

QC 8OZ INSTANT HAND SANITIZER (ORIGINAL)- alcohol gel
Ningbo Liyuan Daily Chemical Products Co., Ltd.

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.

Drug Facts

Medicinal Ingredient

Ethyl Alcohol 62%

Purpose

Antiseptic

Non-Medicinal

Water, Isopropyl Alcohol, Glycerin, Carbomer, Aminomethyl Propanol, Parfum, Propylene Glycol, Isopropyl Myristate, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate

Features

- To decrease bacteria on the skin that could cause disease.
- Recommended for repeated use.
- Use anywhere without water.

Directions

Wet hands thoroughly with product and rub until dry without wiping.

For children under 6, use only under adult supervision.

Not recommended for infants.

Warning

- For external use only-hands
- Flammable, keep away from heat and flame
- Discontinue if skin becomes irritated and ask a doctor

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

When using this product

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- Do not inhale or ingest.
- Avoid contact with broken skin.

This product was not tested on animals

Other information

- Do not store above 105°.

- May discolor some fabrics.
- Harmful to wood finishes and plastics.



QC
QUALITY CHOICE



Compared to Purell® Instant Hand Sanitizer*

Instant Hand Sanitizer

Original

Kills 99.99% of Germs
Leaves hands feeling soft
Moisturizes with Vitamin E

8 FL OZ (236 mL)

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<p>*This product is not manufactured or distributed by GOJO Industry, Inc., owner of the registered trademark Purell®</p>	
	<p>Distributed by C.D.M.A., Inc.® 43157 W. Nine Mile Novi, MI 48376-0995 www.qualitychoice.com Questions: 248-449-9300</p> <p>Made in China</p> <p>This product was not tested on animals</p>

QC 8OZ INSTANT HAND SANITIZER (ORIGINAL)

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76176-035
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)	36 mL in 100 mL
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76176-035-01	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/09/2017	

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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/09/2017	

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Labeler - Ningbo Liyuan Daily Chemical Products Co., Ltd. (530766098)

Registrant - Ningbo Liyuan Daily Chemical Products Co., Ltd. (530766098)

Establishment

Name	Address	ID/FEI	Business Operations
Ningbo Liyuan Daily Chemical Products Co., Ltd.		530766098	manufacture(76176-035)

Revised: 9/2017

Ningbo Liyuan Daily Chemical Products Co., Ltd.