PURIDENCE- ethyl alcohol gel
Ningbo SKL International 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.

PURIDENCE WATERLESS HAND SANITIZER

Active Ingredient

Ethyl Alcohol 70%

Purpose

Antiseptic

Warnings

For external use only
Flammable
Keep away from fire or flame

When using this product:

- Keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- Avoid contact with broken skin.
- Do not inhale or ingest

Stop use and ask a doctor if: irritation, rash or redness develops and persists.

Keep out of reach of children

If swallowed, seek medical help or contact a Poison Control Center immediately.

Directions

Apply enough product to cover all areas of hands.

Rub hands together briskly until dry.

Children under 6 years of age require supervision when using this product.

Uses

To help minimize bacteria on the skin that could cause disease.

Recommended for repeated use.

Other Information

Do not store above 105°F.

May cause fabric discoloration.

May harm plastic and wood finishes.

Inactive ingredient

Perified water, Propylene glycol, Aminomethyl propanol, Carbomer homopolymer, Glycerin

Product labels

PURIDENCE

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74542-011
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: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:74542-011-01	55 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
2	NDC:74542-011-02	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
3	NDC:74542-011-03	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
4	NDC:74542-011-04	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
5	NDC:74542-011-05	980 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
6	NDC:74542-011-06	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
7	NDC:74542-011-07	1300 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
8	NDC:74542-011-08	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	
9	NDC:74542-011-09	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/08/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/08/2020	

Labeler - Ningbo SKL International Co.,Ltd. (548277986)

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