WANCARE ONEHAND HAND SANITIZER- ethyl alcohol solution KAF GRUP SAGLIK HIZMETLERI INSAAT SANAYI VE TICARET LIMITED SIRKETI

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.

WANCARE ONEHAND HAND SANITIZER

DRUG FACTS

Active ingredient(s)

Ethyl Alcohol 80% (v/v)

Purpose

Antiseptic

Use(s)

Hand sanitizer to help reduce bacteria on the skin that could cause disease. Recommended for repeated use.

Warnings

For external use only.

Flammable. Keep away from fire or flame.

Do not use

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

When using this product keep out of eyes, ears and mouth. In case of contact, rinse eyes thoroughly with water. Do not inhale or ingest.

Stop use and ask a doctor if irritation or rash appears and lasts. Condition persists for more than 72 hours.

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

Directions

Place 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.

Other Information

Store between 5-40C(41-104F)

Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Water (Aqua), Glycerin, Fragrance

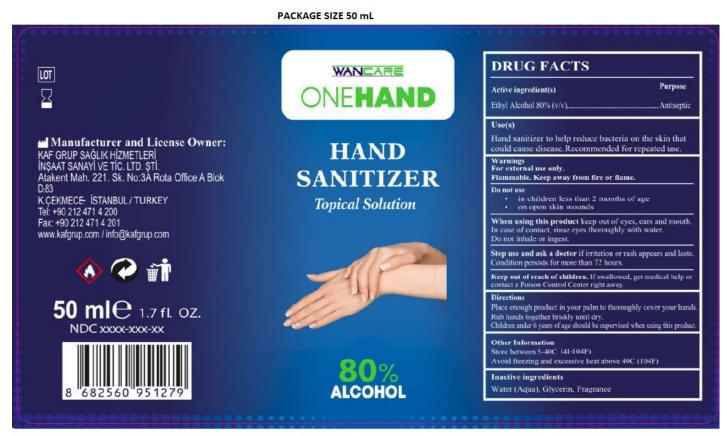
80% ALCOHOL

Topical Solution

Manufacturer and License Owner:

KAF GRUP SAGLIK HIZMETLERI INSAAT SANAYI VE TIC. LTD. STI. Atakent Mah. 221. Sk. No:3A Rota Office A Blok D:83 K.CEKMECE -ISTANBUL/TURKEY Tel: +90 212 471 4 200 Fax: +90 212 471 4 201 www.kafgrup.com/info@kafgrup.com

Packaging



PACKAGE SIZE 100 mL





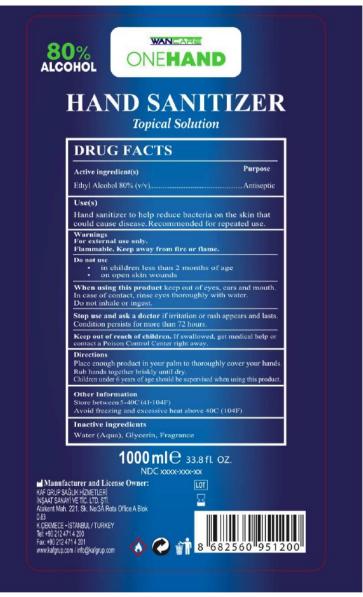
PACKAGE SIZE 250 mL



PACKAGE SIZE 500 mL



PACKAGE SIZE 1000 mL



PACKAGE SIZE 5000 mL



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ethyl alcohol solution		
Product Information		
Product Type	HUMAN OTC DRUG	Item Code (Source)
Route of Administration	TOPICAL	
Route of Administration	TOPICAL	

WANCARE ONEHAND HAND SANITIZER

Route of Administration	TOPICAL		
Active Ingredient/Active Moi	ety		
Ingred	ient Name	Basis of St	trength Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	80 mL in 100 mL
Inactive Ingredients			
	Ingredient Name		Strength
WATER (UNII: 059QF0KO0R)			
GLYCERIN (UNII: PDC6A3C0OX)			

NDC:81314-102

Packaging							
# Item Code		Package Description	Marketing Start Date	Marketing End Date			
1 NDC:81314-102-01	50 mL in	1 BOTTLE; Type 0: Not a Combination Product	12/18/2020				
2 NDC:81314-102-02	100 mL i	n 1 BOTTLE; Type 0: Not a Combination Product	12/18/2020				
3 NDC:81314-102-03	250 mL i	n 1 BOTTLE; Type 0: Not a Combination Product	12/18/2020				
4 NDC:81314-102-04	500 mL i	n 1 BOTTLE; Type 0: Not a Combination Product	12/18/2020				
5 NDC:81314-102-05	1000 mL	in 1 BOTTLE; Type 0: Not a Combination Product	12/18/2020				
6 NDC:81314-102-06	5000 mL	in 1 BOTTLE; Type 0: Not a Combination Product	12/18/2020				
Marketing Information							
Marketing Categ	ory A	pplication Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not	final nar	333A	12/18/2020				

Labeler - KAF GRUP SAGLIK HIZMET LERI INSAAT SANAYI VE TICARET LIMITED SIRKETI (519929115)

Establishment						
	Name	Address	ID/FEI	Business Operations		
KAF GRUP SAGLIK HIZMETLERI INSAAT SANAYI VE TICARET LIMITED SIRKETI			519929115	manufacture(81314-102)		
Revised: 12/2020	KAF GRUP SAGLIK HIZMETLERI INSAA	T SANAY	I VE TICAF	RET LIMITED SIRKETI		