# WINTER VANILLA HAND SANITIZER NONE- winter vanilla hand sanitizer liquid Unique Holding Group 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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#### **Drug Facts**

Put a thumb size amount in your palm and rub your hand briskly until dry.

Do not store in temperatures over 118F children under six years of age should be supervise while using this product. May discolor certain fabrics.

Aloe Barbadensis Gel, Carbomer, Deionized water, Propylene glycol, Glycerin, Triethanolamine, Vitamin E, Fragance, may contain DC red 33, FDC blue 1

To decrease bacteria on the skin that could cause disease. Recommended for repeated use. labelpicture





Drug Facts		Drug Facts (continued)	
Active ingredient Purpose Ethyl Alcohol 62%Sanitizer  Uses ■ to decrease bacteria on		Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a	
the skin that could cause recommended for rep	disease.	Poison Control Center immediately. <b>Directions</b>	
Warnings For external use only-hands. Use only as directed. Excessive use or prolonged exposure may cause irritation to skin. Discontinue use if rash, redness, or itching occurs. Flammable. Keep away from heat and flame.  When using this product with product use if rash, redness, or itching occurs. Flammable is seen and flame.  When using this product leading the with water and call a doctor, and with water and call a doctor. I would contact with broken skin.  Stop use and ask a doctor irritation or redness develops.		■ put a thumb size amount in your palm and rub hands together briskly until dry.  Other information ■ do not store in temperatures over 118° F. ■ dividere under 6 years of a e should be supervised while using this product. ■ may discolor certain fabrics.  Inactive ingredients Aloe Barbadensis Gel, Carbomer, Deionized Water, Fragrance, Glycerin, Propylene Glycol, Triethanolamine, and Vitamin E.	

#### WINTER VANILLA HAND SANITIZER NONE

winter vanilla hand sanitizer liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:25225-011(NDC:None)	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 g in 100 g		

Inactive Ingredients		
Ingredient Name	Strength	
Propylene Glycol (UNII: 6DC9Q167V3)	0.5 g in 100 g	

Glycerin (UNII: PDC6 A3C0 OX)	1 g in 100 g
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	0.01 g in 100 g
CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	0.33 g in 100 g
Vitamin E (UNII: H4N855PNZ1)	0.01 g in 100 g
TROLAMINE (UNII: 9 O 3 K 9 3 S 3 T K)	0.35 g in 100 g
WATER (UNII: 059QF0KO0R)	35.6 g in 100 g

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	VANILLA (0.2% w/w)	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:25225-011-01	59 g in 1 BOTTLE, PLASTIC		
2	NDC:25225-011-02	30 g in 1 BOTTLE, PLASTIC		
3	NDC:25225-011-03	237 g in 1 BOTTLE, PLASTIC		
4	NDC:25225-011-04	500 g in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333	11/10/2009	

## Labeler - Unique Holding Group Inc (529047265)

### Registrant - Unique Holding Group Inc (529047265)

Establishment				
Name	Address	ID/FEI	Business Operations	
Unique Holding Group Inc		529047265	manufacture	

Revised: 11/2009 Unique Holding Group Inc