HAND SANITIZER- alcohol gel Cardinal Health

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

Active ingredient

Ethyl alcohol 70%

purpose

Purpose

Antiseptic

Uses

Uses

- to decrease bacteria on the skin that could cause disease
- recommend for repeated use

Warnings

Warnings

For external use only-hands

Flammable

Flammable. Keep away from heat and flame

When using

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor

Stop use and ask a doctor if skin irritation develops

Keep out of reach of children

Keep out of reach of children. If swallowed get nedical help or contact a Poison Control Center right away

Directions

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

Other information

- do not sore above 1050 F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

Inactive ingredients water glyceryl caprylate/caprate, glycerin, isopropyl myristate, tocopheryl acetate, acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, benzophenone-4

Claims

Effective at eliminating more than 99.99% of many common harmful germs and bacteria in as little as 15 seconds

Dis claimer

This product is no manufactured or distributed by GOJO Industries, Inc., distributor of Purell refreshing Gel Advanced Hand Sanitizer

Adverse Reactions Section

DISTRIBUTED BY: CARDINAL HEALTH, DUBLIN, OHIO 43017

www.myleader.com

Principal Display Panel

NDC 37205-733-34

LEADER

More

Effective

Formula

Compare to Purell

Advanced

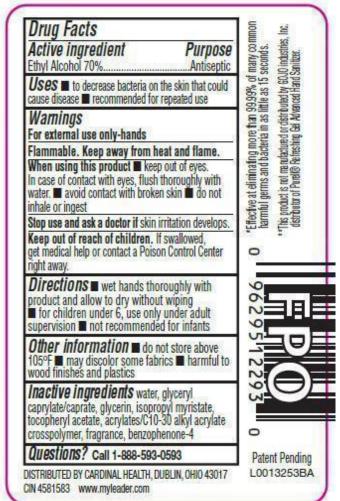
Hand

Sanitizer

Kills more than 99.99% of germs

8 FL OZ (236 mL)





HAND SANITIZER

alcohol gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37205-733	
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)			
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)			
GLYCERIN (UNII: PDC6A3C0OX)			
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)			
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)			

CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809 Y72KV36)

SULISOBENZONE (UNII: 1W6L629B4K)

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l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1 NDC:37205-733-34	236 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	11/09/2012		

Labeler - Cardinal Health (097537435)

Registrant - Vi-Jon (790752542)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(37205-733)

Revised: 11/2012 Cardinal Health