

UZO HAND SANITIZER- hand sanitizer liquid
ANTHOS GROUP, INC, THE

79740-005 □□□□
UZO Hand Sanitizer

Active Ingredient(s)

Alcohol 70% v/v

Purpose

Antiseptic

Use

Hand sanitizer to help reduce bacteria that potentially can cause disease. for use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use
in children less than 2 months of age
on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs.
These may be signs of a serious condition.

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

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Directions

Place enough Product on hands to cover all surfaces. Rub hands together until dry. Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

Store between 15-20C(59-86F)

Avoid freezing and excessive heat about 40C(104F)


Inactive ingredients


water, Glycerin, Carbomer, Aminomethyl Propanol, Parfum, Aloe Leaf Extract

Package Label - Principal Display Panel

Drug Facts	
Active Ingredient(s) <i>Alcohol 70% v/v</i>	Purpose <i>Antiseptic</i>
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Inactive Ingredients Water, Glycerin, Carbomer, Aminomethyl Propanol, Parfum, Aloe Leaf Extract	

Distributed by UZO
www.uzo.la

 CITRUS
240 mL (8 Fl.oz)



UZO HAND SANITIZER

hand sanitizer liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79740-005
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
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
METHYL BENZOATE (UNII: 6618K1VJ9T)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79740-005-01	240 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	03/30/2020	

Labeler - ANTHOS GROUP, INC, THE (117511051)**Establishment**

Name	Address	ID/FEI	Business Operations
ANTHOS GROUP, INC, THE		117511051	manufacture(79740-005)

Revised: 2/2024

ANTHOS GROUP, INC, THE