B-PURE MOISTURIZING HAND SANITIZER TWILIGHT DANCE SCENTED- alcohol liquid
Fourstar Group USA, Inc.

B-Pure Moisturizing Hand Sanitizer Twilight Dance 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 product in hands and rub together.

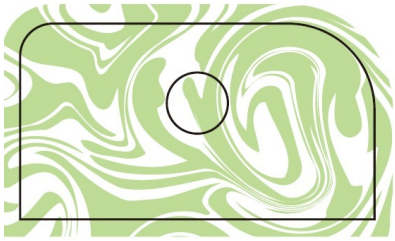
Other information

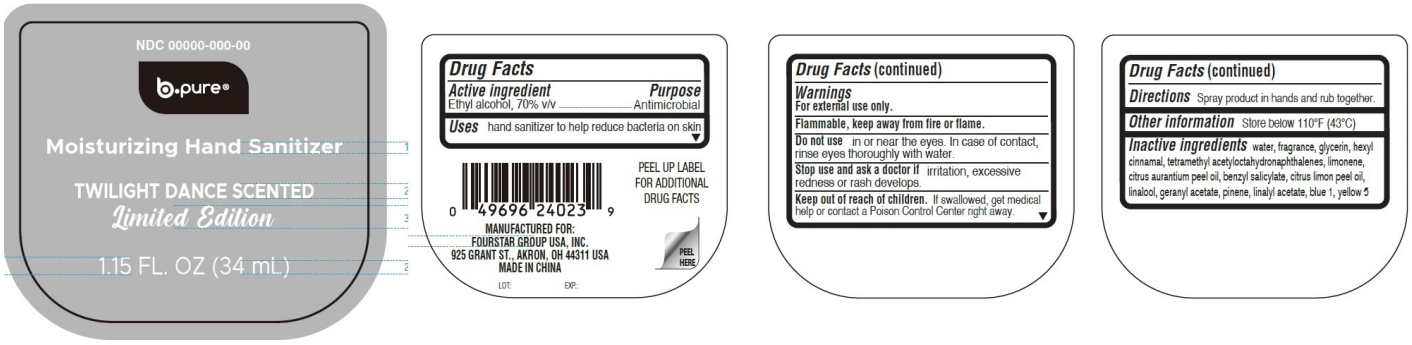
Store below 110°F (43°C)

Inactive ingredients

water, fragrance, glycerin, hexyl cinnamal, tetramethyl acetyloctahydronaphthalenes, limonene, citrus aurantium peel oil, benzyl salicylate, citrus limon peel oil, linalool, geranyl acetate, pinene, linalyl acetate, blue 1, yellow 5

Package Labeling:





B-PURE MOISTURIZING HAND SANITIZER TWILIGHT DANCE SCENTED

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80684-188
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)	
HEXYL CINNAMAL (UNII: 7X6O37OK2I)	
TETRAMETHYL ACETYLOCTAHYDRONAPHTHALENES (UNII: 2JU6ZH6GRE)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
BITTER ORANGE OIL (UNII: 9TLV70SV6I)	
BENZYL SALICYLATE (UNII: WAO5MNK9TU)	
LEMON OIL, COLD PRESSED (UNII: I9GRO824LL)	

LINALOOL, (+/-)- (UNII: D81QY6I88E)	
GERANYL ACETATE (UNII: 3W81YG7P9R)	
PINENE (UNII: 996299PUKB)	
LINALYL ACETATE (UNII: 5K47SSQ51G)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80684-188-01	34 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/20/2025	

Marketing Information

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)

Revised: 12/2025 Fourstar Group USA, Inc.