NEW EGOROUND HAND SANITIZER- 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.

ACTIVE INGREDIENT

Active ingredients: ALCOHOL 70.0%

INACTIVE INGREDIENT

Inactive Ingredients:

Water, Glycerin, Butylene Glycol, Aloe Barbadensis Leaf Extract, Carbomer, Triethanolamine, Flavor

PURPOSE

Purpose: SANITIZER

WARNINGS

Warnings:

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

When using this product
do not use in or near the eyes.
In case of contact, rinse eyes thoroughly with water.

Discontinue use if
irritation or redness develops. If condition persists for more than 72 hours, consult a doctor.

KEEP OUT OF REACH OF CHILDREN

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

Uses

Uses:
Hand sanitizer to help reduce bacteria on the skin.

Directions

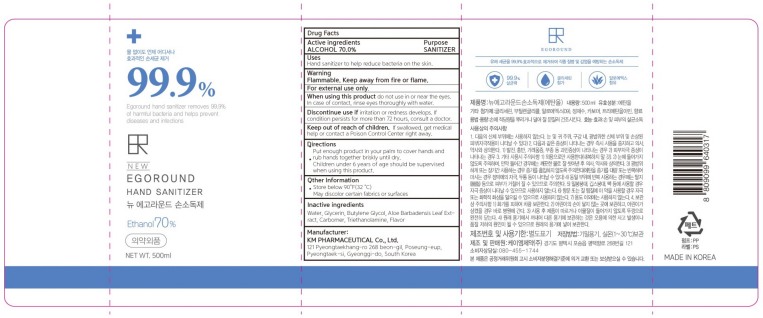
Directions:
Put enough product in your palm to cover hands and rub hands together briskly until dry.
Children under 6 years of age should be supervised when using this product.

Other Information

Other Information:
Store below 90°F(32°C)

May discolor certain fabrics or surfaces

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



NEW EGOROUND HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50555-070
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	350 mL in 500 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Glycerin (UNII: PDC6A3C0OX)	
Butylene Glycol (UNII: 3XUS85K0RA)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50555-070-01	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/01/2020	

Labeler - KMPHARMACEUTICAL Co.,Ltd. (688679158)

Registrant - KMPHARMACEUTICAL Co.,Ltd. (688679158)

Establishment

Name	Address	ID/FEI	Business Operations
KMPHARMACEUTICAL Co.,Ltd.		688679158	manufacture(50555-070)

Revised: 4/2020 KMPHARMACEUTICAL Co.,Ltd.