

ZYLAST ANTISEPTIC- ethyl alcohol liquid
Bocchi Laboratories 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.

Zylas t Antiseptic
Drug Facts

Active Ingredients

Ethyl alcohol 76%

Purpose

Antiseptic

Uses

- Handwash to decrease bacteria on the skin that potentially can cause disease.
- Recommended for repeated use.

Warnings

- For external use only.
- **Flammable. Keep away from flame.**
- Discontinue use if irritation and redness develops. Consult a doctor if condition persists for more than 72 hours.

Keep out of reach of children.

If swallowed, immediately call Poison Control Center or doctor.

Directions

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

Other Information

- Store at 20-25°C (68-77°F)

Inactive ingredients

Water, Polyaminopropyl Biguanide, Panthenol, Hydroxyethyl Ethylcellulose, Farnesol, PEG- 12 Dimethicone, Benzethonium Chloride.

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Package/Label Principal Display Panel

NDC 57702-464-00

Zylast

Broad Spectrum

Antimicrobial

Antiseptic

2 fl oz/ 59.1 mL



Zylast[®]

BROAD SPECTRUM
ANTIMICROBIAL
Antiseptic

FAST ACTING
PERSISTENT

**Kills 99.99%
of Germs**

Non-Irritating

2 fl oz | 59.1 mL



NDC: 57702-464-00

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*Manufactured for Innovative Biodefense Inc.
Lake Forest, CA • Made in USA
www.zylast.com • Patents Pending*



Package/Label Principal Display Panel

NDC 57702-464-03
Zylast
Broad Spectrum
Antimicrobial
Antiseptic
8 fl oz 236.6 mL



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Package/Label Principal Display Panel

NDC 57702-464-13
Zylast
Broad Spectrum
Antimicrobial

Antiseptic
1000mL (33.8oz)



BROAD SPECTRUM
ANTIMICROBIAL
Antiseptic

1000mL (33.8oz)

NDC: 57702-464-13

Kills 99.99% of germs

PERSISTENT

FAST ACTING, NON-IRRITATING



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ZYLAST ANTISEPTIC

ethyl alcohol liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:57702-464

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.76 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
FARNESOL (UNII: EB41QIU6JL)	
WATER (UNII: 059QF0KO0R)	
POLIHEXANIDE (UNII: 322U039GMF)	
PANTHENOL (UNII: WV9CM0O67Z)	
BENZETHONIUM CHLORIDE (UNII: PH41D05744)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57702-464-00	59.1 mL in 1 BOTTLE		
2	NDC:57702-464-03	236.6 mL in 1 BOTTLE		
3	NDC:57702-464-13	1000 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part333E	06/01/2011	

Labeler - Bocchi Laboratories Inc. (078376306)

Revised: 3/2014

Bocchi Laboratories Inc.