STRAWBERRY 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.

Strawberry 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, carbomer, deionized water, fragrance, glycerin, propylene glycol, red 33, 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

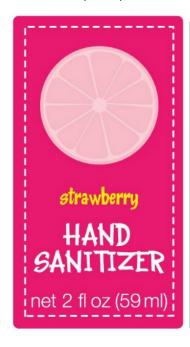
2010 Noteworthy Chatsworth, CA 91311

Made in China

Strawberry

Hand Sanitizer

net 2 fl oz (59ml)





TOPICAL

Drug Facts

Active ingredient Purpose Ethyl Alcohol 62%.....Sanitizer

Uses ■ to decrease bacteria on the skin that could cause disease. ■ recommended for repeated use.

Warnings For external use

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When using this product

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Drug Facts (continued)

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STRAWBERRY HAND SANITIZER

ethyl alcohol liquid

Route of Administration

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:75997-025

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: V5VD430 YW9)				
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
PROPYLENE GLYCOL (UNII: 6 DC9 Q167V3)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
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:75997-025-02	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)

Revised: 2/2011 Papermates, Inc. dba Noteworthy