MEDICATED-PADS HEMORRHOIDAL PADS WITH WITCH HAZEL- witch hazel liquid Yinjing Medical Technology (Shanghai) Co., Ltd.

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.

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Active ingredient

Witch Hazel,50% v/v

Purpose

Astringent

Use

For temporary relief of local discomfort itching, burning, and irritation associated with hermorrhoids.

Warnings

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.

Keep out of reach of children.

Directions

? As a hemorrhoidal treatment for adults and children 12 years of age and older.

? 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 then discard. Apply to the affected area up to 6 times daily or after each bowel movement. Children under 12 years of age: consult a doctor.

Other information

Store at room temperature 15⁰ - 30⁰ C (59⁰ - 86⁰ F)

Inactive Ingredients:

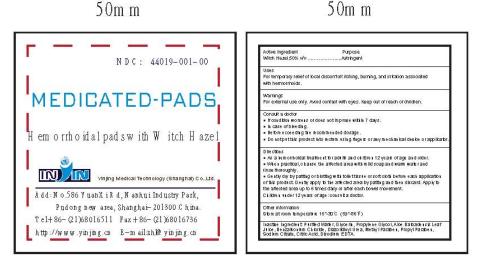
Purified Water, Glycerin, Propylene Glycol, Aloe Barbadensis Leaf Juice, Benzalkonium Chloride,

Diazolidinyl Urea, Methyl Paraben, Propyl Paraben, Sodium Citrate, Citric Acid, Disodium EDTA.

NDC: 44019-001-00

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MEDICATED-PADS Hemorrhoidal Pads with Witch Hazel 1 Wipe (44019-001-00)



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MEDICATED-PADS HEMORRHOIDAL PADS WITH WITCH HAZEL witch hazel liquid **Product Information Product Type** HUMAN OTC DRUG NDC:44019-001 Item Code (Source) **Route of Administration** TOPICAL **Active Ingredient/Active Moiety Basis of Strength Ingredient Name** Strength WITCH HAZEL (UNII: 10114J0U34) (WITCH HAZEL - UNII:10114J0U34) WITCH HAZEL 50 g in 100 g **Inactive Ingredients Ingredient Name** Strength WATER (UNII: 059QF0KO0R) GLYCERIN (UNII: PDC6A3C0OX) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) ALOE VERA LEAF (UNII: ZY81Z83H0X)

DENZALKUNIUM CH	LORIDE (UNII: F5UM2KM3W7)		
DIAZOLIDINYL UREA	(UNII: H5RIZ3MPW4)		
METHYLPARABEN (U	NII: A2I8C7HI9T)		
PROPYLPARABEN (U	NII: Z8IX2SC1OH)		
SODIUM CITRATE (U	NII: 1Q73Q2JULR)		
CITRIC ACID MONOR	HYDRATE (UNII: 2968PHW8QP)		
EDETATE DISODIUM	(UNII: 7FLD91C86K)		
Packaging			
Packaging # Item Code	Package Description	Marketing Start Date	Marketing End Date
00		Marketing Start Date 08/03/2016	Marketing End Date
# Item Code			Marketing End Date
# Item Code 1 NDC:44019-001-00	1 in 1 PACKET		Marketing End Date
# Item Code 1 NDC:44019-001-00	1 in 1 PACKET		Marketing End Date
Item Code NDC:440 19-00 1-00 1	1 in 1 PACKET 0.36 g in 1 PATCH; Type 0: Not a Combination Product		Marketing End Date
# Item Code 1 NDC:44019-001-00	1 in 1 PACKET 0.36 g in 1 PATCH; Type 0: Not a Combination Product		Marketing End Date
Item Code NDC:44019-001-00 1	1 in 1 PACKET 0.36 g in 1 PATCH; Type 0: Not a Combination Product Ormation		Marketing End Date Marketing End Date

Labeler - Yinjing Medical Technology (Shanghai) Co., Ltd. (530501535)

Registrant - Yinjing Medical Technology (Shanghai) Co., Ltd. (530501535)

Revised: 12/2017

Yinjing Medical Technology (Shanghai) Co., Ltd.