

**TROLLS HAND SANITIZER- ethyl alcohol gel**  
**Ashtel Studios, 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.*

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**Smart Care® Hand Sanitizer**

***Drug Facts***

**Active Ingredient**

Ethyl Alcohol 62%

**Purpose**

Antiseptic

**Use**

To help reduce bacteria and germs on the skin.

**WARNING**

Flammable. Keep away from fire or flame. For external use only • Stop use and ask a doctor if irritation or redness develops and persists.

• **Keep out of reach of children.** • In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

**Directions**

▣ Place enough product in palm to cover hands and rub hands together briskly until dry. • Children under 6, use only under adult supervision. • Not recommended for infants.

**Other Information**

• Do not store above 100°F (38°C). • May discolor some fabrics. • Harmful to wood finishes and plastics.

**Inactive Ingredients**

• Deionized Water, Glycerine, Carbomer, Vitamin E, Fragrance, Aloe Barbadensis Gel, Triethanolamine, Propylene Glycol.

**Smart Care®**

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QUESTIONS OR COMMENTS?

1-877-274-8358 Toll Free in USA • 1-909-434-0911 International

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Packaging



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## TROLLS HAND SANITIZER

ethyl alcohol gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70 108-066
<b>Route of Administration</b>	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
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TROLAMINE (UNII: 9O3K93S3TK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70 108-066-01	53 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/29/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/29/2020	

