HAND SANITIZER WITH MOISTURIZERS AND VITAMIN E- ethyl alcohol gel DZA BRANDS

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 65 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 AND LASTS.

KEEP OUT OF REACH OF CHILDREN

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

DIRECTIONS

APPLY A SMALL AMOUNT IN YOUR PALM AND RUB HANDS TOGETHER BRISKLY UNTIL DRY. CHILDREN UNDER 6, SHOULD BE SUPERVISED WHEN USING THIS PRODUCT.

OTHER INFORMATION

DO NOT STORE ABOVE 110^{0} F (43^{0} C).

QUESTIONS OR COMMENTS

1-866-322-2439

INACTIVE INGREDIENTS

WATER, ISOPROPYL ALCOHOL, GLYCERIN, CARBOMER, AMINOMETHYL PROPANOL, FRAGRANCE, PROPYLENE GLYCOL, ISOPROPYL MYRISTATE, TOCOPHERYL ACETATE.

FRONT AND BACK LABELS



HAND SANITIZER WITH MOISTURIZERS AND VITAMIN E

ethyl alcohol gel

Product Information			
Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:55316-267
Route of Administration	TOPICAL		
Active Ingredient/Activ	ve Moiety		
	Ingredient Name	Basis of Strength	Strength
	0M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	65 mL in 100 mL
•		ALCOHOL	
		ALCOHOL	
Inactive Ingredients	Ingredient Name	ALCOHOL	Strength
Inactive Ingredients WATER (UNII: 059QF0KO0R)	Ingredient Name	ALCOHOL	
Inactive Ingredients WATER (UNII: 059QF0K00R) ISOPROPYL ALCOHOL (UN	Ingredient Name	ALCOHOL	
Inactive Ingredients WATER (UNII: 059QF0KO0R) ISOPROPYL ALCOHOL (UN GLYCERIN (UNII: PDC6A3C0	Ingredient Name) NII: ND2M416302) OX)		
Inactive Ingredients WATER (UNII: 059QF0KO0R) ISOPROPYL ALCOHOL (UN GLYCERIN (UNII: PDC6A3C0 CARBOMER 934 (UNII: Z135)	Ingredient Name) NII: ND2M416302) OX) WT9208)		
	Ingredient Name) NII: ND2M416302) OX) WT9208) 4 (UNII: LU49E6626Q)		

.ALPHATO COPHEROL ACETATE, D- (UNII: A7E6112E4N)								
Packaging								
# Item Code	Package Description	Marketii	ıg Start Date	Ma	rketing End Date			
1 NDC:55316-267-08	236 mL in 1 BOTTLE, PUMP							
Marketing Information								
Marketing Category	Application Number or Monogra	ph Citation	Marketing Start I	Date	Marketing End Date			
OTC monograph not final	part333E		10/12/2011					
o re monograph not mar	partooph		10/12/2011					

Labeler - DZA BRANDS (090322194)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

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

Revised: 10/2011

DZA BRANDS