

**ANTISEPTIQUE ADVANCE STRENGTH- alcohol spray**  
**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.*

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**Antiseptique Advance Strength**

**Drug Facts**

**Active ingredient**

Ethyl Alcohol 80% v/v

**Purposes**

Antiseptic handwash

**Uses**

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

**Warnings**

- **For external use only**
- **Flammable**, keep away from heat or flame.

**Do not use**

- In eyes | on children less than 2 months old | on open skin wounds

**When using this product**

- Keep out of eyes, ears and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

**Stop use and ask a doctor if** skin irritation or rash occurs. There may be signs of a serious conditions.

**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)

**Inactive Ingredients**

Purified Water USP, Glycerin, Hydrogen Peroxide

**Questions?**

Email: sales@tri-pac.us

**PRINCIPAL DISPLAY PANEL - 59 ML Bottle Label**

ANTISEPTIQUE<sup>+</sup>  
 ADVANCED STRENGTH  
 sanitizer mist

Alcohol Antiseptic 80%  
 Topical Non-sterile Solution  
 2 FL. OZ. (59 ML)

**ANTISEPTIQUE<sup>+</sup>**  
**ADVANCED STRENGTH**  
 sanitizer mist

**Alcohol Antiseptic 80%**  
**Topical Non-sterile Solution**  
 2 FL. OZ. (59 ML)

NDC 72138-420-20

<b>DRUG FACTS:</b>	
<b>ACTIVE INGREDIENT</b> Alcohol 80% v/v .....	<b>PURPOSES</b> Antiseptic handwash
<b>USES:</b> For handwashing to decrease bacteria on the skin	
<b>WARNINGS:</b> For external use only. Flammable, keep away from fire or flame, heat, sparks and sources of static discharge	
<b>DO NOT USE:</b> In eyes   In children less than 2 months of age   On open skin wounds	
<b>WHEN USING THIS PRODUCT:</b> If in eyes, rinse promptly and thoroughly with water   Discontinue use if irritation and redness develops	
<b>STOP USE AND ASK A DOCTOR:</b> If irritation or rash occurs. These may be signs of a serious condition.	
<b>KEEP OUT OF REACH OF CHILDREN.</b> If swallowed, get medical help or contact a Poison Control Center right away.	
<b>DIRECTIONS:</b> Apply product onto hands, spread thoroughly and rub dry   Supervise children under 6 years of age when using this product to avoid swallowing.	
<b>OTHER INFORMATION:</b> 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)	
<b>INACTIVE INGREDIENTS:</b> Purified Water USP, Glycerin, Hydrogen Peroxide	

5105-NV-20015

QUESTIONS? E-MAIL: SALES@TRIPAC.US  
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 TRI-PAC, INC. | WWW.TRI-PAC.US  
 HUBOT HEALTHCARE | WWW.HUBOT.HEALTH  
 3333 N. KENMORE ST. SOUTH BEND, IN 46628 USA



**ANTISEPTIQUE ADVANCE STRENGTH**

alcohol spray

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:72138-420
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	
Hydrogen Peroxide (UNII: BBX060AN9V)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72138-420-20	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/20/2020	
2	NDC:72138-420-40	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/20/2020	
3	NDC:72138-420-60	177 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/20/2020	
4	NDC:72138-420-16	472 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/20/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	03/19/2020	

**Labeler** - Hubot Healthcare LLC (081084880)

Revised: 3/2020

Hubot Healthcare LLC