

SANITIZING HAND- alcohol spray
BELLA BRANDS INC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Sanitizing Hand

Drug Facts:

Active Ingredient

Ethyl Alcohol 70% V/V

Purpose

Antimicrobial Agent

Use

for cleaning hands and decreasing bacteria on the skin.

Warnings

- For external use only.
- Do not use if you are allergic to any of the ingredients.
- When using this product, do not get into eyes. If contact occurs, rinse thoroughly with water.
- Stop use and consult a doctor if irritation or rash develops and continues for more than 72 hours.
- Keep out of reach of children. If swallowed, get medical help or visit a Poison Control Center right away.

Inactive Ingredients

Water, Glycerin, Aloe Vera Barbadensis Leaf Juice¹, Chamomilla Recutita (Matricaria) Flower Extract¹, Camelia Sinensis Leaf Extract¹, Vitis Vinifera (Grapeseed) Extract¹, Helianthus Annuus (Sunflower) Seed Oil¹, Fragrance (Naturally Derived)

¹ Organic Ingredient

Directions For Use

- Squeeze dispensing sprayer to release product into hands, and rub thoroughly around both sides of hands and between fingers. Do not wipe off. Let air dry.
- For adults and children 2 years and over.
- For children under 2 years of age, consult a doctor before use.

PRINCIPAL DISPLAY PANEL - 400 ml Bottle Label

PURIGEN™

Sanitizing

Hand Spray

- Kills 99.9% of Germs
- Safe for Sensitive Skin
- 70% Alcohol Formula

Paraben Free

No Artificial Fragrance

Made with Organic Ingredients

13.5 fl oz (400 ml)

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Sanitizing Hand Spray

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Pure. Clean. Healthy.

Purigen Sanitizing Hand Spray is convenient, effective and moisturizing. Formula includes organic aloe vera, grapeseed extract and sunflower seed oil to keep skin soft and moisturized.

Also available:
Purigen Hand Sanitizing Lotion, Gel, Soap and Wipes.

Bella Brands Inc.
Aliso Viejo, California
MADE IN THE USA
www.PurigenCare.com

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SANITIZING HAND

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80994-003
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)	
Chamomile (UNII: FGL3685T2X)	
Green Tea Leaf (UNII: W2ZU1RY8B0)	

Vitis Vinifera Seed (UNII: C34U15ICXA)				
Sunflower Oil (UNII: 3W1JG795YI)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80994-003-01	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
2	NDC:80994-003-02	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	05/01/2020		

Labeler - BELLA BRANDS INC (034908755)

Establishment			
Name	Address	ID/FEI	Business Operations
BELLA BRANDS INC		034908755	MANUFACTURE(80994-003)

Revised: 11/2020

BELLA BRANDS INC