

HAND CLEANSER- 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 koraleensis 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	
Drug Facts	
Active Ingredient	Purpose
Ethyl Alcohol 70% v/v	Antibacterial
USES For hand washing to reduce bacteria on skin	
WARNINGS	
Flammable. Keep away from fire or flame	
For External use only	
When using this product do not use in or near the eye. In case of contact with eyes, rinse with water	
Stop use and ask a doctor if irritation or rash appear and lasts	
Keep out of reach of children If swallowed, get medical help or contact a Poison Control center right away	
Directions Put enough product in your palm and rub	

INTENSIVE HA



Dr.LIBEAUTE

17Fl.oz(500ml)

each other's hands until dry

Other Information Store below 105°F
May discolor certain fabrics or surfaces

Inactive Ingredient : Water, Glycerin, Carbomer, Triethanolamine, Tocopheryl Acetate, Pinus Koraiensis Seed Extract, Camellia Sinensis Seed Extract, Aloe Barbadensis Leaf Extract, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Mentha Piperita (Peppermint) Oil, Butylene Glycol, 1,2-Hexanediol.

Manufactured by
Ongoong Co., Ltd
17Fl.oz(500ml)



ISO14001/9001



INTENSIVE HAND CARE SOLUTION



**GEL TYPE
HAND CLEANSER**

Peppermint & Tea tree oil



Dr.LIBEAUTE

3.4Fl.oz(100ml)

Dr.LIBEAUTE

Drug Facts

Active Ingredient	Purpose
Ethyl Alcohol 70% v/v	Antibacterial

USES For hand washing to reduce bacteria on skin

WARNINGS

Flammable. Keep away from fire or flame

For External use only

When using this product do not use in or near the eye. In case of contact with eyes, rinse with water

Stop use and ask a doctor if irritation or rash appear and lasts

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control center right away

Directions Put enough product in your palm and rub each other's hands until dry

Other Information Store below 105°F
May discolor certain fabrics or surfaces

Inactive Ingredient : Water, Glycerin, Carbomer, Triethanolamine, Tocopheryl Acetate, Pinus Koraiensis Seed Extract, Camellia Sinensis Seed Extract, Aloe Barbadensis Leaf Extract, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Mentha Piperita (Peppermint) Oil, Butylene Glycol, 1,2-Hexanediol.

Manufactured by
Ongoong Co., Ltd
3.4Fl.oz(100ml)



ISO14001/9001



HAND CLEANSER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74084-0001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	70 g in 100 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74084-0001-1	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/21/2020	
2	NDC:74084-0001-2	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/21/2020	

Labeler - Ongoong Co Ltd (695625965)

Registrant - Ongoong Co Ltd (695625965)

Establishment

Name	Address	ID/FEI	Business Operations
Ongoong Co Ltd		695625965	label(74084-0001) , manufacture(74084-0001)

Revised: 3/2020

Ongoong Co Ltd