SENTINEL HAND SANITIZER- alcohol gel
DANLAB DANUTA KATRYNSKA

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.

Active Ingredient

Ethyl Alcohol 75% v/v.

Purpose

Antiseptic

Use

- To decrease bacteria on the skin that could cause disease.
- Recommended for repeated use.

Warnings

For external use only:hands.

Flammable. Keep away from heat or flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- Avoid contact with broken skin.
- Do not inhale or ingest.
- **Stop use and ask a doctor if**
- irritation or redness develops.
- Condition 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

- Wet hands thoroughly with product and allow to dry without wiping.
- For children under 6, use only under adult supervision.
- Not recommended for infants.

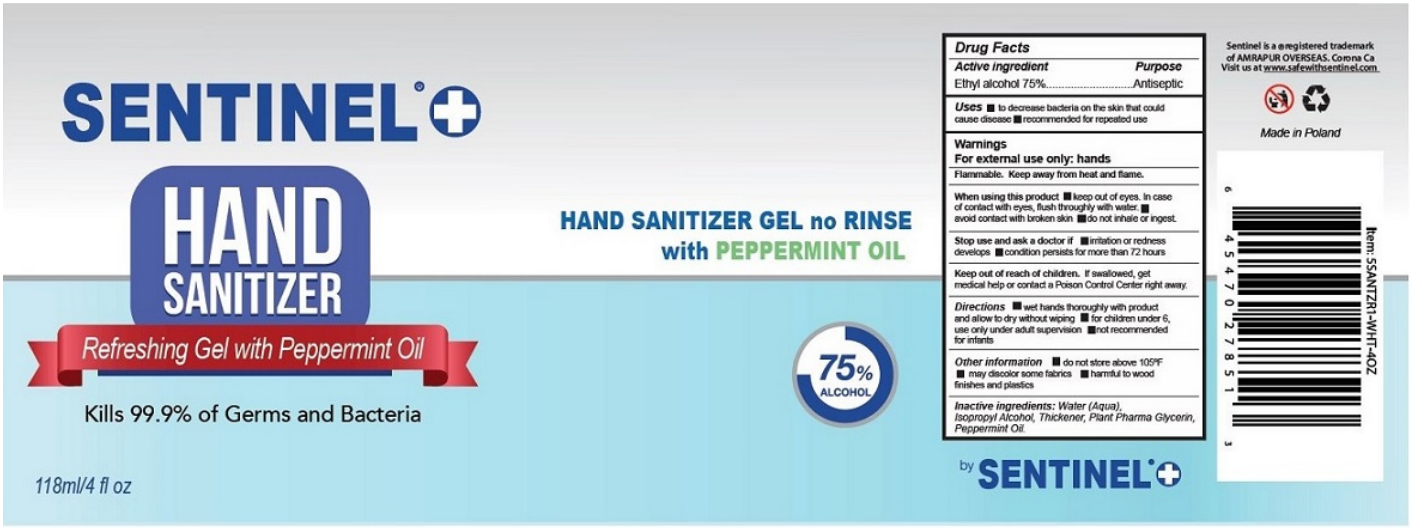
Other information

- Do not store above 105^0F
- May discolor certain fabrics.
- Harmful to wood finishes and plastics

Inactive ingredients

Water (Aqua), Isopropyl alcohol, Thickener, Plant pharma glycerin, Peppermint oil.

Package Label - Principal Display Panel



SENTINEL HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79041-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
GLYCERIN (UNII: PDC6A3C0OX)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
POLYACRYLIC ACID (8000 MW) (UNII: 73861X4K5F)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:79041-001-01	59 mL in 1 TUBE; Type 0: Not a Combination Product	06/18/2020	
2	NDC:79041-001-02	75 mL in 1 TUBE; Type 0: Not a Combination Product	06/18/2020	
3	NDC:79041-001-03	118 mL in 1 TUBE; Type 0: Not a Combination Product	06/18/2020	
4	NDC:79041-001-04	125 mL in 1 TUBE; Type 0: Not a Combination Product	06/18/2020	
5	NDC:79041-001-05	236 mL in 1 TUBE; Type 0: Not a Combination Product	06/18/2020	
6	NDC:79041-001-06	250 mL in 1 TUBE; Type 0: Not a Combination Product	06/18/2020	
7	NDC:79041-001-07	500 mL in 1 TUBE; Type 0: Not a Combination Product	06/18/2020	
8	NDC:79041-001-08	1000 mL in 1 TUBE; Type 0: Not a Combination Product	06/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/18/2020	

Labeler - DANLAB DANUTA KATRYNSKA (422359029)

Revised: 6/2020 DANLAB DANUTA KATRYNSKA