GERM-X ADVANCED- advanced hand sanitizer gel
Vi Jon, Inc

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.

Germ-X 746.000/AA

claims

Introducting the most effective Germ-x formula ever

*Effective at eliminating more than 99.99% of may common harmful germs *& bacteria in as little as 15 seconds

Active ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommend for repeated use

Warnings

For external use only: hands

Flammable. Keep away from heat and flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation or redness develops
- 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

- wet hands thoroghly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 105° F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, glyceryl caprylate/caprate, glycerin, isopropyl myristate, tocopheryl acetate, acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, benzophenone-4

Questions

1-866-MY GERMX - 1-866-694-3769

Adverse reactions

Manufactured By: Vi-Jon, Inc.

8515 Page Ave

St. Louis, MO 63114

www.germx.com

SDS MO-15036

SDA WI-2486

DSP-MO-28

DSP-MO-34

Pat.9,161,982

746.00/746AA

principal display panel

germ-X

ADVANCED

Hand Sanitizer

MORE EFFECTIVE FORMULA

kills more than 99.99% of germs

PATENTED

FORMULA

Original Scent

33.8 FL OZ (1 L)

GERM-X ADVANCED

advanced hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11344-746
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	700 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
SULISOBENZONE (UNII: 1W6L629B4K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-746-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2017	
2	NDC:11344-746-34	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/21/2017	
3	NDC:11344-746-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2017	
4	NDC:11344-746-26	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2017	
5	NDC:11344-746-38	295 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2017	
6	NDC:11344-746-32	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2017	
7	NDC:11344-746-49	443 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2017	
8	NDC:11344-746-88	1999 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/21/2017	
9	NDC:11344-746-86	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/21/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/21/2017	

Labeler - Vi Jon, Inc (150931459)

Registrant - Vi-Jon, Inc. (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, Inc.		088520668	manufacture(11344-746)