QUR LAVENDER ESSENTIAL OIL HAND SANITIZER- alcohol gel K7 Design Group Inc.

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.

QUR Lavender Essential Oil Hand Sanitizer

Drug Facts

Active ingredient

Alcohol 69% v/v

Purpose

Antiseptic

Use

for hand-washing to decrease bacteria on the skin, only when water is not available

Warnings

Flammable, keep away from fire and flames

For external use only

When using this product

- do not get into eyes.
- if contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if

• irritation and redness develop

Keep out of reach of children.

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

Directions

wet hands thoroughly with product and allow to dry without wiping

Inactive ingredients

Water, Glycerin, Propylene Glycol, Carbomer, Aloe Barbadensis Leaf Extract, Aminomethyl Propanol, Tocopheryl Acetate, Lavandula Angustifolia (Lavender) Oil, Denatonium Benzoate.

Company Information

MANUFACTURED FOR & DISTRIBUTED BY QUR NEW YORK, NY 10128

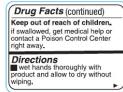
Product Packaging













QUR LAVENDER ESSENTIAL OIL HAND SANITIZER

alcohol gel

 Product Information

 Product Type
 HUMAN OTC DRUG
 Item Code (Source)
 NDC:74177-991

 Route of Administration
 TOPICAL

l	Active Ingredient/Active Moiety		
	Ingredient Name	Basis of Strength	Strength
	ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	69 mL in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)			
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)			
WATER (UNII: 059QF0KO0R)			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)			
DENATO NIUM BENZO ATE (UNII: 4YK5Z54AT2)			
GLYCERIN (UNII: PDC6A3C0OX)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)			
LAVENDER O IL (UNII: ZBP1YXW0H8)			

F	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:74177-991- 01	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/19/2021			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	01/19/2021			

Labeler - K7 Design Group Inc. (080357784)

Revised: 1/2021 K7 Design Group Inc.