CAREONE LEMON- ethyl alcohol liquid AMERICAN SALES COMPANY

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

ETHYL ALCOHOL 70%

PURPOSE

ANTISEPTIC

USES

TO HELP REDUCE BACTERIA ON THE SKIN

WARNINGS

FOR EXTERNAL USE ONLY. FLAMMABLE. KEEP AWAY FROM SOURCE OF HEAT OR FIRE

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE THOROUGHLY WITH WATER

STOP USE AND ASK A DOCTOR IF

IRRITATION OR REDNESS DEVELOPS AND LASTS

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

DIRECTIONS

PUT ENOUGH PRODUCT IN YOUR PALM TO COVER HANDS AND RUB HANDS TOGETHER UNTIL DRY. CHILDREN UNDER 6 YEARS SHOULD BE SUPERVISED WHEN USING THIS PRODUCT

OTHER INFORMATION

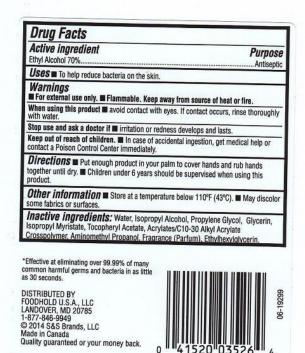
STORE AT A TEMPERATURE BELOW 110°F (43°C). MAY DISCOLOR SOME FABRICS OR SURFACES

INACTIVE INGREDIENTS

WATER, ISOPROPYL ALCOHOL, PROPYLENE GLYCOL, GLYCERIN, ISOPROPYL MYRISTATE, TOCOPHERYL ACETATE, ACRYLATES/C10-30 ALKYL CROSSPOLYMER, AMINOMETHYL PROPANOL, FRAGRANCE (PARFUM), ETHYLHEXYLGLYCERIN

LABEL COPY





CAREONE LEMON

ethyl alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:41520-941

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)
ALCOHOL (UNII: 3K9958V90M) ALCOHOL 700 mg in 1 mL

Inactive Ingredients					
Ingredient Name	Strength				
WATER (UNII: 059QF0KO0R)					
ISOPROPYL ALCOHOL (UNII: ND2M416302)					
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)					
GLYCERIN (UNII: PDC6A3C0OX)					
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)					
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)					
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)					
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)					
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)					

Packaging								
#	Item Code	Package Description	Market	ing Start Date	Marketing End Date			
1	NDC:41520-941-08	236 mL in 1 BOTTLE, PLASTIC						
Marketing Information								
	Marketing Category	Application Number or Monograp	h Citation	Marketing Start Date	Marketing End Date			
O	ΓC monograph not final	part333E		0 1/26/20 14				

Labeler - AMERICAN SALES COMPANY (809183973)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment								
Name	Address	ID/FEI	Business Operations					
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(41520-941)					

Revised: 1/2014 AMERICAN SALES COMPANY