

DONGIL HAND CLEANER HAND SANITIZER- alcohol gel
DONG IL PHARMS 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

Glycerin, Chamomile Extract, Carbomer, Triethanolamine, Water

Antiseptic

KEEP OUT OF REACH OF THE CHILDREN

For the external use only

Do not use on the following body parts.

Around the eyes and ears, in the oral cavity, a wide range of body parts and damaged skin (may have irritating effects)

If the following symptoms appear, stop using them immediately and consult a doctor or pharmacist.

Symptoms of skin irritation

Other precautions

For external use only

Be careful not to get into your eyes, and if so, rinse well with clean water and consult a doctor or pharmacist.

Be careful not to inhale vapor when using it extensively or for a long time. (If you drink ethanol vapor in large quantities or repeatedly, irritation to the mucous membrane, headache, etc. may occur).

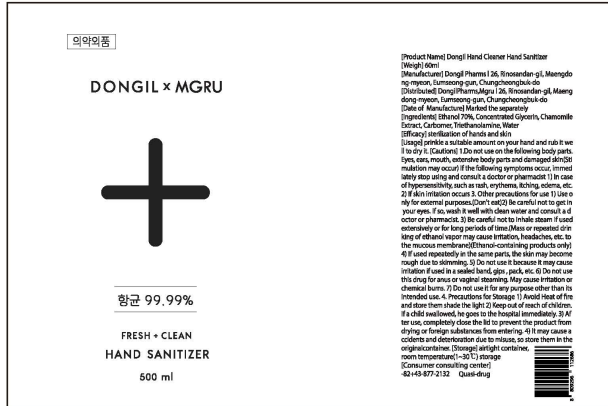
When repeated use on the same site, be careful as the skin may become rough due to degreasing.

When used in sealed bandages, cast bandages, packs, etc., irritation symptoms may appear.

Do not use for any other purpose.

Sprinkle a suitable amount on your hand and rub it well to dry it.

195mm*130



DONGIL HAND CLEANER HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73242-0 150
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	350 mL in 500 mL

Inactive Ingredients

Ingredient Name	Strength
CHAMOMILE (UNII: FGL3685T2X)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	

Packaging

		Marketing Start	Marketing End
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73242-0150-1	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/20/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/20/2020	

Labeler - DONG IL PHARMS CO., LTD. (557810721)

Registrant - DONG IL PHARMS CO., LTD. (557810721)

Establishment			
Name	Address	ID/FEI	Business Operations
DONG IL PHARMS CO., LTD.		557810721	manufacture(73242-0150)

Revised: 4/2020

DONG IL PHARMS CO., LTD.