ANTISEPTIQUE 70- alcohol gel Hubot Healthcare 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.

Antis eptique +70

Drug Facts

Active ingredient

Ethyl Alcohol 70% v/v

Purposes

Antiseptic

Uses

to help reduce bacteria on skin

Warnings

|For external use only | Flammable, keep away from fire or flame, heat, sparks, and sources of static discharge

Do not use

 $|In\ eyes\ |\ On\ children\ less\ than\ 2\ years\ of\ age\ |\ On\ open\ skin\ wounds$

When using this product

|If in eyes, rinse promptly and thoroughly with water | Discontinue use if irritation and redness develops *Stop use and ask a doctor if* skin irritation or redness occurs or persists for more than 72 hours.

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

Directions

|Place enough product on hands to cover all surfaces. Rub hands together until dry | Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

| Store between 15-30C (59-86F). Avoid freezing and excessive heat above 40C (104F) | For additional information, see Safety Data Sheets (SDS)

Inactive Ingredients

water (aqua), propylene glycol, dimethicone PEG-7 Isostearate, acrylates/C10-30 Alkyl Acrylate Crosspolymer, aminomethylpropanediol, tocopherol acetate, aloe barbadensis leaf juice

Questions?

sales@tri-pac.us

PRINCIPAL DISPLAY PANEL - 946 ML Bottle Label

ANTISEPTIQUE⁺70 gel sanitizer antiseptic | rinse-free | 70% alcohol FOR HANDS dye free, fragrance free FOR HEALTHCARE PROFESSIONALS ONLY 32 FL. OZ. (946 ML)

ANTISEPTIQUE[†]70 gel sanitizer

antiseptic | rinse-free | 70% alcohol

FOR HANDS

dye free, fragrance free







Purposes

Antiseptic

FOR HEALTHCARE PROFESSIONALS ONLY

32 FL. OZ. (946 ML)

N 72138-AKR-22

Drug Facts:

Active Ingredient
Ethyl Alcohol 70% v/v.....

Use: to help reduce bacteria on the skin.

Warnings • For external use only • Flammable, keep away from fire or flame, heat, sparks and sources of static discharge.

Do not use • In eyes • On children less than 2 years of age • On open skin wounds.

When using this product: if in eyes, rinse promptly and thoroughly with water I discontinue use if irritation and redness develops

Stop use and ask a doctor if skin irritation or redness occurs, this may be signs of a serious condition.

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

Directions • Place enough product on hands to cover all surfaces. Rub hands together until dry.• Supervise children under 6 years of age when using this product to avoid swallowing.

Other information • Store between 15-30°C (59-86°F), Avoid freezing and excessive heat above 40°C (104°F) • For additional information, see Safety Data Sheets (SDS)

Inactive ingredients: Water (Aqua), Propylene Glycol, Dimethicone Peg-7 Isostearate, Acrylates/ C10-30 Alkyl Acrylate Crosspolymer, Aminomethylpropanedial, Tocopherol Acetate, Aloe Barbadensis Leaf Juice.

Questions? E-mail: sales@tri-pac.us

TRI-PAC, INC. I WWW.TRI-PAC.US HUBOT HEALTHCARE I WWW.HUBOT.HEALTH 3333 N. KENMORE ST. SOUTH BEND, IN 46628 USA









ANTISEPTIQUE 70

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72138-460

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
Water (UNII: 059QF0KO0R)		
Propylene Glycol (UNII: 6DC9Q167V3)		
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)		
DIMETHICO NE PEG-8 ISO STEARATE (UNII: JGV7463UFN)		
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)		
Tocopherol (UNII: R0ZB2556P8)		
Aloe Vera Leaf (UNII: ZY81Z83H0X)		

F	nckaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:72138-460- 06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/25/2020		
2	NDC:72138-460- 08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/25/2020		

3	NDC:72138-460- 16	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/25/2020	
4	NDC:72138-460- 32	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/25/2020	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC MONOGRAPH NOT FINAL	part333E	06/25/2020			

Labeler - Hubot Healthcare LLC (081084880)

Revised: 6/2020 Hubot Healthcare LLC