

GUM DROP 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.

Gum Drop 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. recommended for repeated use.

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, red 33, triethanolamine and vitamin E

Gum Drop

Hand sanitizer

net .95fl oz (28ml)



GUM DROP HAND SANITIZER

ethyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75997-027
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: HBR47K3TBD)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75997-027-01	28 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/28/2011	

Labeler - Papermates, Inc. dba Noteworthy (038734620)

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