

**JUSTICE VANILLA LATTE SCENTED ANTI-BACTERIAL HAND SANITIZER- alcohol gel
Tween 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.

Justice VANILLA Latte Scented Anti-Bacterial Hand Sanitizer

Drug Facts

Active ingredient

Alcohol 62%

Purpose

Antiseptic

Use

- For hand washing to decrease bacteria on the skin.
- Recommended for repeated use.

Warnings

Flammable, keep away from fire or flame.

For external use only.

Do not use

in the eyes or apply over large areas of the body.

In case of deep or puncture wounds, animal bites, or serious burns, consult a doctor.

Use only as directed. Do not drink. If taken internally, it will produce serious gastric disturbances.

Stop use and ask a doctor if

Irritation persists for more than 72 hours.

Keep out of reach of children.

If swallowed get medical help or contact a Poison Control Center right away.

Directions

- Wet hands thoroughly with product and allow to dry without wiping.
- Supervise children in the use of this product.

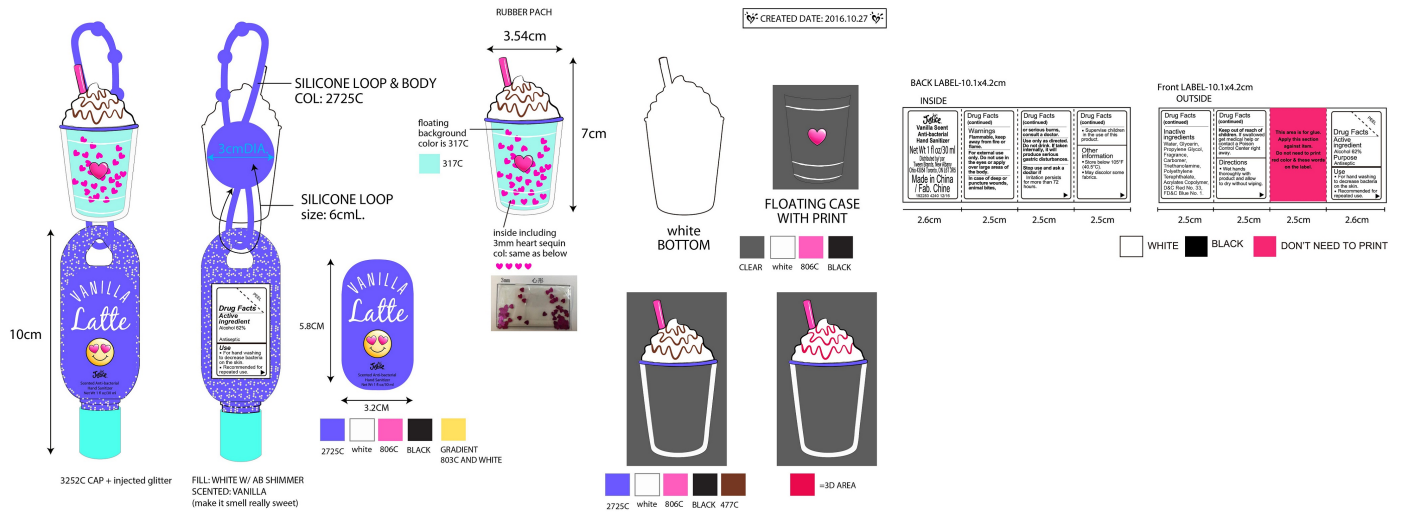
Other information

- Store below 105°F (40.5°C).
- May discolor some fabrics.

Inactive ingredients

Water, Glycerin, Propylene Glycol, Fragrance, Carbomer, Triethanolamine, Polyethylene Terephthalate, Acrylates Copolymer, D&C Red No. 33, FD&C Blue No. 1.

Package Labeling:



JUSTICE VANILLA LATTE SCENTED ANTI-BACTERIAL HAND SANITIZER
alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60637-203
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	620 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60637-203-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/06/2016	

Labeler - Tween Brands, Inc (965758188)

Revised: 12/2016

Tween Brands, Inc