U INSTANT HAND SANITIZER CLASSIC 8OZ- ethyl alcohol gel
Universal Distribution Center LLC

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 percent

Purpose

Antimicrobial

Uses

Hand Sanitizer to help reduce bacteria on skin.

WARNINGS

Flammable. Keep away from fire or flame.

For external use only.

When using this product,

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water. Stop use and ask doctor if irritation or rash appears and lasts.

Keep out of reach for children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Place enough Product in your palm to thoroughly spread on both hands and rub into the skin until dry.
- Children under 6 years of age should be supervised when using this product.

Other Information:

- Store below 106 Fahrenheit (41 Degree Celsius)
- May discolor certain fabrics or surfaces.

INACTIVE INGREDIENTS:

Water (Aqua), Aloe Barbadensis Leaf Juice, Carbomer, Fragrance, Glycerin, Propylene Glycol, Triethanolamine, Tocopheryl Acetate.

U INSTANT HAND SANITIZER CLASSIC 8OZ

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-205
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**
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER 934 (UNII: Z135WT9208)	
TRIETHANOLAMINE BENZOATE (UNII: M3EN4GC19W)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52000-205-01	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

Labeler - Universal Distribution Center LLC (019180459)

Registrant - Universal Distribution Center LLC (019180459)

Revised: 9/2020 Universal Distribution Center LLC