

**P B PURE ANTIBACTERIAL HAND SANITIZER SWEET PEA SCENTED- alcohol gel
FOURSTAR GROUP USA, INC.**

P b pure Antibacterial Hand Sanitizer Gel Sweet Pea Scented

Drug Facts

Active ingredient

Ethyl Alcohol, 65% 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, carbomer, triethanolamine, blue 1, red 33

Package Labeling:

Drug Facts

Active ingredient
Ethyl Alcohol, 65% v/v

Purpose Antimicrobial

Uses hand sanitizer to help reduce bacteria on skin ▼

PEEL UP LABEL FOR
ADDITIONAL DRUG FACTS

SKU#998890
DISTRIBUTED BY:
MIDWOOD BRANDS, LLC
500 VOLVO PARKWAY
CHESAPEAKE, VA 23320 USA
MADE IN CHINA
MIDWOODBRANDSLLC0496965030990920

LOT: EXP:

PEEL
HERE

Drug Facts (continued)

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Drug Facts (continued)

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 **b.pure™**

ANTI-BACTERIAL
HAND SANITIZER GEL



Sweet Pea Scented

1 fl oz (29 mL)

P B PURE ANTIBACTERIAL HAND SANITIZER SWEET PEA SCENTED

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80684-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.65 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)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:80684-002-01	29 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/24/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)		09/24/2020	

Labeler - FOURSTAR GROUP USA, INC. (140099503)

Revised: 10/2023

FOURSTAR GROUP USA, INC.