HAND CARE SET SPICED VANILLA- alcohol
Base4 Ventures, LLC

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.

Hand Care Set Spiced Vanilla

Drug Facts

Active ingredient

Alcohol Denat. 66%

Purpose

Antiseptic

Use

- For handwashing to decrease bacteria on the skin

Warnings

For external use only

- **Flammable, Keep away from fire & flame**
- **Does not contain grain alcohol; do not drink. If taken internally will produce severe gastric disturbances**

When using this product

- Avoid the eyes and mucous membranes
- In the case of eyes or mucous membrane contact, rinse area thoroughly with water

Stop use and ask a doctor if

- Condition worsens
- Redness or irritation develops
- Condition persist for more than 3 days

Keep out of reach of children.

If swallowed, contact a doctor or Poison Control Center immediately.

Directions

- Rub dime sized amount between hands until dry
- Supervise children in the use of this product
- In the case of eye contact rinse eyes thoroughly with water

Other information

- Store below 105°F
- May discolor fabrics

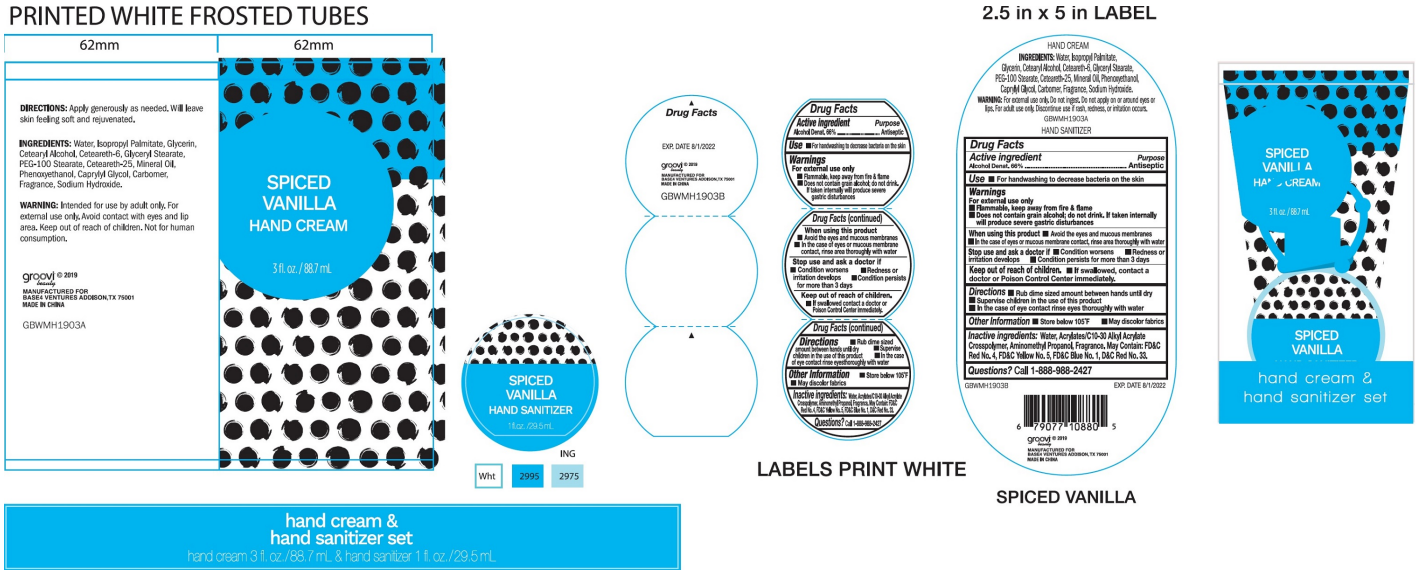
Inactive ingredients:

Water, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance. May Contain: FD&C Red No.4, FD&C Yellow No.5, FD&C Blue No.1, D&C Red No.33.

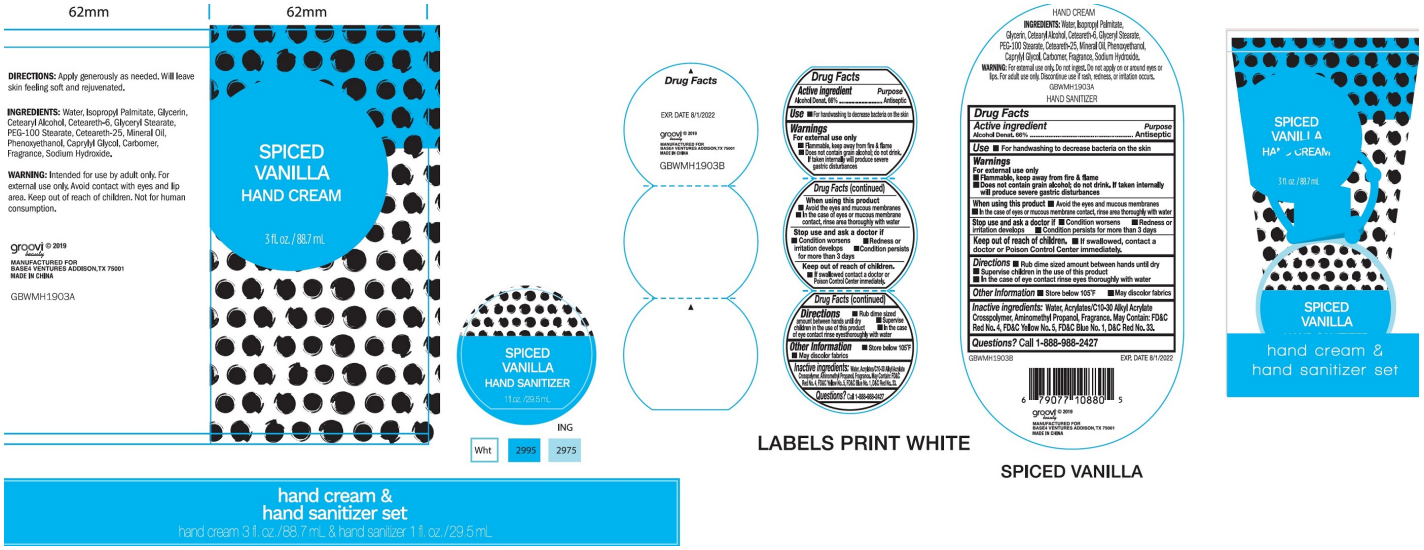
Questions?

Call 1-888-988-2427

Hand Care Set Spiced Vanilla (62651-023-00)



Hand Sanitizer Spiced Vanilla, 1 oz (62651-009-01)



HAND CARE SET SPICED VANILLA

alcohol kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62651-023

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62651-023-00	1 in 1 KIT	10/15/2019	

Quantity of Parts		
Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	29.5 mL

Part 1 of 1

HAND SANITIZER SPICED VANILLA

alcohol gel

Product Information	
Item Code (Source)	NDC:62651-009
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	660 mL in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
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:62651-009-01	29.5 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/15/2019	

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Labeler - Base4 Ventures, LLC (137316126)

Revised: 6/2019 Base4 Ventures, LLC