

BEIS CLEAN KIT- alcohol

Beis LLC

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.

BEIS Clean Kit

Drug Facts

Active Ingredient

Ethyl Alcohol 75% v/v

Purpose

Antiseptic

Uses

- For hand washing to help reduce bacteria that potentially can cause disease. Recommended for repeated use. For use when soap and water are not available.

Warnings

For external use only

Flammable, Keep away from heat or flame

When using this product

do not use in or near eyes, ears and mouth, In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or rash appears on the skin.

Keep out of reach of children.

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

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Children under 6 years of age should be supervised by adult when applying this product.
- Do not use on children less than 2 months of age or on open skin wounds.

Other information

- Store between 15-30°C (59-86°F)
- Avoid freezing and excessive heat above 40°C (104°F)
- May discolor some fabrics or surfaces

Inactive ingredients

Water, Glycerin, Squalane, Panthenol, Inulin, Sodium Hyaluronate, Fructose, Tocopheryl Acetate, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol.

Drug Facts**Active Ingredient**

Ethyl alcohol 75%

Purpose

Antiseptic

Use

For hand washing to decrease bacteria on the skin

Warnings

For external use only

Flammable, keep away from fire or flame.

Do not use

- in the eyes

Stop use and ask a doctor if

- irritation and redness develop
- 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

wet hands thoroughly with wipe and allow to dry without wiping

Inactive Ingredients

Purified water

Package Labeling:81263-001-00

BÉIS

ADVANCED
HAND
SANITIZER

WITH MOISTURIZERS
& VITAMIN E

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Manufactured by
Zhejiang iColor Biotech Co., Ltd
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New York, NY 10168
Tel: +1 212-551-2657
Made in China

2 FL.OZ/ 60mL

Package Labeling:81263-002-00



BEIS CLEAN KIT

alcohol kit

Product Information					
Product Type		HUMAN OTC DRUG	Item Code (Source)		NDC:81263-003
Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:81263-003-01	1 in 1 KIT	08/01/2021		
Quantity of Parts					
Part #	Package Quantity		Total Product Quantity		
Part 1	1 BOTTLE		60 mL		
Part 2	10 PATCH		40 mL		
Part 1 of 2					
BEIS ADVANCED HAND SANITIZER					
alcohol gel					

Product Information	
Item Code (Source)	NDC:81263-001
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SQUALANE (UNII: GW89575KF9)	
PANTHENOL (UNII: WW9CM0O67Z)	
INULIN (UNII: JOS53KRJ01)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
FRUCTOSE (UNII: 6YSS42VSEV)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81263-001-00	60 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/01/2021	

Part 2 of 2
BEIS HOT MESS ANTIBACTERIAL WIPES alcohol cloth

Product Information	
Item Code (Source)	NDC:81263-002
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81263-002-00	10 in 1 PACKET		
1		4 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	08/01/2021	

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OTC monograph not final	part333E	08/01/2021	

Labeler - Beis LLC (117648333)

Revised: 11/2021

Beis LLC