

**ANTIBACTERIAL HAND SANITIZER SUNFLOWER AND CITRUS- ethyl alcohol gel  
APOLLO HEALTH AND BEAUTY CARE**

*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.*

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**DRUG FACTS**

**ACTIVE INGREDIENT**

ETHYL ALCOHOL 62 PERCENT

**PURPOSE**

ANTISEPTIC

**USES**

TO HELP REDUCE BACTERIA ON THE SKIN.

**WARNINGS**

FOR EXTERNAL USE ONLY.

FLAMMABLE. KEEP AWAY FROM FIRE OR FLAME.

**WHEN USING THIS PRODUCT**

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

**STOP USING THIS PRODUCT AND ASK DOCTOR IF**

IRRITATION OR REDNESS DEVELOPS OR LASTS.

**KEEP OUT OF REACH OF CHILDREN**

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

**DIRECTIONS**

WET HANDS THOROUGHLY AND RUB TOGETHER UNTIL DRY.

**QUESTIONS OR COMMENTS**

1-866-695-3030

**INACTIVE INGREDIENTS**

WATER, PROPYLENE GLYCOL, ISOPROPYL MYRISTATE, ACRYLATES/C10-30 ALKYL ACRYLATES CROSSPOLYMER, GLYCERIN, FRAGRANCE, AMINOMETHYL PROPANOL, TOCOPHERYL ACETATE, MANNITOL, CELLULOSE, HYDROXYPROPYL METHYLCELLULOSE, CAPRYLIC/CAPRIC TRIGLYCERIDE, IRON OXIDES (CI 77491, CI 77492, CI 77891), MICA (CI 77019), TITANIUM DIOXIDE (CI 77891), YELLOW 5 (CI 19140), YELLOW 6 (CI 15985).



## ANTIBACTERIAL HAND SANITIZER SUNFLOWER AND CITRUS

ethyl alcohol gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63148-270
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
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
MANNITOL (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
CAPRYLIC/CAPRIC MONO/DIGLYCERIDES (UNII: U72Q2I8C85)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
MICA (UNII: V8A1AW0880)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63148-270-09	259 mL in 1 BOTTLE, PUMP		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/15/2011	

**Labeler** - APOLLO HEALTH AND BEAUTY CARE (201901209)

**Registrant** - APOLLO HEALTH AND BEAUTY CARE (201901209)

## Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture

Revised: 4/2011

APOLLO HEALTH AND BEAUTY CARE