ENZYSEPT- alcohol gel
HP&C, 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

Active ingredient

Ethyl Alcohol 83% w/w

Purpose

Antiseptic

Uses

Hand Sanitizer to help reduce bacteria that potentially can cause disease.

For use when soap and water are not available.

Warnings

For external use only. Flammable.

Keep away from heat or flame

Do not use

- on open skin wounds

When using this product avoid contact with face, eyes, and broken skin. If eye contact occurs, flush 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.

Directions

- 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

Other information

- Do not rinse or wipe off the hand sanitizer before it's dry. Store in a cool dry place 1□(34□)~30□(86□)
- Avoid freezing and excessive heat above 40□(104□)

Inactive ingredients

Water, Aloe Barbadensis Leaf Extract, Aminomethyl Propanol, Polyacrylic Acid, Glycerin

Questions?

1-424-252-7887

Principal Display Panel – 60 mL Bottle Label

Enzysept

PREMIUM HAND SANITIZER

Kills 99.99% of germs in just 15 seconds

INSTANT KILL

Kills
99.99%
of germs

2 fl oz / 60 mL

Principal Display Panel – 500 mL Bottle Label

Enzysept

PREMIUM HAND SANITIZER

Kills 99.99% of germs in just 15 seconds

INSTANT KILL

Kills
99.99%
of germs

500 mL / 16.9 fl oz

Enzysept

PREMIUM HAND SANITIZER

INSTANT
KILL

의약외품

Kills

99.99%

of germs

of germs

엔지셉트
프리미엄 손 소독제
500mL / 16.9 fl oz

ENZYSEPT
alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71040-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	86 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Aloe Vera Leaf (UNII: ZY81Z83H0X)	
Aminomethylpropanol (UNII: LU49E6626Q)	
Polyacrylic Acid (250000 MW) (UNII: 9G2MAD7J6W)	
Glycerin (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:71040-101-01	60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	05/01/2020	
2	NDC:71040-101-02	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

Labeler - HP&C, Ltd. (689847430)

Establishment

Name	Address	ID/FEI	Business Operations
HP&C, Ltd.		689847430	MANUFACTURE(71040-101)

Revised: 4/2020 HP&C, Ltd.