SUNKISSED HONEYDEW HAND SANITIZER - ethyl alcohol liquid
Papermates, Inc. dba Noteworthy

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.

Sunkissed Honeydew Hand Sanitizer

Active Ingredient

Ethyl Alcohol 62%

Purpose

Sanitizer

Uses

To decrease 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 skin. Discontinue use if rash 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 immediately flush 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 6 years of age should be supervised while using this product.

may discolor certain fabrics.

Inactive Ingredients

Aloe barbadensis gel, blue 1, carbomer, deionized water, fragrance, glycerin, propylene glycol, triethanolamine, vitamin E, and yellow 5

Warning this item is for external use only. Do not ingest.

Use anytime, anyplace, without water

Feel refreshed without stickiness or residue

Manufactured for Noteworthy

2010 Noteworthy
Chatsworth, CA 91311

Made in China

Sunkissed Honeydew

Hand Sanitizer

net 2 fl oz (59ml)

SUNKISSED HONEYDEW HAND SANITIZER

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75997-021
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE (UNII: V5VD430YW9)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:75997-021-02	59 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/02/2011	

Labeler - Papermates, Inc. dba Noteworthy (038734620)

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