ANTISEPTIQUE- 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

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

Questions?

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

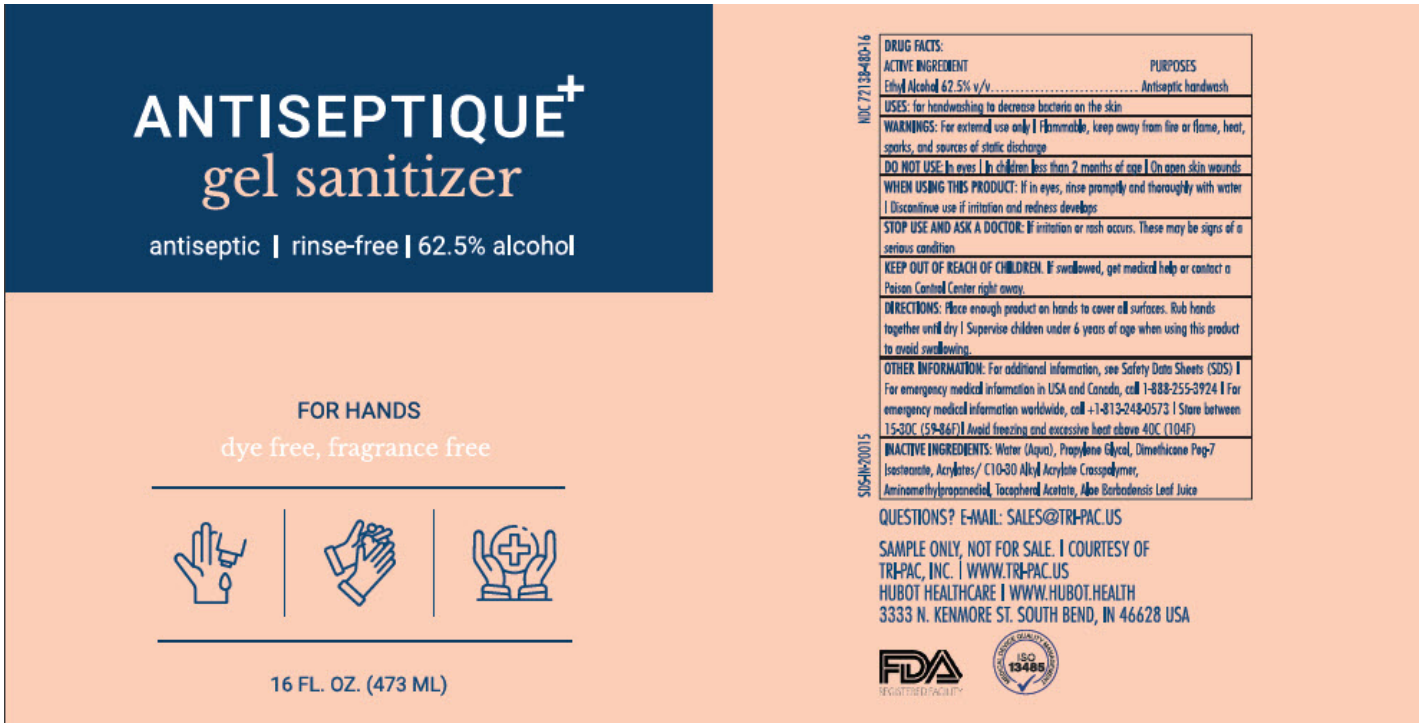
PRINCIPAL DISPLAY PANEL - 473 ML Bottle Label

ANTISEPTIQUE+
gel sanitizer

antiseptic | rinse-free | 62.5% alcohol

FOR HANDS
dye free, fragrance free

16 FL. OZ. (473 ML)



ANTISEPTIQUE

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72138-480
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)	
Propylene Glycol (UNII: 6DC9Q167V3)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
Dimethicone PEG-7 Isostearate (UNII: JVS3399FNW)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
Tocopherol (UNII: R0ZB2556P8)	
Aloe Vera Leaf (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72138-480-06	177 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/12/2020	
2	NDC:72138-480-16	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/12/2020	
3	NDC:72138-480-32	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/12/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/12/2020	

Labeler - Hubot Healthcare LLC (081084880)

Revised: 3/2020 Hubot Healthcare LLC