributed by Pfizer Consumer Healthcare, owner of the registered trademark Preparation H®.



NOT SEPTIC SAFE.

Drug Facts
Active ingredient Purpose
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 ■ itching ■ burning ■ irritation
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Drug Facts (continued)
When using this product
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Stop use and ask a doctor if
 ■ rectal bleeding occurs.
 ■ condition worsens or does not improve within 7 days.

Drug Facts (continued)
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.

NDC 41250-988-96
 meijer
 Compare to Preparation H® active ingredient*
 medicated wipes
 Hemorrhoidal Wipes with Witch Hazel

MAXIMUM STRENGTH

Provides Temporary Relief of Itching, Burning and Irritation

For Use with Hemorrhoids and for Vaginal Care

Convenient, Easy to Use

96 WIPES (2 x 48 CT. PACKS)

6 IN x 5 IN (15.2 cm x 12.7 cm)



MEDICATED WIPES

witch hazel patch

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41250-988
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WITCH HAZEL (UNII: 10114J0U34) (WTCH HAZEL - UNII:10114J0U34)	WTCH 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: PDC6A3COOX)	
PHENOXYETHANOL (UNII: H1E492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41250-988-48	48 in 1 BOX	10/02/2019	
1		500 mg in 1 PATCH; Type 0: Not a Combination Product		
2	NDC:41250-988-96	96 in 1 BAG	10/02/2019	
2		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(41250-988)

Revised: 1/2024

Meijer Distribution Inc.