

ALCO-GEL- alcohol gel
Brainerd Chemical Company 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.

Alco-Gel

Drug Facts

Active ingredient[s]

Isopropyl alcohol 75% 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.

Directions

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

Inactive ingredients

hydrogen peroxide, purified water USP, d-limonene, cocamidopropyl betaine, hydroxypropyl methylcellulose

Distributed By:

Nyco Products Company
5332 Dansher Rd. • Countryside, IL 60525

PRINCIPAL DISPLAY PANEL - 3.785 L Jug Label

Nyco®
Hand Care

Alco-Gel Plus

Hand Sanitizer

- Kills 99.99% of common germs that can cause illness
- Evaporates Quickly • Leaves hands smooth and refreshed

KEEP OUT OF THE REACH OF CHILDREN.
FOR INSTITUTIONAL AND INDUSTRIAL USE

NET CONTENTS: 1 GAL / 3.785 L

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Isopropyl Alcohol Antiseptic 75% (Gel)

Topical Solution
Hand Sanitizer
Non- Sterile Solution



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Inactive Ingredients: Hydrogen Peroxide, Purified Water USP, d-Limonene, Cocamidopropyl Betaine, Hydroxypropyl methylcellulose

DANGER

Hazard Statements: Highly flammable liquid and vapor. Causes serious eye irritation. May cause drowsiness or dizziness.

Precautionary Statements: Keep away from heat, open flames, sparks - No smoking. Keep container closed. Ground/bond container and receiving equipment. Use explosion proof electrical, ventilation away from clothing. Use non-sparking tools. Take precautionary measures against static discharge. Avoid breathing dust, fume, gas, mist, vapors and spray. Wear protective gloves, eye and face protection.

IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. If eye irritation persists, seek medical advise/attention.

SPECIFIC TREATMENT: (See First Aid section on the SDS.) Rinse mouth. Wash contaminated clothing before reuse.

IN CASE OF FIRE: Use dry san, dry chemical or alcohol resistant foam to extinguish. Store in a well ventilated place. Keep container tightly closed. Store in well ventilated place. Keep cool. Store locked up. Dispose of contents/container to licensed waste handling facility.



Distributed By:
Nyco Products Company
5332 Dansher Rd. • Countryside, IL 60525
(800) 752-4754 • nycoproducts.com
MADE IN THE U.S.A.



ALCO-GEL

alcohol gel

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:54555-131

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)	alcohol	750 mL in 1 L

Inactive Ingredients

Ingredient Name	Strength
Cocamidopropyl Betaine (UNII: 5OCF3011KX)	
hydrogen peroxide (UNII: BBX060AN9V)	
water (UNII: 059QF0K00R)	
LIMONENE OXIDE, TRANS-(+)- (UNII: 8VUQ1B30IK)	
HYDROXYMETHYL CELLULOSE (UNII: 273FM27VK1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54555-131-04	1040.9 L in 1 TANK; Type 0: Not a Combination Product	07/01/2020	
2	NDC:54555-131-03	208.1 L in 1 DRUM; Type 0: Not a Combination Product	07/01/2020	
3	NDC:54555-131-02	18.9 L in 1 JUG; Type 0: Not a Combination Product	07/01/2020	
4	NDC:54555-131-01	3.785 L in 1 JUG; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333A	07/01/2020	

Labeler - Brainerd Chemical Company Inc., (787442417)

Establishment

Name	Address	ID/FEI	Business Operations
Brainerd Chemical Company Inc.,		787442417	MANUFACTURE(54555-131) , LABEL(54555-131) , PACK(54555-131) , REPACK(54555-131)

Revised: 6/2020

Brainerd Chemical Company Inc.,