

**B-PURE MOISTURIZING HAND SANITIZER BLOSSOM DREAM SCENTED LIMITED EDITION- alcohol liquid**  
**Fourstar Group USA, Inc.**

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**B-Pure Moisturizing Hand Sanitizer Blossom Dream Scented Limited Edition**

***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 product in hands and rub together.

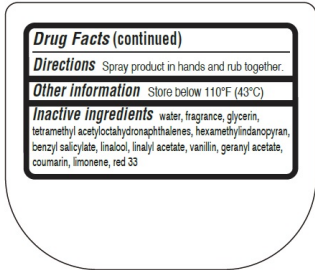
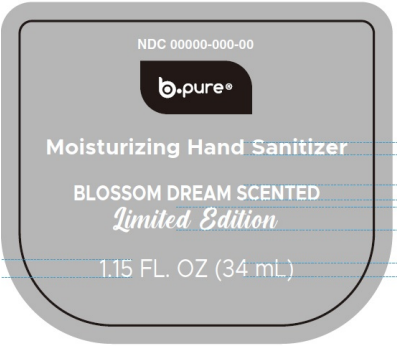
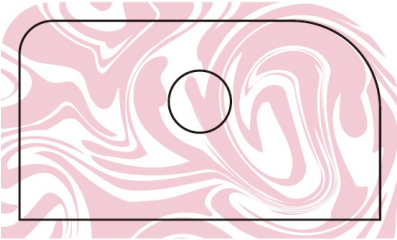
***Other information***

Store below 110°F (43°C)

***Inactive ingredients***

water, fragrance, glycerin, tetramethyl acetyloctahydronaphthalenes, hexamethylindanopyran, benzyl salicylate, linalool, linalyl acetate, vanillin, geranyl acetate, coumarin, limonene, red 33

Package Labeling:



B-PURE MOISTURIZING HAND SANITIZER BLOSSOM DREAM SCENTED LIMITED EDITION

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80684-189
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)			
TETRAMETHYL ACETYLOCTAHYDRONAPHTHALENES (UNII: 2JU6ZH6GRE)			
HEXAMETHYLINDANOPYRAN (UNII: 14170060AT)			
BENZYL SALICYLATE (UNII: WAO5MKN9TU)			
LINALOOL (UNII: D81QY6I88E)			
LINALYL ACETATE (UNII: 5K47SSQ51G)			
VANILLIN (UNII: CHI530446X)			

GERANYL ACETATE (UNII: 3W81YG7P9R)				
COUMARIN (UNII: A4VZ22K1WT)				
LIMONENE, (+)- (UNII: GFD7C86Q1W)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80684-189-01	34 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2025	
<b>Marketing Information</b>				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		505G(a)(3)	11/20/2025	

**Labeler** - Fourstar Group USA, Inc. (140099503)

**Registrant** - Fourstar Group USA, Inc. (140099503)