GLITTER RITZ ICED BERRY- alcohol gel
Landy International

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.

Active Ingredient ***Purpose***

Alcohol denat. 63% ..... Antiseptic

Uses

to decrease bacteria on the skin

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Hand Sanitizer

30 ml 1.0 fl.oz.

Warnings

For external use only.

Flammable.

Keep away from heat and flame.

When using this product

- Avoid contact with eyes. If contact occurs, flush eyes with water.
- Avoid contact with broken skin.

Stop use and ask a doctor if

irritation and redness develop and persist for more than 72 hours.

Directions

- Wet hands with product and allow to dry without wiping.
- Not recommended for infants.

Inactive Ingredients

alcohol denat., water, propylene glycol, acrylates/c10-30 alkyl acrylate crosspolymer, aminomethyl propanol, fragrance, lactose, cellulose, hydroxypropyl methylcellulose, jojoba esters, tocopheryl acetate, yellow 5 (CI 19140), red 4 (CI 14700)

Drug Facts(Continued)

Directions

- Wet hands with product and allow to dry without wiping.
- Not recommended for infants.

Other information

- Store at 68° to 77°F (20° to 25°c).
- Do not store above 105°F.
- May discolor some fabrics.
- Harmful to wood finishes and plastics.

Expired Date 05/2015

Tracking number: OF0503-01
FDA number: 545291775

Inactive ingredients

Alcohol Denat., Water, Propylene Glycol, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aminomethyl Propanol, Fragrance, Lactose, Cellulose, Hydroxypropyl Methylcellulose, Jojoba Esters, Tocopheryl Acetate, Red 33 (CI 17200), Red 4 (CI 14700).

PEEL UP TO SEE ADDITIONAL DRUG FACTS

Walmart.com

MADE IN CHINA

Distributed by Wal-Mart Stores, Inc.
Bentonville, AR 72716

8 43736 02148 8

WSS15A010

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alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51706-512
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	63 in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
(C10-C30)ALKYL METHACRYLATE ESTER (UNII: XH2FQZ38D8)	
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)	
LACTOSE (UNII: J2B2A4N98G)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
HYDROLYZED JOJOBA ESTERS (ACID FORM) (UNII: UDR641JW8W)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51706-512-01	63 g 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	04/15/2015	

Labeler - Landy International (545291775)

Establishment

Name	Address	ID/FEI	Business Operations
Landy International		545291775	manufacture(51706-512)

Revised: 4/2015 Landy International