

HAND SANITIZER CA75- alcohol liquid
Pharmco Laboratories, Inc.

Hand Sanitizer
CA75

Drug Facts

Active Ingredient

Ethyl Alcohol 70% v/v

Purpose

Antimicrobial

Uses

- Hand sanitizer to help reduce bacteria on the skin that could cause disease.
- Recommended for repeated use.

Warnings

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

Do not use

- on 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.

Discontinue use if irritation or redness develops.

If condition persists for more than 72 hours, consult a doctor.

Keep out of reach of children. If swallowed get medical help or call a poison control center immediately.

Directions

- Place enough product in your palm to thoroughly cover your hands.
- Rub hands together briskly until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other Information

- Store below 40°C (104°F)
- May discolor certain fabrics or surfaces.

Inactive Ingredients

Purified Water USP, Glycerin USP, Jojoba Oil

Questions or comments?

call 1.800.635.0712 Monday through Friday 8:00 AM to 4:30 PM

Manufactured by:
Pharmco Laboratories, Inc.
3520 South St. Titusville, FL 32780

PRINCIPAL DISPLAY PANEL - 59 mL Tube Label

pharmco
MANUFACTURER OF PREMIUM SKIN CARE PRODUCTS

HAND
SANITIZER

2 fl oz. (59 mL)



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<p align="center">Manufactured by: Pharmco Laboratories, Inc. 3520 South St. Titusville, FL 32780 www.pharmcolabs.com</p>	



PLOTG-REV0001

HAND SANITIZER CA75			
alcohol liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58400-012
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
Alcohol (UNII: 3K9958V90M) (Alcohol - UNII:3K9958V90M)		Alcohol	70 mL in 100 mL
Inactive Ingredients			
Ingredient Name			Strength
Water (UNII: 059QF0KO0R)			
Glycerin (UNII: PDC6A3C0OX)			

Jojoba Oil (UNII: 724GKU717M)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58400-012-02	59 mL in 1 TUBE; Type 0: Not a Combination Product	06/29/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M003	05/01/2020	

Labeler - Pharmco Laboratories, Inc. (096270814)

Establishment

Name	Address	ID/FEI	Business Operations
Pharmco Laboratories, Inc.		096270814	MANUFACTURE(58400-012) , LABEL(58400-012) , PACK(58400-012)

Revised: 1/2024

Pharmco Laboratories, Inc.