ANTIBACTERIAL HAND SANITIZER- alcohol spray
TEKWELD SOLUTIONS, INC.

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.

Antibacterial Hand Sanitizer

Drug Facts

Active Ingredients

Ethyl Alcohol 62%

Purpose

Antibacterial

Uses

For hand washing to decrease bacteria on the skin.

Warnings Flammable.

Keep away from fire or flame. For external use only.

When using this product

Keep out of eyes. In case of contact with eyes, rinse with water.

Stop use and ask doctor if

irritation and redness develops and persists.

Keep out of reach of children.

If swallowed, get medical help promptly and contact Poison Control.

Directions

Wet hands throughly with product and allow to dry without wiping.

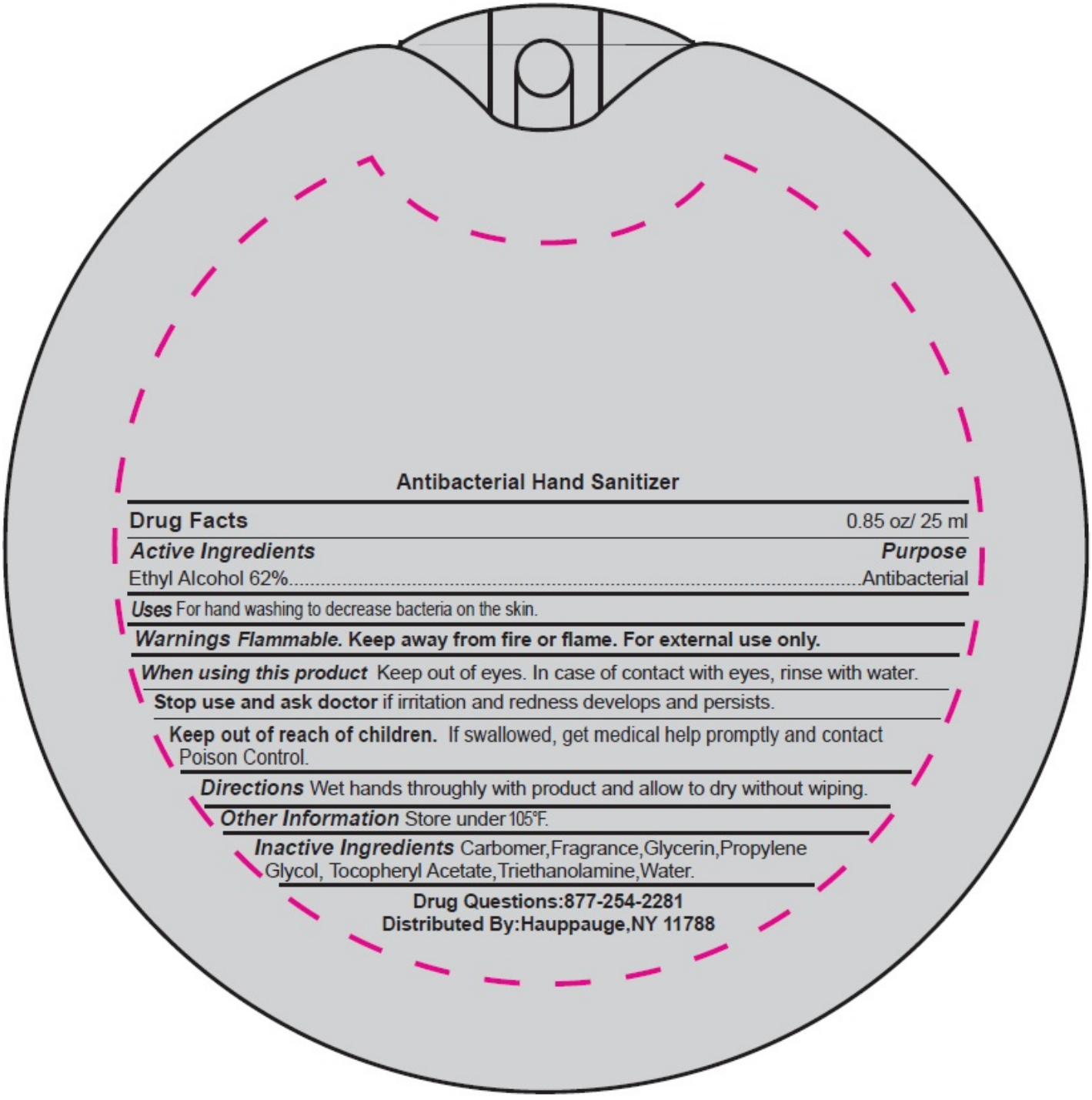
Other Information

Store under 105°F.

Inactive Ingredients

Carbomer,Fragrance,Glycerin,Propylene Glycol, Tocopheryl Acetate,Triethanolamine,Water.

Package Labeling:



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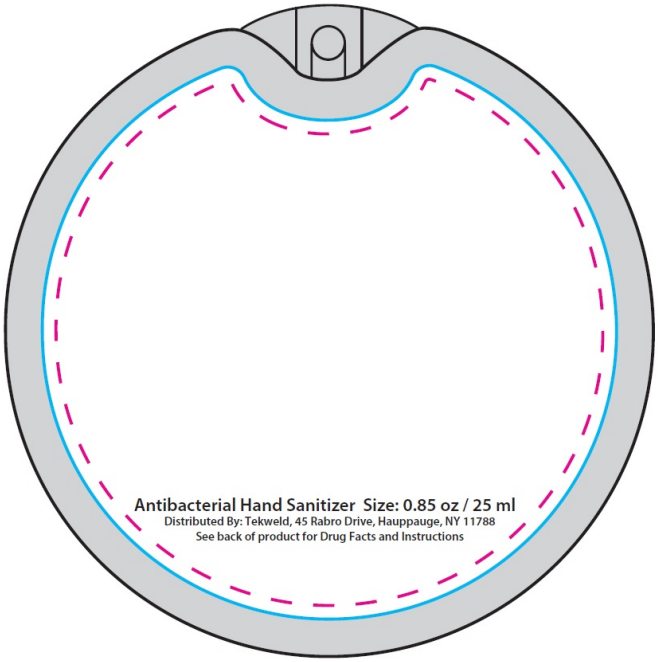
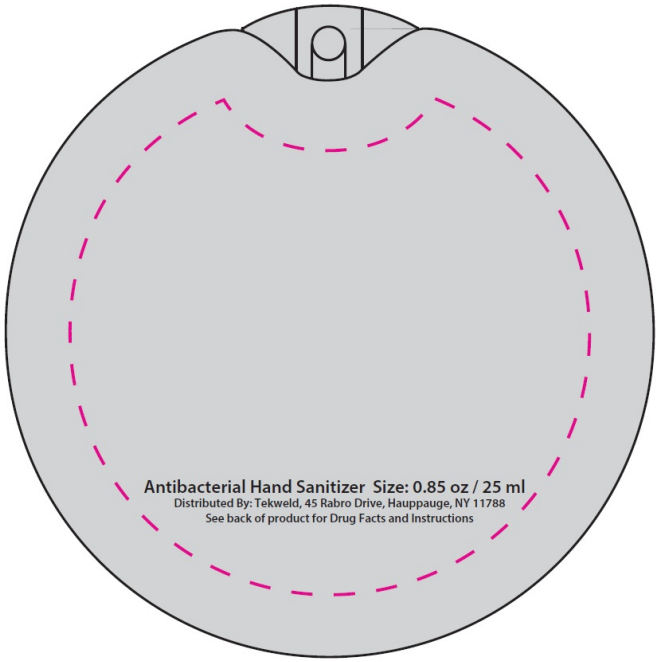
Drug Questions:877-254-2281

Distributed By:Hauppauge,NY 11788





ANTIBACTERIAL HAND SANITIZER

alcohol spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71160-189
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	620 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71160-189-10	10 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/09/2017	
2	NDC:71160-189-05	5 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/09/2017	
3	NDC:71160-189-25	25 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/09/2017	
4	NDC:71160-189-20	20 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/09/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/09/2017	

Labeler - TEKWELD SOLUTIONS, INC. (029077754)

Revised: 11/2018 TEKWELD SOLUTIONS, INC.