CLEAN RX- ethyl alcohol solution
Quality Innovations 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.

Active Ingredient(s)

Ethyl Alcohol 75%

Purpose

Antiseptic

Use

Helps decrease bacteria on skin. Recommended for repeated use.

Warnings

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

Do not apply

around eyes

When using this product

avoid contact with eyes. In case of contact flush eyes with water.

Stop use and ask a doctor

If irritation or redness develop and conditions 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. Children must be supervised in use of this product.

Directions

pump as needed into your palmsand throughly spread on both the hands.

Other information

- Store at 20°C (68° to 77°F).
- May discolor fabrics

Inactive ingredients

acrlates/C10-30 alkyl acrylates crosspolymer, benzophenone-4, glycerin, tocopherol, water

Principal Display Panel

NDC: 74235-075-02

Clean Rx^{TM} 2 oz

A percentage of the proceeds from this product will be donated towards the coronavirus epidemic

HAND SANITIZER

Kills 99.999% of GERMS

With Moisturizer and Vitamin E

2 FL OZ (60 mL)

NDC: 74235-075-08

Clean Rx^{TM} 8 oz

A percentage of the proceeds from this product will be donated towards the coronavirus epidemic

HAND SANITIZER

Kills 99.999% of GERMS

With Moisturizer and Vitamin E

8 FL OZ (240 mL)

CLEAN RX

ethyl alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74235-075
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**
GLYCERIN (UNII: PDC6A3C0OX)	
SULISOBENZONE (UNII: 1W6L629B4K)	
WATER (UNII: 059QF0KO0R)	
TOCOPHEROL (UNII: R0ZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74235-075-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2020	
2	NDC:74235-075-08	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/06/2020	

Labeler - Quality Innovations Inc. (079258818)

Registrant - Quality Innovations Inc. (079258818)

Revised: 4/2020 Quality Innovations Inc.