

**B PURE HAND SANITIZER EUCALYPTUS SCENTED- alcohol liquid
FOURSTAR GROUP USA, INC**

b.pure Hand Sanitizer Eucalyptus Scented

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.

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, acrylates/c10-30 alkyl acrylate

crosspolymer, aminomethyl propanol, yellow 6, yellow 5

Package Labeling:



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MANUFACTURED FOR:
FOURSTAR GROUP USA, INC.
925 GRANT ST., AKRON, OH 44311 USA
MADE IN CHINA

\$1.25



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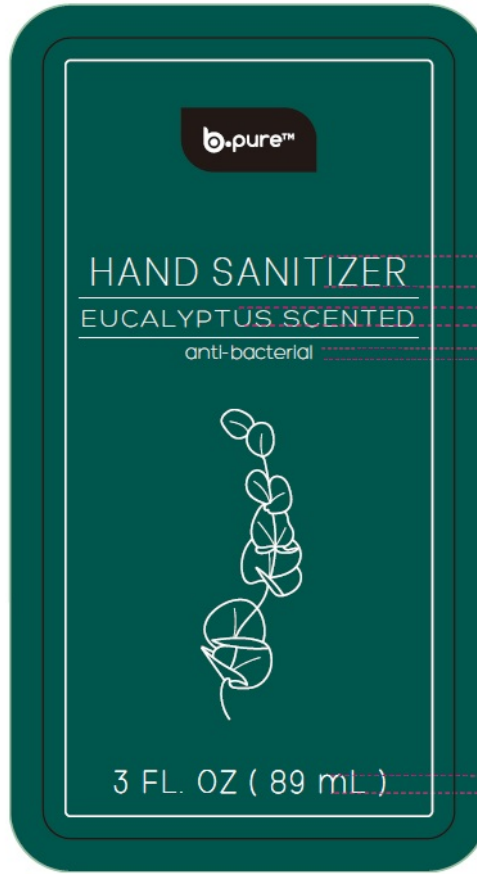
LOT

EXP



3 FL. OZ (89 mL)

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MANUFACTURED FOR: FOURSTAR GROUP USA, INC. 925 GRANT ST., AKRON, OH 44311 USA MADE IN CHINA LOT: EXP:	

B PURE HAND SANITIZER EUCALYPTUS SCENTED

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80684-115
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80684-115-00	1 in 1 BOX	08/25/2024	
1		89 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/25/2024	

Labeler - FOURSTAR GROUP USA, INC (140099503)

Revised: 6/2024

FOURSTAR GROUP USA, INC