CRANBERRY CAKE- antibacterial hand sanitizer gel Brands 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.

CRANBERRY CAKE ANTIBACTERIAL HAND SANITIZER

JACTIVE INGREDIENT: ETHYL ALCOHOL 62% PURPOSE ANTISEPTIC

ANTISEPTIC

USES: TO DECREASE BACTERIA ON THE SKIN AND CLEAN HANDS. RECOMMENDED FOR REPEATED USE

UWARNING: FOR EXTERNAL USE ONLY. FLAMMABLE, UKEEP AWAY UFROM FIRE OR FLAME.

DO NOT GET INTO EYES. IF CONTACT OCCURS, RINSE THROUGHLY WITH WATER.

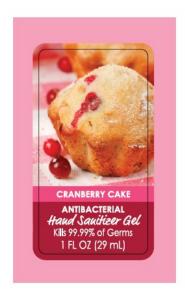
DISCONTINUE USE IF IRRITATION OR REDNESS DEVELOP. IF IRRITATION PERSISTS FOR MORE THEN 72 HOURS, CONSULT A DOCTOR.

SUPERVISE CHILDREN IN THE USE OF THIS PRODUCT.

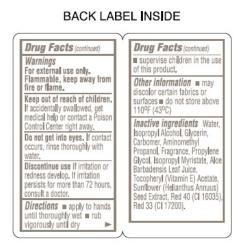
OTHER INFORMATIONMAY DISCOLOR CERTAIN FEBRICS OR SURFACES.

INACTIVE INGREDIENTS: WATER, PISOPROPYL ALCOHOL, GLYCERIN, CARBOMER, AMINOMETHYL PROPANOL, PROPYLENE GLYCOL, ISOPROPYL MYRISTATE, ALOE BARBADENSIS LEAF JUICE, TOCOPHEROL ACETATE, SUNFLOWER (HELIANTHUS ANNUUS) SEED EXTRACT, MAY CONTAIN: FD&C BLUE NO. 1(CI 42090), FD&C RED NO. 40(CI 16035), FD&C YELLOW NO. 5 (CI 19140)

CRANBERRY CAKE ANTIBACTERIAL HAND SANITIZER







CRANBERRY CAKE

antibacterial hand sanitizer gel

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50157-112		
Route of Administration	TOPICAL				

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	620 mL in 1000 mL			

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
ISOPROPYL ALCOHOL (UNII: ND2M416302)				
GLYCERIN (UNII: PDC6A3C0OX)				
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)				
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
ISOPROPYL MYRISTATE (UNII: 0 RE8 K4LNJS)				
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)				
.ALPHATO COPHEROL ACETATE (UNII: 9E8X80D2L0)				
HELIANTHUS ANNUUS SEED WAX (UNII: 42DG15CHXV)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:50157- 112-01	29 mL in 1 BOTTLE, PLASTIC; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)	10/01/2015	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	10/01/2015				

Labeler - Brands International (243748238)

Establishment					
Name	Address	ID/FEI	Business Operations		
Brands International		243748238	manufacture(50157-112)		

Revised: 10/2015 Brands International