

HILLYARD INSTANT HAND SANITIZER- ethyl alcohol gel
Hillyard GMP

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.

Hillyard Instant Hand Sanitizer Gel

Active Ingredients

Ethyl Alcohol 62.0%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use.

Warnings

For External Use Only.

Flammable. Keep away from fire or flame.

When using this product

avoid contact with eyes. In case of eye contact, flush eyes thoroughly with water.

Stop use and ask a doctor if

irritation or redness develops, or if condition persists for more than 72 hours.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Pump small amount into palm of hand.
- Rub thoroughly over all surfaces of both hands.
- Rub hands together briskly until dry.
- Children under 6 years of age should be supervised when using this product.

Inactive Ingredients

Water, Acrylates/C10-C30 Alkyl Acrylate Crosspolymer, Tetrahydroxypropyl Ethylenediamine, Glycerine, Fragrance, Citric Acid, Vitamin E, Aloe Barbadensis Leaf Juice.

Package/Label Principal Display Panel

Hillyard
**Instant Hand
Sanitizer Gel**

HIL0040203

affinity

Place cartridge into dispenser, label facing back, and gently push pump into receiver until it clicks into place.

To remove the cartridge and pump, rotate the green ring clockwise and pull the pump out.

HILLYARD INDUSTRIES

PO Box 909,

St. Joseph, MO

64502-0909 U.S.A.

Telephone: 816-233-1321

www.hillyard.com

NET CONTENTS 33.8 FL. OZ. 1L

HIL0040203

0216

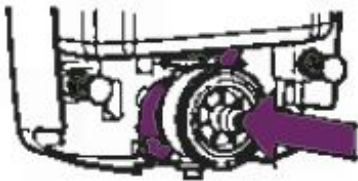
PEEL HERE

HILLYARD

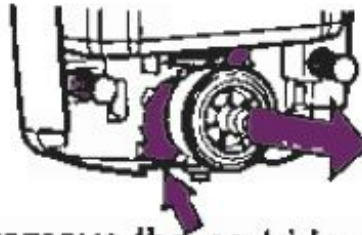
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Label

HILLYARD INSTANT HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76402-402
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL
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Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
EDETOL (UNII: Q4R969U9FR)	
GLYCERIN (UNII: PDC6A3C0OX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
.ALPHA.-TOCOPHEROL, D- (UNII: N9PR3490H9)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76402-402-06	3785 mL in 1 JUG; Type 0: Not a Combination Product	05/23/2012	
2	NDC:76402-402-03	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/21/2012	
3	NDC:76402-402-02	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/27/2013	
4	NDC:76402-402-83	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/27/2013	
5	NDC:76402-402-44	119 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/21/2012	

Labeler - Hillyard GMP (969081483)

Establishment

Name	Address	ID/FEI	Business Operations
Hillyard GMP		969081483	MANUFACTURE(76402-402)

Revised: 12/2017

Hillyard GMP