# RELIEF - witch hazel swab H and P Industries, Inc. dba Triad Group

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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### **DRUG FACTS**

### ACTIVE INGREDIENT

Witch Hazel 50%

### **PURPOSE**

Astringent

#### **USES**

Temporarily relieves these symptoms associated with hemorrhoids:

- itching
- burning
- irritation

### WARNINGS

For external use only.

### Stop use and ask 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

If swallowed, get medical help or contact a Poison Control Center right away.

### DIRECTIONS

- 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.

# **Inactive ingredients**

citric acid, diazolidinyl urea, glycerin, methylparaben, propylene glycol, propylparaben, sodium citrate, water

# **Package Information**

CAT. NO. 10-7540

NDC 50730-7502-1

### Relief

Pre-Moistened Hemorrhoidal Pads Comforts - Cools - Soothes

Compares to the leading brand

40 PADS

Triad Disposables, Inc. Brookfield, WI 53045 Made In USA



### **RELIEF** witch hazel swab **Product Information** Product Type HUMAN OTC DRUG NDC:50730-7502 Item Code (Source) **Route of Administration** RECTAL, TOPICAL **Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength witch hazel (UNII: 101I4J0U34) (witch hazel - UNII:101I4J0U34) witch hazel .5 mL

Inactive Ingredients			
Ingredient Name	Strength		
citric acid monohydrate (UNII: 2968PHW8QP)			
diazolidinylurea (UNII: H5RIZ3MPW4)			
glycerin (UNII: PDC6A3C0OX)			
methylparaben (UNII: A218 C7H19 T)			
propylene glycol (UNII: 6DC9Q167V3)			
propylparaben (UNII: Z8IX2SC1OH)			
sodium citrate (UNII: 1Q73Q2JULR)			
water (UNII: 059QF0KO0R)			

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:50730-7502-1	40 in 1 JAR					

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph final	part346	0 1/14/19 9 7				

# Labeler - H and P Industries, Inc. dba Triad Group (050259597)

Registrant - Hand P Industries, Inc. dba Triad Group (050259597)

Establishment					
Name	Address	ID/FEI	<b>Business Operations</b>		
Hand P Industries, Inc. / Triad Group		050259597	manufacture		

Revised: 11/2009 Hand P Industries, Inc. dba Triad Group