## BODYCOLOGY VANILLA CUPCAKE- ethyl alcohol gel Wal-Mart Stores, 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.

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#### Vanilla Cupcake Hand Sanitizer

Active Ingredient purpose
Ethyl Alcohol 70% Antiseptic

Uses: to decrease bacteria on the skin.

Keep out of reach of children.

Stop use and consult a doctor if irritation and redness develop and persist for more than 72 hours.

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.

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

Directions: Wet hands with product and allow to dry without wiping. For children under 6, use only under adult supervision. Not recommended for infants.

Inactive ingredients: Water (aqua, eau), fragrance (parfum), Propylene Glycol, Acrylates/C10-30 alkyl acrylate crosspolymer, triethanolamine, Gelatin, Tocopheryl Acetate, Aloe Barbadensis (Aloe Vera leaf juice), Ultramarines (CI77007)

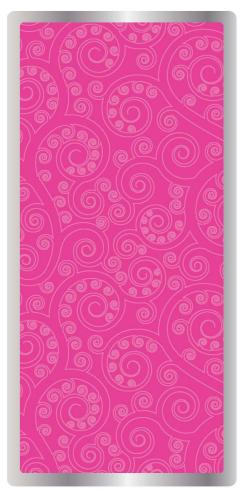
May contain: FD C Red No.4(CI 14700, Aka504), FD C Yellow No. 5 (CI 19140,Ki4), FD C Blue No.1 (CI 42090, Ao1), D C Red No 33(CI 17200,Aka227)



Bodycology Moisturizing Hand Sanitizer

30 mL 1 Fl Oz







#### **BODYCOLOGY VANILLA CUPCAKE**

ethyl alcohol gel

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-992		
Route of Administration	TOPICAL				

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL (To ml in 100 ml

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
PROPYLENE GLYCOL (UNII: 6 DC9 Q16 7 V3)		
CARBOMER INTERPOLYMER TYPE A (55000 MPA.S)) (UNII: 59TL3WG5CO)		
TROLAMINE (UNII: 9O3K93S3TK)		

GELATIN (UNII: 2G86QN327L)	
ALOE VERA LEAF (UNII: ZY81Z83H0 X)	
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:49035-992-02	1 in 1 BOTTLE				
1	NDC:49035-992-01	30 mL in 1 BOTTLE				

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	04/09/2011			

### Labeler - Wal-Mart Stores, Inc. (051957769)

Revised: 4/2011 Wal-Mart Stores, Inc.