

I FAM SKIN HAND SANITIZER GEL- alcohol gel
Rainbow Co Ltd

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.

Drug Facts

alcohol

WATER

DIPROPYLENE GLYCOL

CARBOMER

GLYCERIN

TRIETHANOLAMINE

PEG-60 HYDROGENATED CASTOR OIL

PANTHENOL

CAMELLIA SINENSIS LEAF EXTRACT

ALOE 8AR8ADENSIS LEAF EXTRACT

DIETHANOLAMINE

CITRUS PARADIS! (GRAPEFRUIT) PEEL OIL

Sterilization of hands and skin

KEEP OUT OF REACH OF THE CHILDREN

Apply to clean, dry hands. Apply sufficient amount to thoroughly wet all surfaces of hands and fingers.
Rub onto hands until dry.

Supervise children in the use of this product.

- Flammable. Keep away from fire or flame.
- For external use only.
- Do not use in eyes.
- If swallowed, get medical help promptly.
- Stop use, ask doctor if irritation occurs.
- Keep out of reach of children.

for external use only

I FAM SKIN

Hand Sanitizer



I Fam Skin Hand Sanitizer Quasi drugs

70% ETHANOL !

- Non sticky after use
- Contains Aloe Extracts and D-Panthenol
- Advanced Hand Sanitizer

500 ml / 16.90 fl. oz. e



[Product name] I Fam Skin Hand Sanitizer Gel
 [Active ingredient] Ethanol
 [Other additives] Water, Dipropylene Glycol, Carbomer, Glycerin, Triethanolamine, PEG-60 Hydrogenated Castor Oil, Panthenol, Camellia Sinensis Leaf Extract, Aloe Barbadosis Leaf Extract, Diethanolamine, Citrus Paradisi (Grapefruit) Peel Oil
 [Efficacy and effectiveness] Sterilization of hands and skin
 [Usage and dosage] Apply an appropriate amount on your hands and rub well to dry.
 [Volume] 500 ml / 16.90 fl. oz.
 [How to store] Airtight container, storage at room temperature (1 – 30 °C)
 [Manufacturing number and expiry date] Notation separately
 [Precautions for use] 1. Do not use on the following body parts. A wide range of body parts and damaged skin around the eyes and ears in the oral cavity may have 2. If the following symptoms appear, stop using them immediately and consult a doctor or pharmacist. 1) Hypersensitivity symptoms such as rash, erythema, itching, and edema 2) Skin irritation symptoms 3. Other precautions 1) For external use only (do not underwear). 2) Be careful not to get into your eyes, and if so, rinse well with clean water and consult a doctor or pharmacist. 3) Be careful not to inhale the vapor when using it extensively or for a long period of time (irritation to the mucous membranes, headaches, etc. may occur if ethanol vapor is consumed in large quantities or repeatedly). 4) If repeated use on the same site, be careful as the skin may become rough due to degreasing. 5) Do not use sealed bandages, cast bandages, packs, etc., as irritation may occur. 6) Do not use this medicine for anal or vaginal compresses as it may cause irritation or chemical burns. 7) Do not use for any other purpose. 4. Precautions for storage 1) Avoid shading and keep in shading. 2) Keep it out of reach of children, and if a child swallows it, go to the hospital right away. 3) After use, close the product completely with a lid to prevent the product from drying out or entering foreign objects. 4) Taking it out of the original container and storing it in another container may cause accidents due to misuse or deterioration of quality, so store it in the original container. In case of any abnormality in this product, compensation will be made according to the consumer dispute resolution standards announced by the Fair Trade Commission.
 [Customer Service Center] +82 32 813 3364 MADE IN KOREA
 [Manufacturer] Rainbow Co.,Ltd.
 [Manufacturer's address] 60, 347beon-gil, Hambakmo-ro, Namdong-gu, Incheon (Namchon-dong)

I FAM SKIN HAND SANITIZER GEL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74247-0007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
GLYCERIN (UNII: PDC6A3C00X)	
GRAPEFRUIT PEEL (UNII: 3582N05Q44)	
PEG-60 HYDROGENATED CASTOR OIL (UNII: 02NG325BQG)	
DIMETHICONE (UNII: 92RU3N3Y10)	
PANTHENOL (UNII: WV9CM0O67Z)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74247-0007-1	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	
2	NDC:74247-0007-2	3 mL in 1 POUCH; Type 0: Not a Combination Product	06/13/2020	
3	NDC:74247-0007-3	15 mL in 1 POUCH; Type 0: Not a Combination Product	06/13/2020	
4	NDC:74247-0007-4	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/13/2020	

Labeler - Rainbow Co Ltd (690423720)**Registrant** - Rainbow Co Ltd (690423720)**Establishment**

Name	Address	ID/FEI	Business Operations
Rainbow Co Ltd		690423720	manufacture(74247-0007)

Revised: 6/2020

Rainbow Co Ltd