ALCOHOL WIPE- isopropyl alcohol swab ALCOHOL PREP PAD- isopropyl alcohol swab Honeywell Safety Products USA, Inc

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

0498-0142 & 0498-0143: Alcohol Wipes

Active ingredient

Isopropyl alcohol 70%

Purpose

First aid antiseptic

Uses

first aid to help prevent infection in minor cuts, scrapes, and burns

Warnings

For external use only

Flammable, keep away from fire and flame

Do not use

- in or near eyes
- over large areas of the body

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

When using this product

• do not use longer than 1 week unless directed by a doctor

Stop use and consult a doctor if

• condition persists or gets worse

Keep out of the reach of children

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

Directions

- clean the affected area
- may be covered with a sterile bandage
- apply wipe to affeted are 1 to 3 times daily
- discard wipe after single use

Other information

- store at room temperature 15 ° to 25 ° C (59 ° to 77 °F)
- do not use if packet is torn or opened

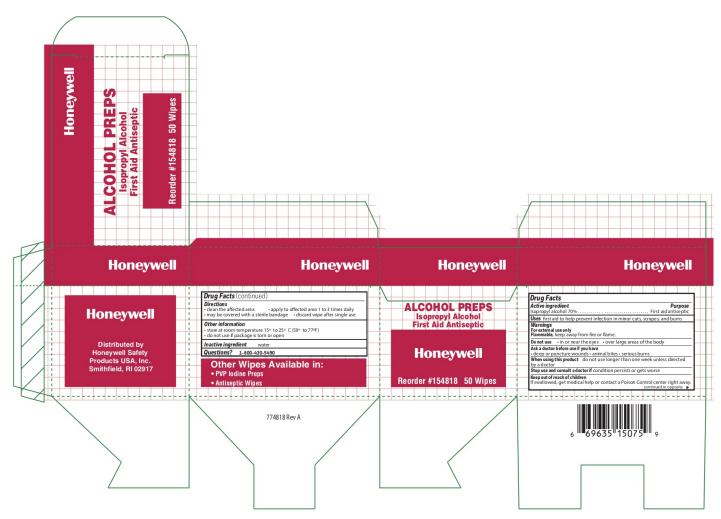
Inactive ingredient

water

Questions

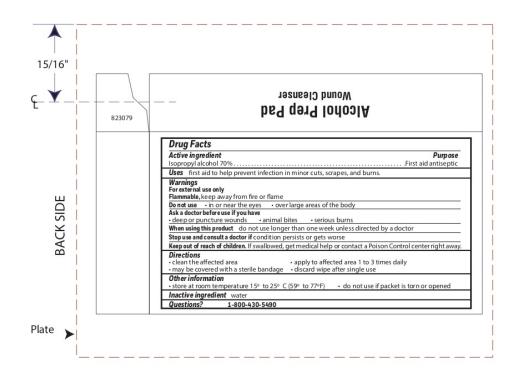
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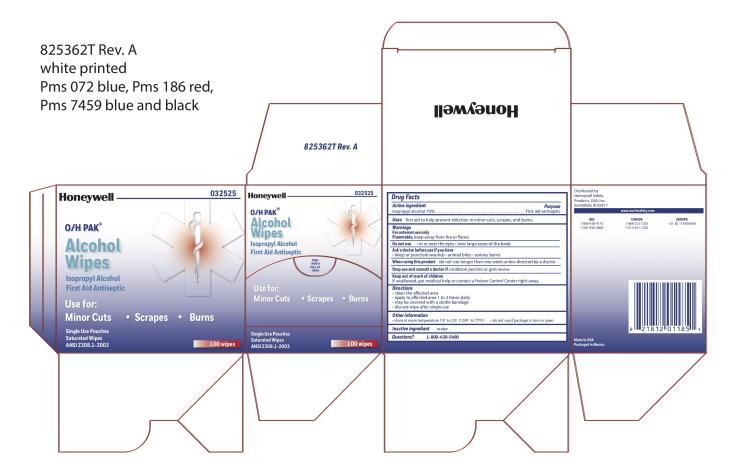
alcohol wipes





796390 Rev. B Unit Carton Printing Plate for "B" size carton.





ALCOHOL WIPE							
isopropyl alcohol swab							
Product Information							
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:0498-0143			
Route of Administration	TOPICAL						
Active Ingredient/Active Moiety							
Ingredient Name			Basis o Streng		Strength		
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)			ISOPROPYL ALCOHOL		0.7 mL in 1 mL		
Inactive Ingredients							
Ingredient Name			Strength				
WATER (UNII: 059QF0K00R)							
Packaging							

#	ltem Code	Pa	ckage Description		Marketing Start Date	Marketing End Date
1	NDC:0498-0143- 04	0.4 mL in 1 PO Product	OUCH; Type 0: Not a Combination		09/18/2018	
2	NDC:0498-0143- 34	10 in 1 BOX		(09/18/2018	
2		0.4 mL in 1 PO Product	UCH; Type 0: Not a Combir	nation		
3	NDC:0498-0143- 05	50 in 1 BOX		(09/18/2018	
3		0.4 mL in 1 PO Product	UCH; Type 0: Not a Combir	nation		
4	NDC:0498-0143- 10	100 in 1 BOX		(09/18/2018	
4		0.4 mL in 1 PO Product	UCH; Type 0: Not a Combir	nation		
5	NDC:0498-0143- 33	3000 in 1 BOX		(09/18/2018	
5			UCH; Type 0: Not a Combir	nation		
5		Product				
	arkating		ion			
	arketing	Informat		Iraph	Marketing Start	Marketing End
M	Marketing Category	Informat	ion tion Number or Monog Citation	ıraph	Marketing Start Date	Marketing End Date
M	Marketing	Informat	tion Number or Monog	ıraph	-	-
M	Marketing Category approved drug	Informat	tion Number or Monog	ıraph	Date	-
M un oth	Marketing Category approved drug	Informat Applicat	tion Number or Monog	ıraph	Date	-
M un otř	Marketing Category approved drug	Informat Applicat	tion Number or Monog	Jraph	Date	-
un otř Al	Marketing Category approved drug her	Informat Applicat	tion Number or Monog	Jraph	Date	-
un otř Al	Marketing Category approved drug her LCOHOL P propyl alcohol	Informat Applicat	tion Number or Monog		Date	-

Active Ingredient/Active Moiety						
Ingredient Name		Basis o Streng		strength		
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)		ISOPROPYL ALCOHOL	.	′mL 1mL		
Inactive Ingredients						
Ingredient Name			Strength			
WATER (UNII: 059QF0KO0R)						
Packaging						
	Maulsat		Maulzat	ing Find		

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:0498-0142- 34	10 in 1 BOX	01/01/2017					
1		0.4 mL in 1 POUCH; Type 0: Not a Combination Product						
Μ	Marketing Information							
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
	approved drug ner		01/01/2017					

Labeler - Honeywell Safety Products USA, Inc (118768815)

Registrant - Honeywell Safety Products USA, Inc (118768815)

Revised: 1/2024

Honeywell Safety Products USA, Inc