AER PRE-MOISTENED WITCH HAZEL PAD- witch hazel solution
Birchwood Laboratories LLC

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.

AER 40's Drug Facts

Active Ingredient

Witch Hazel 50%

Purpose

Astringent

Uses

Temporary relief of anal itching and burning

Temporarily protects irritated tissues

Warnings

For external rectal use only

Stop use and ask doctor if

rectal bleeding or continued irritation occurs

Keep Out Of Reach of Children

Keep out of reach of children

Directions

Gently apply to affected areas by patting and then discard

Inactive Ingredients

Benzalkonium Chloride, Citric Acid, Glycerin, Methylparaben, Purified Water, Sodium Citrate

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AER PRE-MOISTENED WITCH HAZEL PAD

witch hazel solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50289-3250
Route of Administration	TOPICAL, RECTAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
WITCH HAZEL (UNII: 101I4J0U34) (WITCH HAZEL - UNII:101I4J0U34)	WITCH HAZEL	500 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Product Characteristics

Color		**Score**	
Shape		**Size**	108mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50289-3250-1	40 in 1 JAR	06/28/2018	
1		2.8 mL in 1 APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

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

Labeler - Birchwood Laboratories LLC (096488432)

Revised: 1/2023

Birchwood Laboratories LLC