

ENMOTION FOAM HIGH FREQUENCY USE- ethyl alcohol solution
Georgia-Pacific Consumer Products LP

enMotion High Frequency Foam Hand Sanitizer Fragrance Free 54622-121

☐ **Active ingredient**

Ethyl Alcohol 70% v/v

☐ **Purpose**

Antiseptic

Uses

- To decrease bacteria on skin that could cause disease
- Recommended for repeated use

☐ **Warnings**

- **FLAMMABLE, keep away from fire or flame**
- **For external use only**

When using this product do not use in or near eyes.

Stop use and ask a doctor if irritation or redness develops. If condition persists for more than 72 hours consult a doctor.

Keep out of reach of children.

If swallowed, seek immediate medical attention or call a poison control center.

Directions

- Wet hands thoroughly with product and allow to dry without wiping.

☐ **Inactive ingredients**

Water, Isopropyl Alcohol*, PEG-12 Dimethicone, Glycerin, Hydrolyzed Jojoba Esters, Caprylic/Capric Triglyceride, Panthenol, Isopropyl Myristate, Aloe Barbadensis Leaf Juice, Chamomilla Recutita (Matricaria) Flower Extract, Camellia Sinensis Leaf Extract, Avena Sativa (Oat) Kernel Extract, Euterpe Oleracea Fruit Extract, Butylene Glycol, Bisabolol, Phenoxyethanol, Citric Acid, Sodium Benzoate, Potassium Sorbate, Acetyl Hexapeptide-49

*may contain this ingredient

Rear label text

enMotion

High Frequency Use

Foam Hand Sanitizer

Fragrance Free SKU 42336

Manufactured for

Georgia-Pacific Consumer Products. Atlanta, GA 30303

Questions? Call 1-866-HELLOGP (435-5647)

or visit us online at www.gppro.com

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alcohol permit SDS-KY-15002

5000008656/000/02

principal display panel

High Frequency Use

Foam Hand Sanitizer

1000 mL (33.8 FL OZ)





High Frequency Use
Foam Hand Sanitizer
Fragrance Free • SKU 42336

Drug Facts

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Drug Facts (continued)

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ENMOTION FOAM HIGH FREQUENCY USE

ethyl alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54622-121
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.7 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PEG-12 DIMETHICONE (UNII: ZEL54N6W95)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROLYZED JOJOBA ESTERS (ACID FORM) (UNII: UDR641JW8W)	
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)	
PANTHENOL (UNII: WW9CM0O67Z)	
ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
MATRICARIA CHAMOMILLA FLOWERING TOP OIL (UNII: SA8AR2W4ER)	
CAMELLIA SINENSIS FLOWER (UNII: 9I2BJY2J17)	
AVENA SATIVA WHOLE (UNII: 5P8D0Z74RG)	
EUTERPE OLERACEA WHOLE (UNII: Y57H6218HP)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
.BETA.-BISABOLOL (UNII: LP618AV2EA)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
ACETYL HEXAPEPTIDE-49 (UNII: 4055X1S509)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54622-121-01	2 in 1 BOX	06/10/2021	
1		1000 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/10/2021	

