

EHS7000- ehs7000 gel**EPIC Chemistry LLC**

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.

8 OZ**Active Ingredient**

Ethyl Alcohol 70.05 v/v% Antimicrobial

Uses

Help reduce bacteria on the skin

Warnings

For external use only

Flammable. Keep away from heat or flame.

When using this product avoid contact with eyes. If contact occurs, flush thoroughly with water. Avoid contact with broken skin. Do not ingest or inhale.

Stop use and ask doctor if skin irritation or redness occurs.

Keep out of reach of children. If swallowed, seek medical attention or contact Poison Control Center immediately.

Directions

Place a generous amount of product on hands and rub hands together until dry.

Other information

Do not store above 100° F. May cause fabric discoloration.

Inactive ingredients

Water (Agua), Sorbitol, Methycellulose, Polyquaternium-10, Fragrance, Glycerin

Dosage

Use a liberal amount in palm of hand.

Notice

Keep out of reach of children.

Indications & Usage

For use when soap and water are not available, or in addition to.

Label

Drug Facts
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8 FL OZ

Manufactured by:

EPIC Chemistry, LLC

3879 Maple Ave Ste 300

Dallas, TX 75219

www.epicchemistry.com

EHS7000

ehs7000 gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:770 22-1113
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70.05

Inactive Ingredients

Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)			33.12	
METHYLCELLULOSE, UNSPECIFIED (UNII: Z944H5SN0H)			0.6	
POLYQUATERNIUM-10 (30000 MPA.S AT 2 %) (UNII: C7RDC8Y4JS)			0.6	
SORBITOL (UNII: 506T60A25R)			1.13	
GLYCERIN (UNII: PDC6A3C0OX)			0.38	
Product Characteristics				
Color		Score		
Shape		Size		
Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77022-1113-8	1 in 1 BOTTLE; Type 0: Not a Combination Product	09/17/2020	

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4 FL OZ

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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/17/2020	

Labeler - EPIC Chemistry LLC (106734930)

Registrant - Epic Chemistry LLC (106734930)

Establishment

Name	Address	ID/FEI	Business Operations
EPIC Chemistry LLC		106734930	manufacture(77022-1113)

Revised: 9/2020

EPIC Chemistry LLC