

FRESH WATER HAND SANITIZER- alcohol gel
THESKINFACTORY 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 62% w/w

Water, Butylene Glycol, Glycerin, Carbomer, Triethanolamine, Sodium Hyaluronate, Camellia Sinensis Leaf Extract, Artemisia Princeps Extract

Antiseptic

KEEP OUT OF REACH OF THE CHILDREN

Put enough product in your palm to thoroughly cover your hands.

Rub hands together briskly until dry.

Children under 6 years of age should be supervised when using 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

주식회사 더 스킨팩토리 THE SKIN FACTORY		인쇄사양	
품목명	쿤달_손소독제_100ml_(아마존, 화인Ver)	<input type="checkbox"/> 칼선	
수정날짜	2020 / 08 / 12	<input checked="" type="checkbox"/> WHITE	
작업자	컨텐츠팀 최민용 010-2950-4855	<input type="checkbox"/> PANTONE 199 C	

작업지시사항 HISTORY	
2020-08-12	쿤달_손소독제_100ml_(아마존, 화인Ver)_ 신규

※ 답답은 본 이미지를 사내 폴더에 올렸음을 확인합니다.
 ※ 본 이미지는 사내 폴더에 올린 최종 데이터 이미지와 동일합니다.





FRESH WATER HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74773-0015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
ARTEMISIA PRINCEPS LEAF (UNII: SY077EW02G)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74773-0015-1	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/01/2020	

Labeler - THESKINFACTORY Co., Ltd. (694804099)

Registrant - THESKINFACTORY Co., Ltd. (694804099)

Establishment

Name	Address	ID/FEI	Business Operations
WHAIN COSMETIC		557837595	manufacture(74773-0015)

Establishment

Name	Address	ID/FEI	Business Operations
THESKINFACTORY Co., Ltd.		694804099	label(74773-0015)

Revised: 8/2020

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