

AZ HAND SANITIZER- az hand sanitizer spray

Ageless Zen Inc.

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.

Ageless Zen

AZ Hand Sanitizer

Active Ingredient

Ethyl alcohol 80% v/v. Purpose: Antiseptic

Benzalkonium Chloride 0.14% v/v. Purpose: Antiseptic

Purpose

Alcohol Antiseptic Spray

Uses

Hand Sanitizer to help reduce bacteria that potentially can cause disease

Warnings

For external use only: hands.

Flammable, keep away from fire or flame.

When using this product

Keep out of eyes. In case of contact with eyes, flush thoroughly with water. Avoid contact with broken skin. Do not inhale or ingest

Stop use and ask a doctor if

Irritation and redness develop 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

Spray into palm and rub hands together until dry. For children under 6 years of age, use only under adult supervision. Not recommended for infants

Other information

Do not store above 105F. May discolor some fabrics. Harmful to wood finishes and plastics.

Inactive Ingredients

Distilled Water, Glycerin, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Lavandula Angustifolia (Lavender) Oil, Citrus Medica Limonum (Lemon) Peel Oil

Ageless Zen AZ Hand Sanitizer

Alcohol Antiseptic Spray

Moisturizes hands with natural oils

80% Alcohol

0.14% Benzalkonium Chloride

200ML/ 6.76 FL.OZ

Made in the USA

NDC: 90080-200-01



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AZ HAND SANITIZER

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:90080-200
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 200 mL
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.14 in 200 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
LEMON OIL (UNII: I9GRO824LL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:90080-200-01	200 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/17/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/17/2020	

Labeler - Ageless Zen Inc. (117375334)**Registrant** - Ageless Zen Inc. (117375334)**Establishment**

Name	Address	ID/FEI	Business Operations
Ageless Zen Inc.		117375334	manufacture(90080-200)

Revised: 11/2020

Ageless Zen Inc.