JUSTICE VANILLA SCENT 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 Scent 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.

Supervise children in the use of this product.

Other Information

- Store beow 105 ⁰F (40.5 ⁰C)
- M ay discolor some fabrics.

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.

Inactive Ingredients

Water, Glycerin, Propylene Glycol, Fragrance, Carbomer, Triethanolamine, FD&C Red No. 33, FD&C Yellow No. 5

Product Labels







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Drug Facts

(continued)

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Drug Facts

(continued)

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Other information

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

Drug Facts

(continued)

Inactive ingredients

Water ,Glycerin , Propylene Glycol . Fragrance , Carbomer, Triethanolamine, D&C Red No. 33 (CI 17200). FD&C Yellow No. 5 (CI 19140).

Drug Facts

(continued)

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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.

This area is for glue. Apply this section against item. Do not need to print red color & these words on the label.



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alcohol gel

Product Information

HUMAN OTC DRUG NDC:60637-030 **Product Type** Item Code (Source)

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength **ALCOHOL** ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M) 620 mg in 1 mL

Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R) **GLYCERIN** (UNII: PDC6A3C0OX) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) TROLAMINE (UNII: 903K93S3TK) FD&C YELLOW NO. 5 (UNII: I753WB2F1M)

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:60637-030-	1 in 1 CARTON	08/18/2015		
1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333E	08/18/2015		

Labeler - Tween Brands Inc (965758188)

Revised: 10/2022 Tween Brands Inc