

YUN HAND SANITIZER GEL- alcohol gel
KMPHARMACEUTICAL 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.

KMPHARMACEUTICAL Co.,Ltd. (JY COMMERCE) - Yun Hand Sanitizer Gel

Alcohol

water, carbomer, etc

Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

keep out of reach of the children

Place enough product on hands to cover all surfaces. Rub hands together until dry.

Supervise children under 6 years of age when using this product to avoid swallowing.

For external use only. Flammable. Keep away from heat or flame.

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

for external use only

YUN HAND SANITIZER GEL

PRODUCT NAME | YOON HAND SANITIZER GEL(ETHANOL)

ACTIVE INGREDIENT ■ ETHYL ALCOHOL 70%.....ANTIBACTERIAL AGENT(PURPOSE)

INACTIVE INGREDIENT ■ GLYCERINE, BUTYLENE GLYCOL, ALOEX (09), PURIFIED WATER, CARBOMER, TRIETHANOLAMINE, FRAGRANCES

EFFICACY EFFECT ■ STERILIZATION OF HANDS AND SKIN

USES ■ SPRINKLE OR REMOVE THE RIGHT AMOUNT ON YOUR HAND AND RUB IT WELL TO DRY

MEASURE OF CAPACITY ■ 500ML/ 16.9 FL. OZ

WARNINGS ■ CAUTIONS FOR THE USE OF EXTERNAL DISINFECTANTS ARE AS FOLLOWS.

1. DO NOT USE ON THE FOLLOWING BODY PARTS.AROUND THE EYES AND EARS,WITHIN THE MOUTH, EXTENSIVE BODY PARTS AND DAMAGED SKIN (CAN CAUSE IRRITATION). 2. IF THE FOLLOWING SYMPTOMS OCCUR, STOP USING THEM IMMEDIATELY AND CONSULT WITH DOCTORS AND PHARMACISTS. 1) IN CASE OF IRRITATION SUCH AS RASH, ERYTHEMA, ITCHING, EDEMA, ETC.) IN CASE OF SKIN IRRITATION 3. OTHER PRECAUTIONS TO BE TAKEN WHEN USING 1) USE IT FOR EXTERNAL USE ONLY (DO NOT WEAR IT INTERNALLY). 2) BE CAREFUL NOT TO GET INTO THE EYE. IF IT GOES IN, WASH IT WELL WITH CLEAN WATER AND CONSULT A DOCTOR OR PHARMACIST.) BE CAREFUL NOT TO INHALE STEAM WHEN USED EXTENSIVELY OR FOR LONG PERIODS OF TIME (AN IRRITATION TO THE MUCOUS MEMBRANE, HEADACHE, ETC. MAY APPEAR IF ETHANOL VAPOR IS DRUNK IN LARGE OR REPEATEDLY.)) WHEN USING THE SAME AREA REPEATEDLY, BE CAREFUL BECAUSE THE SKIN CAN GET ROUGH WITH DISLOCATION, ETC 5) DO NOT USE THE SEALED, CAST, OR PACK AS IT MAY CAUSE IRRITATION. 6) DO NOT USE THIS MEDICINE FOR ANAL OR VAGINAL STEAMING BECAUSE IT MAY CAUSE IRRITATION OR CHEMICAL BURNS.7) IT IS NOT USED EXCEPT FOR ITS PURPOSE. 4. STORAGE PRECAUTIONS 1) AVOID FIRE AND KEEP IT IN THE SHADE. 2) KEEP IT OUT OF THE REACH OF CHILDREN AND GO TO THE HOSPITAL IMMEDIATELY IF THE CHILD SWALLOWED IT. 3) AFTER USE, CLOSE THE LID COMPLETELY TO PREVENT ANY DRYNESS OR FOREIGN SUBSTANCES FROM ENTERING THE PRODUCT. 4) IF THE CONTAINER IS REMOVED FROM THE ORIGINAL CONTAINER AND STORED IN ANOTHER CONTAINER, IT CAN CAUSE ACCIDENTS CAUSED BY MISUSE OR DECREASE IN QUALITY, SO KEEP IT IN THE ORIGINAL CONTAINER.

OTHER INFORMATION ■ CLASSIFIED CONTAINER, ROOM TEMPERATURE (130 °C) STORAGE

BATCH NO./MFG DATE/EXP DATE ■ MARKED ON EACH PRODUCT (DD.MM.YYYY)

MANUFACTURED BY K.M. PHARMACEUTICALS
268 PYEONGTAEK PORT ROAD, POSEONG-EUP, GYEONGGI-DO

MADE IN KOREA BY JY COMMERCE
51ST STREET TO GANGNAM-DAERO, SEOCHO-GU, SEOUL

CONSUMER COUNSELING NUMBER +82 70-8065-5339

IF THERE IS A PROBLEM WITH THIS PRODUCT, COMPENSATION SHALL BE MADE IN ACCORDANCE WITH THE FAIR TRADE COMMISSION NOTICE "CONSUMER DISPUTE RESOLUTION STANDARDS."



YUN HAND SANITIZER GEL			
alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50555-002
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		
WATER (UNII: 059QF0K00R)			
Packaging			

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50555-002-01	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/06/2020	

Labeler - KMPHARMACEUTICAL Co.,Ltd. (688679158)

Registrant - KMPHARMACEUTICAL Co.,Ltd. (688679158)

Establishment

Name	Address	ID/FEI	Business Operations
KMPHARMACEUTICAL Co.,Ltd.		688679158	manufacture(50555-002) , label(50555-002) , pack(50555-002)

Revised: 4/2020

KMPHARMACEUTICAL Co.,Ltd.