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.

75132-013

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

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Directions

Scope of application: Household and personal cleaning.

Main Effect Effective inhibition of microorganisms category intestinal bacteria, pyogenic coccus.

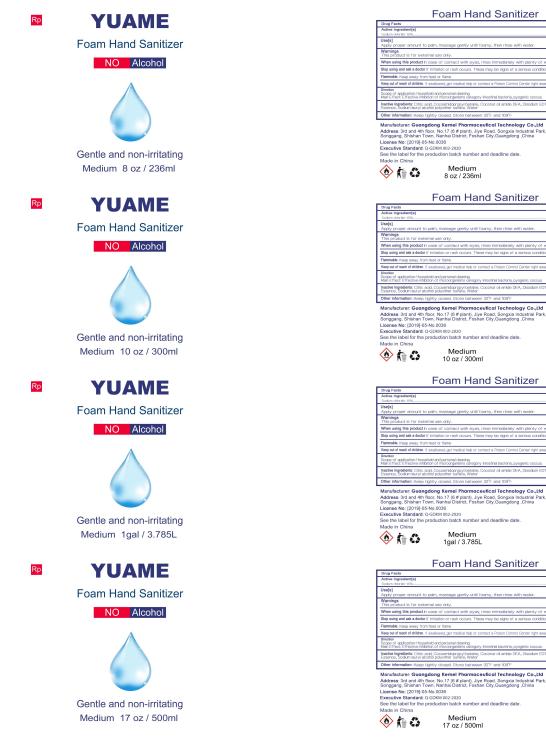
Other information

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

NO Alcohol



Gentle and non-irritating Medium 34 oz / 1L



Foam Hand Sanitizer

NO Alcohol



Gentle and non-irritating Medium 15 oz / 450ml



Foam Hand Sanitizer

NO Alcohol



Gentle and non-irritating Medium 1.3 gal / 5L

Foam Hand Sanitizer

g Facts	
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Manufacturer: Guangdong Kemel Pharmaceutical Technology Co.,Ltd Address: 3rd and 4th floor, No.17 (6 # plant), Jiye Road, Songxia Industrial Park, Songgang, Shishan Town, Nanhai District, Foshan City, Guangdong, China License, No.; (2019.10.F.N.), 0038

Executive Standard: Q-GDKM 002-2020

See the label for the production batch number and deadline date.



Medium 34 oz / 1L

Foam Hand Sanitizer

Active Ingredient(s) Sodum offorite 15%	Purpose
Use[s] Apply proper amount to palm, massage gently until foamy, then rinse with water.	
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Other information: Keep tightly closed. Store between 32°F and 109°F	

Manufacture Change (uptn) coses Store between 32°F and 100°F
Manufacture Changelong Kernel Phemaceutiled Technology Co.,Ltd
Address: 2rd and 4th floor, No.17 (6 s plant), Uye Road, Songosi netuurial Park,
Songosi, Shiham Town, Nahan District, Foshan City, Guangdong, China
License No; [2019]-05-No.0036
Executive Standard' C-GCMX 002°2020
See the label for the production batch number and deadline date.
Made in China



Medium 15 oz / 450ml

Foam Hand Sanitizer

Drug Facts	
Active Ingredient(s)	Purpos
Sedium chicride 16%. Use(s)	Antisopti
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Manufacturer: Guongdong Kemei Pharmaceutlical Technology Co.,Lid Address: 3rd and 4th floor, No.17 (6 # plant), Jlye Road, Songxia Industrial Park, Songsang, Shishan Town, Nanhai District, Foshan City, Guangdong, China License No; 2019-95-No.0036 Executive Standard: 0-GOM.002-2020 See the label for the production batch number and deadline date.





Medium 1.3 gal / 5L

FOAM HAND SANITIZER

foam hand sanitizer liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:75132-013

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)			
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3 O 11 KX)			
SODIUM LAURETH SULFATE (UNII: BPV390 UAP0)			

DMDM HYDANTO IN (UNII: BYR0546 TOW)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
EDETATE DISO DIUM ANHYDRO US (UNII: 8 NLQ36 F6 MM)	
METHYL BENZOATE (UNII: 6618K1VJ9T)	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:75132-013-03	450 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020		
2	NDC:75132-013-01	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020		
3	NDC:75132-013-02	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020		
4	NDC:75132-013-04	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020		
5	NDC:75132-013-07	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020		
6	NDC:75132-013-05	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020		
7	NDC:75132-013-06	3785 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(75132-013)		

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