PHOENIX HAND SANITIZER GEL- ethyl alcohol gel
THANH CONG PHARMACEUTICAL AND TRADING 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.

Phoenix Hand Sanitizer Gel

Drug Facts

Active ingredient

Ethanol 70% ± 3%

Purpose

Antiseptic

Uses

• Help to clean hand quickly.

• Resist bacteria without water, deodorizer.

Warnings: **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 eyes 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 at **1 - 800 -222 - 1222.**

Directions

• Use about 5 ml of solution, massage lightly whole hand that need to be cleaned until dry.

• Do not need to wash again with water.

Other information

• Store between 15-30°C (59-86°F).

• Avoid freezing and excessive heat above 40°C (104°F).

Inactive ingredients: Carbomer, Triethanolamine (TEA), Water.

Alcohol 70%

• *Clean hands quickly*

• *Resist bacteria*

• *Deodorizer*

DO NOT DRINK

NO NEED TO WASH AGAIN WITH WATER

Manufactured & packed for:

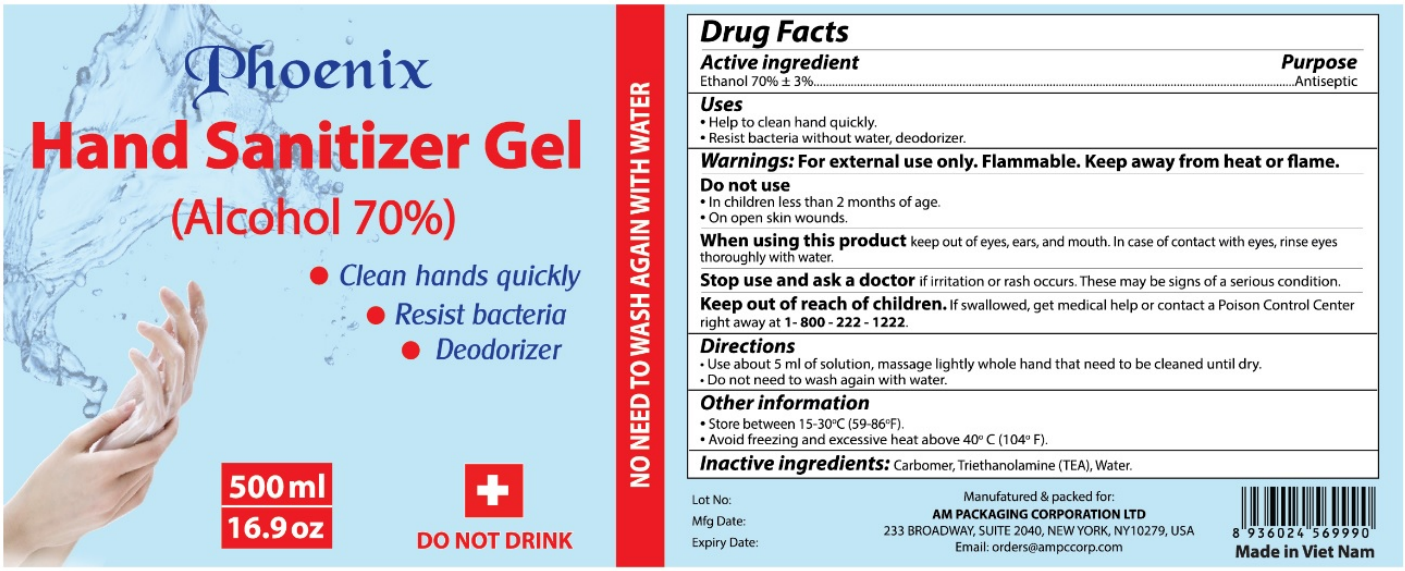
AM PACKAGING CORPORATION LTD

233 BROADWAY, SUITE 2040, NEW YORK, NY 10279, USA

Email: orders@ampccorp.com

Made in Viet Nam

Packaging



PHOENIX HAND SANITIZER GEL

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80666-555
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**
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	

WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80666-555-85	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2020	
2	NDC:80666-555-80	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2020	
3	NDC:80666-555-55	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2020	
4	NDC:80666-555-50	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2020	
5	NDC:80666-555-56	3800 mL in 1 CAN; Type 0: Not a Combination Product	10/01/2020	
6	NDC:80666-555-58	5000 mL in 1 CAN; Type 0: Not a Combination Product	10/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/01/2020	

Labeler - THANH CONG PHARMACEUTICAL AND TRADING CO.,LTD (555289565)

Establishment

Name	Address	ID/FEI	Business Operations
THANH CONG PHARMACEUTICAL AND TRADING CO.,LTD		555289565	manufacture(80666-555)

Revised: 10/2020 THANH CONG PHARMACEUTICAL AND TRADING CO.,LTD