DR. LIBEAUTE HAND SANITIZER- alcohol gel
Ongoong 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

ethyl alcohol

Water, Glycerin, carbomer, triethanolamine, tocopheryl acetate, pinis koralensis seed extract, camellia sinensis seed extract, aloe barbadensis left extract, melaleuca alternifolia (tea tree) leaf oil menthol, peppermint oil, butylene glycol, 1,2-hexanediol

Antiseptic

instant healthcare personnel hand antiseptic to reduce bacteria that potentially can cause disease

instant hand antiseptic to decrease bacteria on the skin
recommended for repeated use

hand sanitizer to help reduce bacteria on the 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.

For external use only.

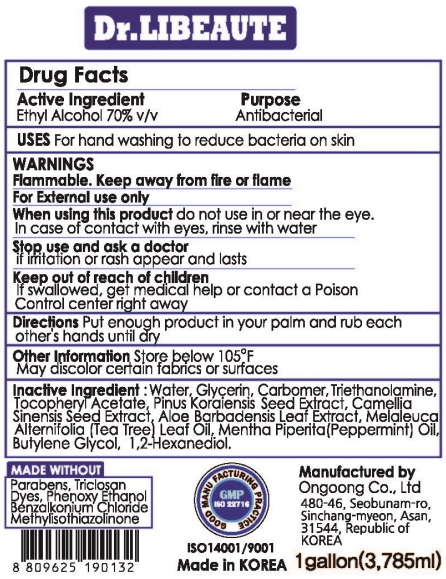
Flammable, keep away from fire or flame.

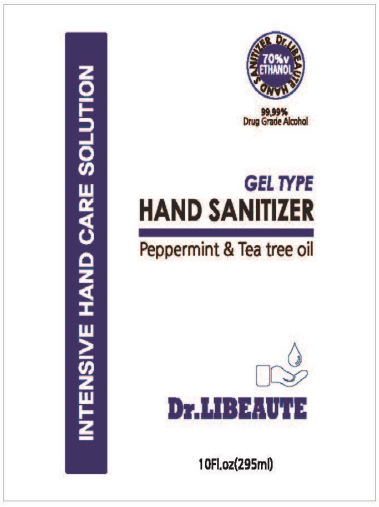
When using this product keep out of eyes. If contact with eyes occurs, rinse promptly and thoroughly with water.

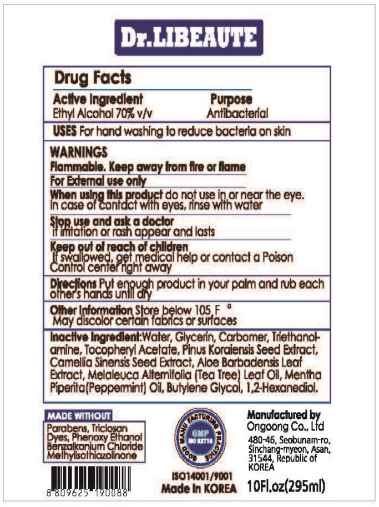
Stop use and ask a doctor if significant irritation or sensitization develops.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

for external use only







DR. LIBEAUTE HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74084-0009
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:74084-0009-1	3785 mL in 1 CONTAINER; Type 0: Not a Combination Product	05/16/2020	
2	NDC:74084-0009-2	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/16/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	05/16/2020	

Labeler - Ongoong Co Ltd (695625965)

Registrant - Ongoong Co Ltd (695625965)

Establishment

Name	Address	ID/FEI	Business Operations
Ongoong Co Ltd		695625965	label(74084-0009) , manufacture(74084-0009)

Revised: 5/2020 Ongoong Co Ltd