AQUA SUNSET HAND SANITIZER - alcohol 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

Active Ingredient

Ethyl Alcohol 62%

Purpose:

Sanitizer

Uses:

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

Warnings:

For external use only-hands.
Use only as directed.
Excessive use or prolonged exposure may cause irritation to the skin.
Discontinue use if irritation, redness, or itching occurs.
Flammable. Keep away from heat and flame

When Using This Product:

Keep out of eyes. In case of contact with eyes, flush immediately with water and call a doctor.

Avoid contact with broken skin

Stop Use And Ask A Doctor if irritation or redness develops

Keep Out of Reach of Children:

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

Directions:

Put a thumb size amount in your palm and rub hands together briskly until dry

Other Information

Do not store in temperatures over 118F

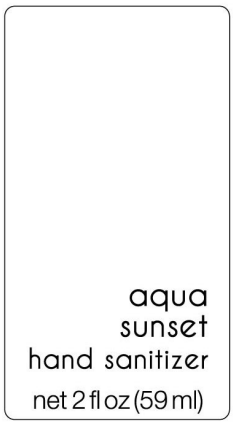
Children under six years of age should be supervised while using this product.

May discolor certain fabrics

Inactive Ingredients

aloe barbadensis gel, carbomer, deionized water, Fragrance, glycerin, propylene glycol, D and C Blue No. 1, triethanolamine, and vitamin E, FD and C Yellow No. 5

lab



AQUA SUNSET HAND SANITIZER

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:25225-023(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**
Water (UNII: 059QF0KO0R)	35.5998 g in 100 g
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
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	0.01 g in 100 g
TROLAMINE (UNII: 9O3K93S3TK)	0.35 g in 100 g
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	0.0001 g in 100 g
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	0.0001 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:25225-023-01	28 g in 1 BOTTLE, PLASTIC		
2	NDC:25225-023-02	59 g in 1 BOTTLE, PLASTIC		
3	NDC:25225-023-04	237 g in 1 BOTTLE, PLASTIC		
4	NDC:25225-023-05	500 g in 1 BOTTLE, PLASTIC		
5	NDC:25225-023-03	222 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	11/13/2010	

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: 11/2010 Unique Holding Group Inc