PRE-MOISTENED MEDICATED PADS, PRE-MOISTENED MEDICATED PADS HEMORRHOIDAL PADS WITH WITCH HAZEL, WALGREENS PRE-MOISTENED MEDICATED PADS, WALGREENS PRE-MOISTENED MEDICATED PADS HEMORRHOIDAL PADS WITH WITCH HAZEL- witch hazel patch
Walgreens Company

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 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

citric acid, glycerin, phenoxyethanol, potassium sorbate, sodium citrate, water

Principal Display Panel

NDC 0363-1250-10

Walgreens

Compare to the active ingredient in Tucks® ††

Pre-Moistened Medicated Pads

HEMORRHOIDAL PADS WITH WITCH HAZEL

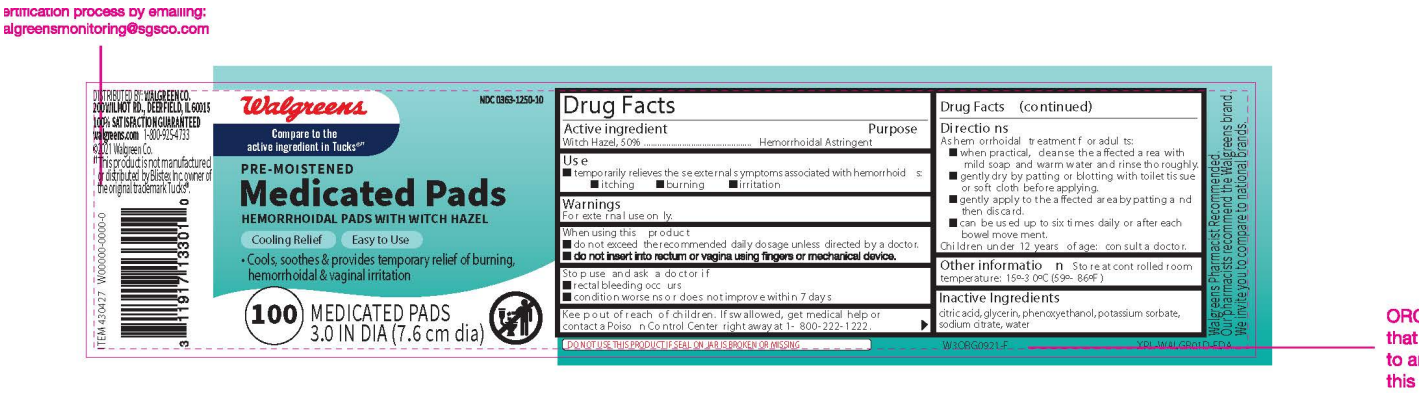
Cooling Relief

Easy to Use

Cools, soothes & provides temporary relief of burning, hemorrhoidal & vaginal irritation

100 Medicated Pads

3.0 IN DIA (7.6 cm dia)



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witch hazel patch

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-1250
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WITCH HAZEL (UNII: 101I4J0U34) (WITCH HAZEL - UNII:101I4J0U34)	WITCH HAZEL	50 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-1250-10	100 in 1 JAR	03/29/2009	
1		50 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	03/29/2009	

Labeler - Walgreens Company (008965063)

Revised: 1/2025 Walgreens Company