

**DR.URBAN HAND SANITIZER- alcohol gel  
NEW SKIN**

*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.*

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**ACTIVE INGREDIENT**

Alcohol 70% w/w

**INACTIVE INGREDIENTS**

Water, Glycerin, Carbomer, Triethanolamine

**PURPOSE**

Antiseptic

**WARNINGS**

For external use only. Flammable. Keep away from heat or flame

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Do not use

- in children less than 2 months of age
- on open skin wounds

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When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

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Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

**KEEP OUT OF REACH OF CHILDREN**

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

**Uses**

Hand sanitizer to help reduce bacteria that potentially can cause disease.

**Directions**

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

**Other information**

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

**PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**

**의약품**

**닥터어반 손소독제겔 (에탄올)**

평균 99.9% 닥터어반 손소독제겔은 언제 어디서나 간편하게 손, 피부 등의 상균소독으로 위생상태를 절감하게 유지시켜주며, 각종 손에 의한 질병감염예방에도 효과적입니다.

**상표명**: 닥터어반 손소독제겔(에탄올) **용량**: 500ml **성상**: 무색의 투명인 겔  
**용법**: 손에 적당량을 뿌리거나 얇은 층을 바르십시오. **효능/효과**: 손, 피부 등의 상균소독 **주성분**: 100g 중 에탄올 70% 에탄올(카미르산) 70%  
**제조번호 및 사용기한**: 별도표기 **저장방법**: 실온보관(1~30°C)보관, 기밀용기  
**제조판매업**: 뉴스킨 / 경기도 고양시 일산서구 송호로 425번길92-5  
**소비자상담전화**: 032-715-4448

**사용 주의사항**: 1. 어린이의 손바닥에 사용되지 않습니다. 눈, 피부, 코, 구강에 들어가지 않도록 주의하십시오. 2. 알레르기 반응이 나타나지 않는 한, 계속 사용할 수 있습니다. 3. 알레르기 반응이 나타나면, 사용을 중단하십시오. 4. 사용 후, 손을 깨끗이 씻어주세요. 5. 알레르기 반응이 나타나면, 사용을 중단하십시오. 6. 알레르기 반응이 나타나면, 사용을 중단하십시오. 7. 알레르기 반응이 나타나면, 사용을 중단하십시오. 8. 알레르기 반응이 나타나면, 사용을 중단하십시오. 9. 알레르기 반응이 나타나면, 사용을 중단하십시오. 10. 알레르기 반응이 나타나면, 사용을 중단하십시오.

**전성분**: 주성분(에탄올 70%), 장가제(정제수, 글리세린, 카보머, 트리에탄올아민)  
 본 제품에 이상이 있을 경우, 고객센터(032-715-4448) 또는 소외차별감정기관에 의해 보상에 드립니다.

MADE IN KOREA



**HAND  
SANITIZER**

70% ETHANOL  
KILLS 99.9% OF GERMS

닥터 어반  
손소독제겔(에탄올)

500ml (19.6 fl. oz.)

손, 피부 등의 살균소독제

**DR.URBAN HAND SANITIZER GEL (gel type)**

<b>Drug Facts</b>	
<b>Active ingredients</b>	<b>Purpose</b>
Alcohol 70% w/w	Antiseptic
<b>Uses</b> Hand sanitizer to help reduce bacteria that potentially can cause disease.	
<b>Warnings</b>	
For external use only. Flammable. Keep away from heat or flame.	
Do not use - in children less than 2 months of age - on open skin wounds	
<b>When using this product</b> Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.	
<b>Stop use and ask a doctor</b> if irritation or rash occurs. These may be signs of a serious condition.	
<b>Keep out of reach of children.</b> If swallowed, get medical help or contact a Poison Control Center right away.	
<b>Directions</b>	
Place enough product on hands to cover all surfaces. Rub hands together until dry.	
Supervise children under 6 years of age when using this product to avoid swallowing.	
<b>Other information</b>	
Store between 15-30C (59-86F)	
Avoid freezing and excessive heat above 40C (104F)	
<b>Inactive ingredients</b>	
Water, Glycerin, Carbomer, Triethanolamine	

**Manufacturer**: NEWSKIN / 92-5, Songporo 425beom-gil, Ilsanseo-gu, Gyeonggi-do, Republic of Korea  
**Distributor**: Root and fruit  
**Net Wt**: 500ml (19.6 Fl. Oz.)



**DR.URBAN HAND SANITIZER**

alcohol gel

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:79 282-0 10
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	Alcohol	350 g in 500 mL

**Inactive Ingredients**

Ingredient Name	Strength
Glycerin (UNII: PDC6A3C0OX)	
CARBOMER HO MO POL YMER, UNSPECIFIED TYPE (UNII: 0 A5MM30 7FC)	
TROLAMINE (UNII: 9O3K93S3TK)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79 282-0 10-01	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	06/01/2020	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/01/2020	

Labeler - NEW SKIN (695538784)

**Registrant** - NEW SKIN (695538784)

**Establishment**

Name	Address	ID/FEI	Business Operations
NEW SKIN		695538784	manufacture(79282-010)

Revised: 6/2020

NEW SKIN