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.

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

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: PDC6A3C0OX)	1 g in 100 g
ALOE VERA LEAF (UNII: ZY81Z83H0X)	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: 9O3K93S3TK)	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	Marketing Start Date	Marketing End Date
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