LEADER MEDICATED WIPE - witch hazel solution CARDINAL HEALTH

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active ingredient Purpose

Witch Hazel, 50% v/v......Astringent

Uses

For temporary relief of local discomfort itching, burning, and irritation associated with hemorrhoids For external use only. Avoid contact with eyes.

Consult a doctor

- if condition worsens or does not improve within 7 days
- in case of bleeding
- before exceeding the recommended dosage

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

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults and children 12 years and older, up to 6 times daily or after each bowel movement

Children under 12, consult a doctor

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

Other information

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

Inactive ingredients

aloe barbadensis gel, citric acid, diazolidinyl urea, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate

DISTRIBUTED BY CARDINAL HEALTH

DUBLIN, OHIO 43017

CIN 4246294

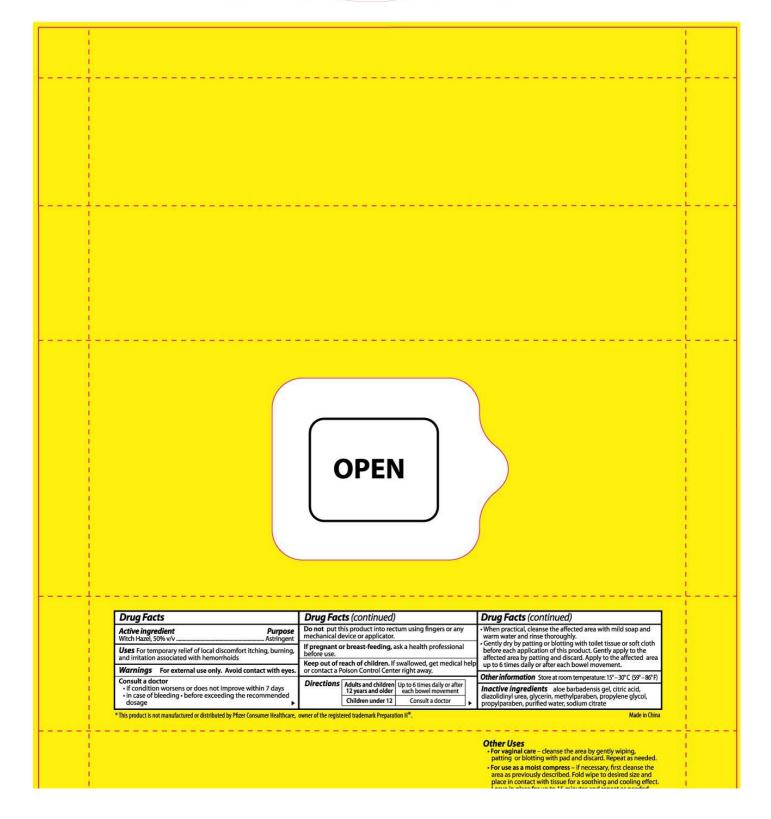
www.myleader.com

1-800-200-6313

Made in China







LEADER MEDICATED WIPE

witch hazel solution

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Product Type HUMAN OTC DRUG Item Code (Source) NDC:37205-741

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient NameBasis of StrengthStrengthWITCH HAZEL (UNII: 10 114J0 U34) (WITCH HAZEL - UNII: 10 114J0 U34)WITCH HAZEL5 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

 $\textbf{CITRIC ACID MO NO HYDRATE} \ (\text{UNII: } 2968 \, \text{PHW8 QP})$

DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)

GLYCERIN (UNII: PDC6 A3C0 OX)

METHYLPARABEN (UNII: A2I8 C7HI9 T)

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

PROPYLPARABEN (UNII: Z8IX2SC1OH)

SODIUM CITRATE (UNII: 1Q73Q2JULR)

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37205-741-67	48 in 1 POUCH		

Marketing Information

	ation Marketing Start Date Marketing End Date
OTC monograph final part346	03/01/2012

Labeler - CARDINAL HEALTH (097537435)

Revised: 3/2012 CARDINAL HEALTH