

OCEAN BREEZE SCENTED ANTIBACTERIAL HAND SANITIZER- alcohol gel
Fourstar Group USA, Inc.

Ocean Breeze Scented Antibacterial Hand Sanitizer Gel

Drug Facts

Active ingredient

Ethyl Alcohol, 70% v/v

Purpose

Antimicrobial

Uses

hand sanitizer to help reduce bacteria on skin

Warnings

For external use only.

Flammable, keep away from fire or flame.

Do not use

in or near the eyes In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if

irritation, excessive redness or rash develops.

Keep out of reach of children.

Supervise children under 6 years of age when using this product to avoid swallowing. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Put a dime sized drop onto hands and rub together briskly until dry.

Other information

Store below 110°F (43°C)

Inactive ingredients

water, glycerin, propylene glycol, fragrance, carbomer, triethanolamine, blue 1

Package Labeling:

Drug Facts
Active ingredient
 Ethyl Alcohol, 70% v/v
Purpose Antimicrobial
Uses hand sanitizer to help reduce bacteria on skin ▼

PEEL UP LABEL FOR ADDITIONAL DRUG FACTS



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328424 2011
 DISTRIBUTED BY:
 IN U.S.A.: FOURSTAR GROUP USA, INC.
 925 GRANT STREET AKRON,
 OH 44311
 MADE IN CHINA

PEEL HERE

Drug Facts (continued)
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hand sanitizer
 Anti-Bacterial Hand Gel

OCEAN BREEZE SCENTED

1 fl oz / 29 mL

OCEAN BREEZE SCENTED ANTIBACTERIAL HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80684-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

		Marketing Start	Marketing End
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#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80684-005-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/07/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	11/07/2020		

Labeler - Fourstar Group USA, Inc. (140099503)

Revised: 10/2023

Fourstar Group USA, Inc.