

PUMPKIN SPICE SCENTED HAND SANITIZER- alcohol gel
Fourstar Group USA, Inc.

PUMPKIN SPICE SCENTED HAND SANITIZER

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.

Supervise children under 6 years of age when using this product to avoid swallowing. 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.

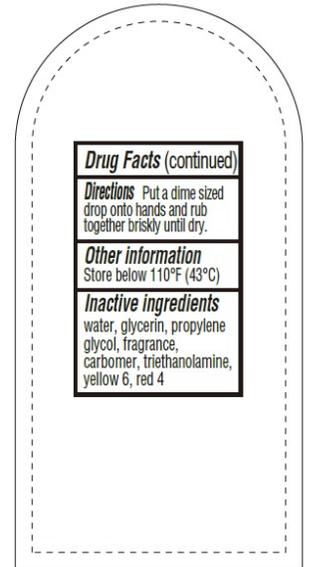
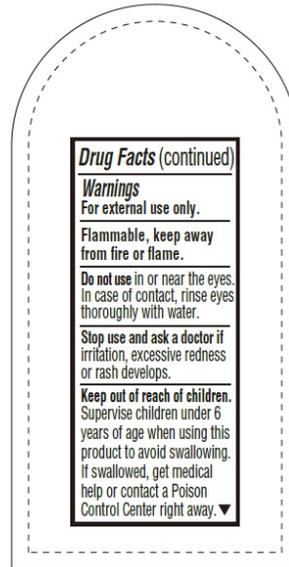
Other information

Store below 110°F (43°C)

Inactive ingredients

water, glycerin, propylene glycol, fragrance, carbomer, triethanolamine, yellow 6, red 4

Package Labeling:59ml



Package Labeling:236ml

94- 820935

\$3.49

PUMPKIN SPICE SCENTED



HAND SANITIZER

ANTI-BACTERIAL HAND GEL

8 FL. OZ (236 mL)

3.4mm

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820935 2204 MANUFACTURED FOR:
FOURSTAR GROUP USA, INC.
925 GRANT ST., AKRON, OH 44311 USA
MADE IN CHINA

LOT: EXP:

PUMPKIN SPICE SCENTED HAND SANITIZER 8OZ (236mL)



PUMPKIN SPICE SCENTED HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80684-065
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)	

CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)

TROLAMINE (UNII: 9O3K93S3TK)

FD&C YELLOW NO. 6 (UNII: H77VEI93A8)

FD&C RED NO. 4 (UNII: X3W0AM1JLX)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80684-065-02	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2022	
2	NDC:80684-065-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/10/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/10/2022	

Labeler - Fourstar Group USA, Inc. (140099503)

Revised: 10/2023

Fourstar Group USA, Inc.