HAND SANITIZER- ethyl alcohol gel
Ultra Distributors 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.

HAND SANITIZER Cars

Active ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only: hands

Flammable. Keep away from fire 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(1-800-222-1222)right away.

Directions

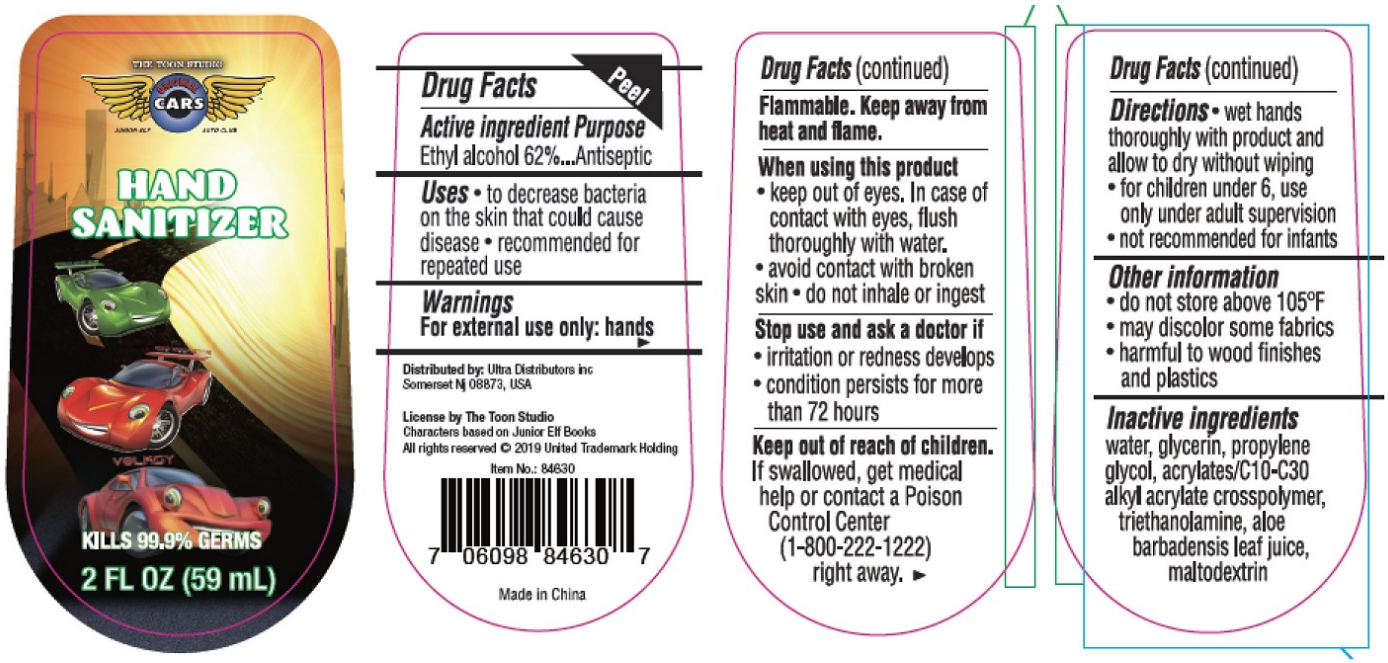
- wet hands thoroughly with product and allow to dry without wiping.
- for children under 6 , use only under adult supervision.
- not recommended for infants .

Other information

- do not store above 105°F
- may discolor some fabics
- harmful to wood finished and plastics

Inactive ingredients

water, glycerin, propylene glycol, acrylates/C10-C30 alkyl acrylate crosspolymer, triethanolamine, aloe barbadensis leaf juice, maltodextrin



HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78495-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:78495-002-01	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	09/04/2020	

Labeler - Ultra Distributors Inc (007160073)

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