DR.S CLEAN HAND SANITIZER- ethyl alcohol gel
Erf Exhibit 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.

Dr.s CLEAN HAND SANITIZER

Drug Facts

Active ingredients

Ethyl Alcohol 70.0%

Purpose

ANTISEPTIC

Uses

for hand-washing to decrease bacteria on the skin. only when water is not available

Warnings

Flammable. Keep away from fire and flames. For external use only.

When using this product • Do not get into eyes.

• If contact occurs, rinse eyes thoroughly with water.

Stop use and ask a doctor ifirritation or redness develops.

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.

Inactive Ingredients

Purified Water, Aloe Extract, Glycerin, Sodium Hyaluronate, Carbomer, Butylene Glycol, Triethanolamine, Flavoring

Questions?

310.537.7662 or visit http://www.megaexhibit.com

HAND GEL

Simply KILL 99.9% Germ

Leaves Hands Feeling Soft

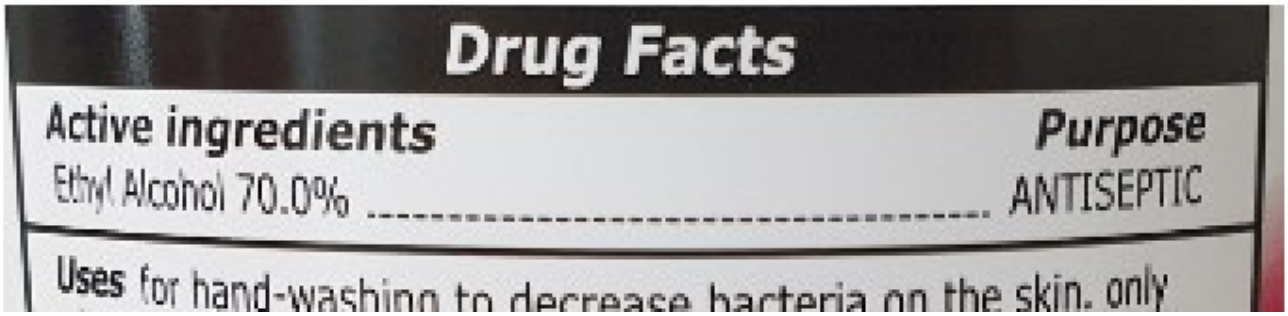
ADVANCED

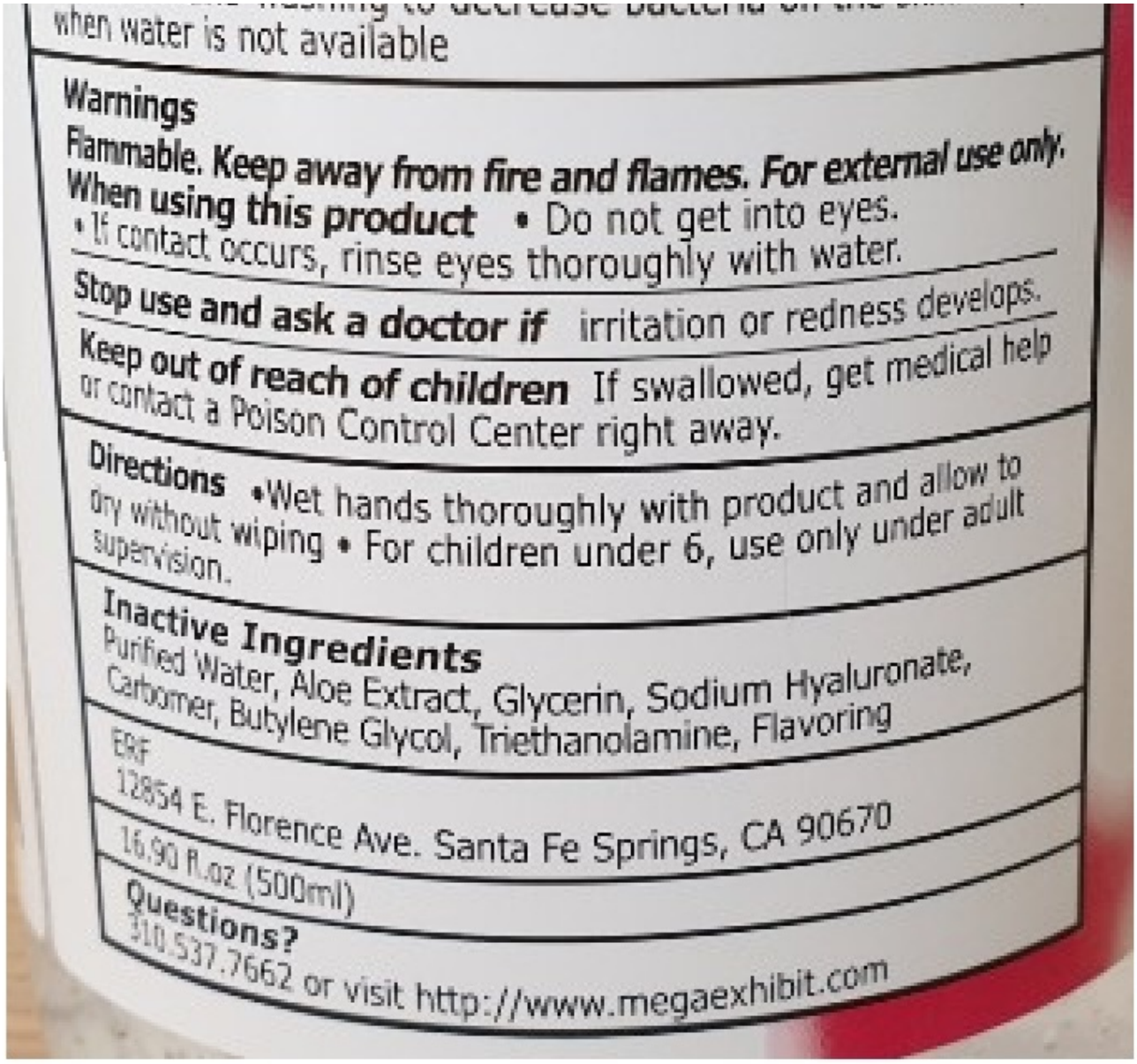
(ethanol) contented

ERF

12854 E. Florence Ave. Santa Fe Springs, CA 90670

Packaging





DR.S CLEAN HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76724-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALOE (UNII: V5VD430YW9)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:76724-001-05	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/04/2020	

Labeler - Erf Exhibit Inc (099461452)

Revised: 5/2020 Erf Exhibit Inc