HANNA HAND SANITIZER- alcohol gel
Rainbow 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
DIPROPYLENE GLYCOL
CARBOMER
GLYCERIN
TRIETHANOLAMINE
PEG-60 HYDROGENATED CASTOR OIL
PANTHENOL
CAMELLIA SINENSIS LEAF EXTRACT
ALOE 8AR8ADENSIS LEAF EXTRACT
DIETHANOLAMINE
CITRUS PARADIS! (GRAPEFRUIT) PEEL OIL

Sterilization of hands and skin

KEEP OUT OF REACH OF THE CHILDREN

Apply to clean, dry hands. Apply sufficient amount to thoroughly wet all surfaces of hands and fingers. Rub onto hands until dry.

Supervise children in the use of this product.

■ Flammable. Keep away from fire or flame.

■ For external use only.

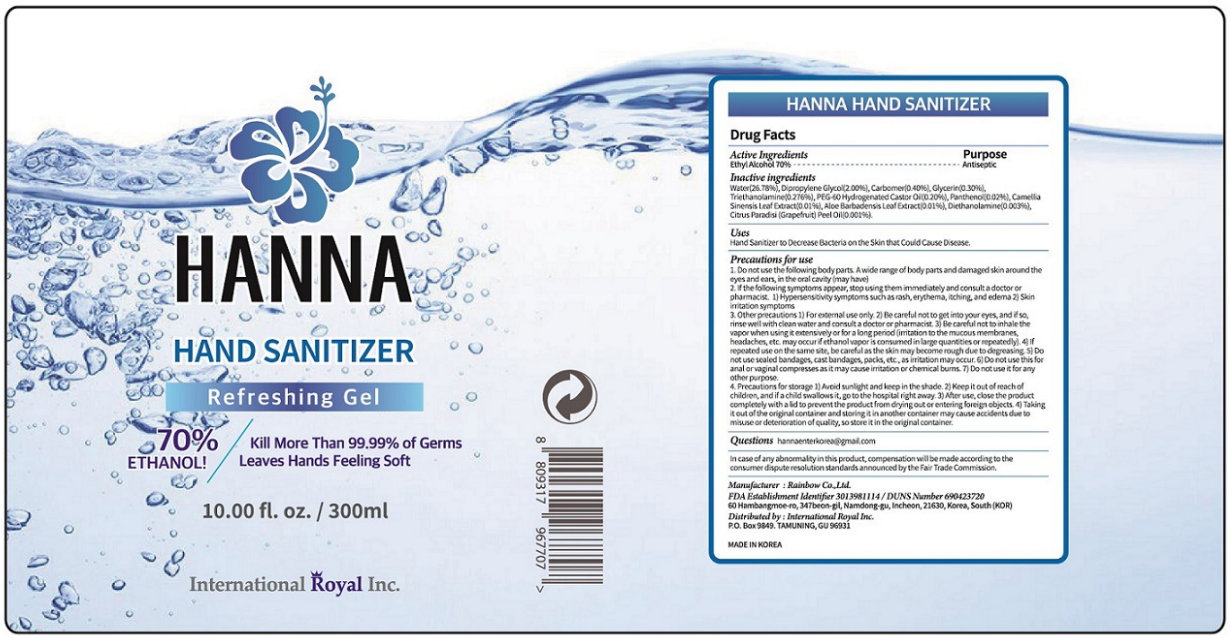
■ Do not use in eyes.

■ lf swallowed, get medical help promptly.

■ Stop use, ask doctor lf irritation occurs.

■ Keep out of reach of children.

for external use only



HANNA HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74247-0012
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**
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
GLYCERIN (UNII: PDC6A3C0OX)	
GRAPEFRUIT PEEL (UNII: 3582N05Q44)	
PEG-60 HYDROGENATED CASTOR OIL (UNII: 02NG325BQG)	
DIETHANOLAMINE (UNII: AZE05TDV2V)	
PANTHENOL (UNII: WV9CM0O67Z)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74247-0012-1	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
2	NDC:74247-0012-2	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
3	NDC:74247-0012-3	55 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
4	NDC:74247-0012-4	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
5	NDC:74247-0012-5	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
6	NDC:74247-0012-6	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
7	NDC:74247-0012-7	3 mL in 1 POUCH; Type 0: Not a Combination Product	06/13/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/13/2020	

Labeler - Rainbow Co Ltd (690423720)

Registrant - Rainbow Co Ltd (690423720)

Establishment

Name	Address	ID/FEI	Business Operations
Rainbow Co Ltd		690423720	manufacture(74247-0012)

Revised: 7/2020 Rainbow Co Ltd