INSTANT ANTISEPTIC HAND SANITIZER- alcohol solution
United Laboratories Inc.

Instant Antiseptic Hand Sanitizer 63998-364

Drug Facts

Active ingredient

Ethyl Alcohol 70%v/v

Purpose

Sanitizer

Uses

For hand sanitizing to decrease bacteria on the skin. Recommended for repeated use.

Warnings

Flammable. Keep away from fire or flame.

For external use only

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, or if condition persists for more than 72 hours.

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

Directions

Apply gel to dry hands and rub thoroughly until hands are dry. Do not rinse or wipe gel off. Should solution enter eyes, rinse immediately with water.

Other Information

Store below 110°F (43°C). May discolor certain fabrics or surfaces.

Inactive ingredients

Water, Propylene Glycol, PEG-33 (and) PEG-8 Dimethicone (and) PEG-14, Triethanolamine, Carbopol and Fragrance.

PRINCIPAL DISPLAY PANEL - 750 ml Bottle Label

UNITED
LABORATORIES

United 364

CONTACT PLUS

Instant Antiseptic
Hand Sanitizer

Kills 99.99% of common germs that can cause disease. Evaporates quickly.

750 ml (25 fl. oz.)

Made in USA 1018

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INSTANT ANTISEPTIC HAND SANITIZER

alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63998-364
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)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYETHYLENE GLYCOL 1600 (UNII: 1212Z7S33A)	
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
POLYETHYLENE GLYCOL 700 (UNII: 762678AC5R)	
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63998-364-01	750 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/2019	

Labeler - United Laboratories Inc. (001759737)

Establishment

Name	Address	ID/FEI	Business Operations
Woodbine Products Company		004220323	manufacture(63998-364)

Revised: 11/2024 United Laboratories Inc.