

ANTISEPTIQUE PLUS- 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.

Antiseptique Plus

Drug Facts

Active ingredient

Ethyl Alcohol 62.5% w/w

Purposes

Antiseptic handwash

Uses

for handwashing to decrease 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

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

- Apply product onto hands, spread thoroughly and rub dry

Other Information

- For additional information, see Safety Data Sheets (SDS)
- For emergency medical information in USA and Canada, call 1-888-255-3924
- For emergency medical information worldwide, call 1-813-248-0573

Inactive Ingredients

water (aqua), ethyl acrylate/ methacrylic acid/ steareth-20 methacrylate copolymer, aminomethylpropanediol, ppg-14 butyl ether, tocopherol acetate, aloe barbadensis leaf juice

Questions?

Call 1-855-CLEAN-55 (253-2655)

PRINCIPAL DISPLAY PANEL - 473 ML Bottle Label

ANTISEPTIQUE⁺
PLUS
gel sanitizer

antiseptic | rinse-free | 62.5% alcohol

FOR HANDS
dye free, fragrance free

16 FL. OZ. (473 ML)


ANTISEPTIQUE⁺ PLUS

gel sanitizer

antiseptic | rinse-free | 62.5% alcohol

FOR HANDS

dye free, fragrance free



16 FL. OZ. (473 ML)

NDC 72138-473-16

DRUG FACTS:	
ACTIVE INGREDIENT	PURPOSES
Ethyl Alcohol 62.5% v/v	Antiseptic handwash
USES: For handwashing to decrease 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 In children less than 2 months 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 irritation or rash occurs. These 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: For additional information, see Safety Data Sheets (SDS) For emergency medical information in USA and Canada, call 1-888-255-3924 For emergency medical information worldwide, call +1-813-248-0573 Store between 15-30C (59-86F) Avoid freezing and excessive heat above 40C (104F)	
INACTIVE INGREDIENTS: Aqua (Water), Acrylates/styrene-20 Methacrylate Crosspolymer, Poly-4 Butyl Ether, Aminomethylpropanol, Aloe Barbadensis Leaf Juice, Tocopherol Acetate	



SDS#H20015

QUESTIONS? E-MAIL: SALES@TRI-PAC.US

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3333 N. KENMORE ST. SOUTH BEND, IN 46628 USA

ANTISEPTIQUE PLUS

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72138-470
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62.5 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
ETHYL ACRYLATE/METHACRYLIC ACID/STEARETH-20 METHACRYLATE COPOLYMER (UNII: EPA1872R1N)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
PPG-14 BUTYL ETHER (UNII: R199TJT95T)	
Aloe Vera Leaf (UNII: ZY81Z83H0X)	
Tocopherol (UNII: R0ZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72138-470-06	177 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/13/2020	
2	NDC:72138-470-16	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/13/2020	
3	NDC:72138-470-32	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/13/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	03/13/2020	

Labeler - Hubot Healthcare LLC (081084880)

Revised: 3/2020

Hubot Healthcare LLC