

LIQUID HAND SANITIZER- alcohol solution**Absara Cosmetics S.A.P.I DE C.V.**

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.

Liquid Hand Sanitizer**Active Ingredient(s)**

Alcohol 80% v/v. Purpose: Antimicrobial

Purpose

Antiseptic, Hand Sanitizer

Use

Hand sanitizer to help reduce bacteria on the skin.

Warnings

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

Children under 6yrs old to use under adult supervision.

In case of irritation, discontinue use and consult your doctor.

Do not ingest, if swallowed seek medical help. Avoid contact with eyes. Keep out of reach of children.

Directions

Apply the Gel on the hands and rub gently for 20 seconds until the product evaporates.

Other information

Store below 106 F (41 C). May discolor certain fabrics or surfaces.

Inactive ingredients

demineralized water, glycerin, dmdm hydantoin, aloe barbadensis extract, iodopropynyl butylcarbamate, butylene glycol.

Package Label - Principal Display Panel

CITRIC HINT

vlanc
+piür

HAND SANITIZER

Liquid

RINSE FREE

WITH
**& ALOE VERA
& GLYCERIN**

FORMULATED WITH

80%

OF ALCOHOL

DRUG FACTS

Active ingredient

Ethyl alcohol 80% v/v

Purpose

Antimicrobial

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Warnings

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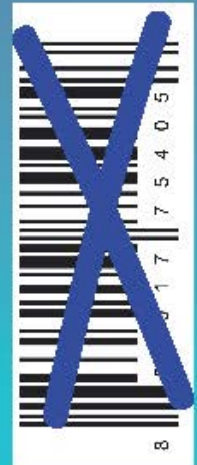
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MANUFACTURED IN MEXICO. Distributed by SHASON INC. VERNON, CA. 90058.

1 gal (3.78 L)

LIQUID HAND SANITIZER

alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70956-500
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 L in 100 L

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70956-500-01	3.78 L in 1 BOTTLE; 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 - Absara Cosmetics S.A.P.I DE C.V. (816161236)**Establishment**

Name	Address	ID/FEI	Business Operations
Absara Cosmetics S.A.P.I DE C.V.		816161236	manufacture(70956-500)

Revised: 5/2020

Absara Cosmetics S.A.P.I DE C.V.