

OZCARE ANTISEPTIC HAND SANITIZER- alcohol spray
Reed Holdings Australia, Pty. Ltd.

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.

Ozcare Antiseptic Hand Sanitizer Spray

Drug Facts

Active ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Use

- for hand washing to decrease bacteria on the skin.

Warnings

For external use only

Flammable, keep away from fire or flame

Do not use

- in the eyes

Stop use and ask a doctor if

- irritation and redness develop
- 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

- Wet hands thoroughly with product and allow to dry without wiping

Inactive ingredients

Aloe Vera Extract, Disodium EDTA, Eucalyptus Oil, Glycerin, Lavender Oil, Tocopherol Acetate, Triethanolamine, Water

Questions or Comments:

1-800-888-0776

Package Labeling:



Antiseptic Hand Sanitizer Spray 70% Ethyl Alcohol

No Water Needed



8.5 fl oz (250 mL)



B1063 EXP04042022



Drug Facts

Active ingredient	Purpose
Ethyl Alcohol 70%	Antiseptic
Use	
■ for hand washing to decrease bacteria on the skin.	
Warnings	
For external use only	
Flammable, keep away from fire or flame	
Do not use ■ in the eyes	
Stop use and ask a doctor if ■ irritation and redness develop	
■ 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	
■ Wet hands thoroughly with product and allow to dry without wiping	
Inactive ingredients	
Aloe Vera Extract, Disodium EDTA, Eucalyptus Oil, Glycerin, Lavender Oil, Tocopherol Acetate, Triethanolamine, Water	
Questions or Comments: 1-800-888-0776	

Manufactured By: Reed Holdings Australia, Pty Ltd
9 Whitfield Boulevard
Cranbourne West, Victoria 3977, Australia
Made in Australia

OZCARE ANTISEPTIC HAND SANITIZER

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75584-001
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
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
EDETATE DISODIUM ANHYDRO US (UNII: 8NLQ36F6MM)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75584-001-01	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/23/2020	

Labeler - Reed Holdings Australia, Pty. Ltd. (748340262)

Revised: 7/2020

Reed Holdings Australia, Pty. Ltd.