HANNA HAND SANITIZER LIQUID- alcohol liquid
Rainbow Beauty Cosmetic Co., Ltd.

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.

Drug Facts

alcohol

Water, Glycerin, Sodium Hyaluronate, Citrus Limon (Lemon) Fruit Extract, Fragrance, CI 15985, CI 16255

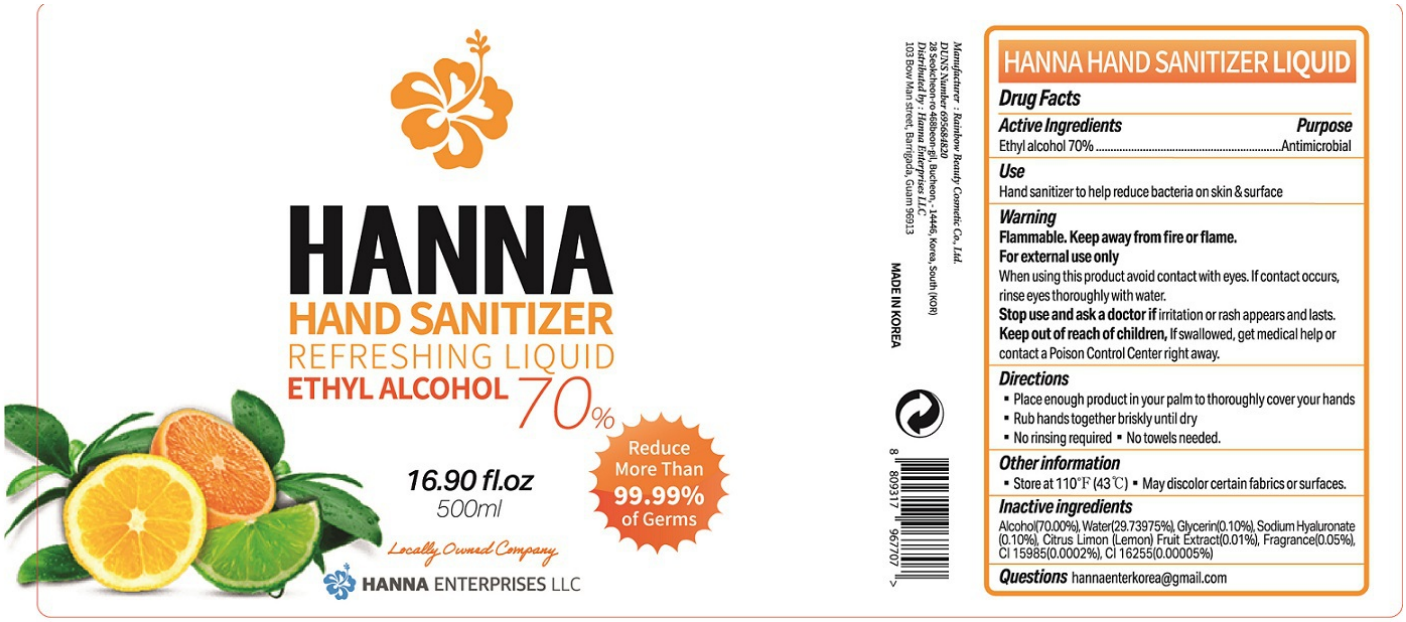
Sterilization of hands and skin

KEEP OUT OF REACH OF THE CHILDREN

Apply an appropriate amount on your hands and rub well to dry.

1. Do not use on the following body parts. A wide range of body parts and damaged skin around the eyes and
ears, in the oral cavity (may have irritating effects)
2. If the following symptoms appear, stop using them immediately and consult a doctor or pharmacist.
1) Hypersensitivity symptoms such as rash, erythema, itching, and edema
2) Skin irritation symptoms
3. Other precautions
1) For external use only (do not underwear).
2) Be careful not to get into your eyes, and if so, rinse well with clean water and consult a doctor or pharmacist.
3) Be careful not to inhale the vapor when using it extensively or for a long period of time (irritation to the
mucous membranes, headaches, etc. may occur if ethanol vapor is consumed in large quantities or repeatedly).
4) If repeated use on the same site, be careful as the skin may become rough due to degreasing.
5) Do not use sealed bandages, cast bandages, packs, etc., as irritation may occur.
6) Do not use this medicine for anal or vaginal compresses as it may cause irritation or chemical burns.
7) Do not use for any other purpose.
4. Precautions for storage
1) Avoid shading and keep in shading.
2) Keep it out of reach of children, and if a child swallows it, go to the hospital right away.
3) After use, close the product completely with a lid to prevent the product from drying out or entering foreign
Objects.

for external use only



HANNA HAND SANITIZER LIQUID

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71193-102
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**
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRUS LIMON FRUIT OIL (UNII: 0HNC1J1YED)	
WATER (UNII: 059QF0KO0R)	
D&C BLUE NO. 4 (UNII: 0KSY80VYS3)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71193-102-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph not final	part333A	09/03/2020	

Labeler - Rainbow Beauty Cosmetic Co., Ltd. (695684820)

Registrant - Rainbow Beauty Cosmetic Co., Ltd. (695684820)

Establishment

Name	Address	ID/FEI	Business Operations
Rainbow Beauty Cosmetic Co., Ltd.		695684820	manufacture(71193-102) , label(71193-102)

Revised: 9/2020 Rainbow Beauty Cosmetic Co., Ltd.