FOAM HAND SANITIZER- foam hand sanitizer liquid
Guangdong Kemei Pharmaceutical Technology 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.

75132-012
Foam Hand Sanitizer

ACTIVE INGREDIENTS

Sodium chloride 1.5%

Purpose

Antiseptic

Uses

Apply proper amount to palm, massage gently until foamy, then rinse with water.

Warnings

This product is for external use only.

When using this product In case of contact with eyes, rinse immediately with plenty of water.

Stop using and ask a doctor if irritation or rash occurs. These may be signs of a serious condition Flammable. Keep away from heat or flame

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

/

When using this product In case of contact with eyes, rinse immediately with plenty of water.

Stop using and ask a doctor if irritation or rash occurs. These may be signs of a serious condition Flammable. Keep away from heat or flame

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Stop using and ask a doctor if irritation or rash occurs. These may be signs of a serious condition Flammable. Keep away from heat or flame

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Directions

Scope of application: Household and personal cleaning.
Main Effect Effective inhibition of microorganisms category intestinal bacteria, pyogenic coccus.

Other information

Keep tightly closed. Store between 32°F and 109°F

Inactive ingredients

Citric acid, Cocoamidopropyl betaine, Coconut oil amide DEA, Disodium EDTA, DMDM hydantoin, Essence, Sodium lauryl alcohol polyether sulfate, Water

Package Label - Principal Display Panel

FOAM HAND SANITIZER

foam hand sanitizer liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75132-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM CHLORIDE	1.5 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	

SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
METHYL BENZOATE (UNII: 6618K1VJ9T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75132-012-03	950 mL in 1 BAG; Type 0: Not a Combination Product	04/12/2020	
2	NDC:75132-012-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
3	NDC:75132-012-02	950 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part349	04/12/2020	

Labeler - Guangdong Kemei Pharmaceutical Technology Co., Ltd. (554528507)

Registrant - Guangdong Kemei Pharmaceutical Technology Co., Ltd. (554528507)

Establishment

Name	Address	ID/FEI	Business Operations
Guangdong Kemei Pharmaceutical Technology Co., Ltd.		554528507	manufacture(75132-012)

Revised: 9/2020 Guangdong Kemei Pharmaceutical Technology Co., Ltd.