

LEVEL 1 HEALTH HAND SANITIZER LIQUID NON-STERILE SOLUTION- alcohol hand sanitizer liquid

Continental Manufacturing Chemist, 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.

Drug Facts

Active Ingredient[s]

Alcohol 80% v/v

Purpose

Antiseptic

Use[s]

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat 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 with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children.

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

Direction

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

- Store between 15 °-30 ° C (59 ° 86 °F)
- Avoid freezing and excessive heat above 40 °C (104 °F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Principal Display Panel

Level 1 Health

Hand Sanitizer

Liquid

Non-Sterile Solution

Topical Solution

80% Alcohol Antiseptic

XX fl oz (X mL)

Manufactured by

Continental Manufacturing Chemist, Inc.

1501 Blue Sky Blvd, Huxley, IA 50124

www.cmchemist.com

515.795.2000

Made in USA

UPC Code

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LEVEL 1 HEALTH

**HAND
SANITIZER**

LIQUID
Non-sterile Solution

TOPICAL SOLUTION

80% | Alcohol
Antiseptic

32 FL OZ (946.37 mL)



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LEVEL 1 HEALTH HAND SANITIZER LIQUID NON-STERILE SOLUTION

alcohol hand sanitizer liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:49794-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49794-002-01	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/08/2020	

Labeler - Continental Manufacturing Chemist, Inc. (005278007)

Establishment

Name	Address	ID/FEI	Business Operations
Continental Manufacturing Chemist, Inc.		005278007	repack(49794-002) , manufacture(49794-002)

Establishment

Name	Address	ID/FEI	Business Operations
Continental Manufacturing Chemist		081171390	manufacture(49794-002) , repack(49794-002)

Revised: 5/2020

Continental Manufacturing Chemist, Inc.