

**B-PURE HAND SANITIZER COCONUT VANILLA SCENTED- alcohol liquid**  
**Fourstar Group USA, Inc.**

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**B-Pure Hand Sanitizer Spray, Coconut Vanilla 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***

Spray into hands and rub together briskly until dry.

***Other information***

Store below 110°F (43°C)

***Inactive ingredients***

water, fragrance, glycerin, cocos nucifera (coconut) oil, benzyl salicylate, hexyl cinnamal,

coumarin

## Package Labeling:

NDC 00000-000-00

**b.pure®**

COCONUT VANILLA SCENTED  
**HAND  
SANITIZER**  
SPRAY

Infused with Coconut Oil



**2 FL. OZ (59 mL)**

**Drug Facts**

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

LOT:    EXP:



0 49696 19887 5

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LOT:  
EXP:



Infused with  
Coconut Oil



**0.33 FL. OZ**  
**(10 mL)**

## B-PURE HAND SANITIZER COCONUT VANILLA SCENTED

alcohol liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:80684-178
<b>Route of Administration</b>	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
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>COCONUT OIL</b> (UNII: Q9L0O73W7L)	
<b>BENZYL SALICYLATE</b> (UNII: WAO5MNMK9TU)	
<b>.ALPHA.-HEXYLCINNAMALDEHYDE</b> (UNII: 7X6O37OK2I)	
<b>COUMARIN</b> (UNII: A4VZ22K1WT)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80684-178-01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/07/2025	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/07/2025	

**Labeler** - Fourstar Group USA, Inc. (140099503)**Registrant** - Zhejiang Meimi Technology Co., Ltd. (413668440)

Revised: 12/2025

Fourstar Group USA, Inc.