HAND SANITIZER- alcohol gel
Urban Electric Power

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.

The hand sanitizer is manufactured using only the following:

a. Alcohol (Ethyl Alcohol) (70%, volume/volume (v/v))
b. Polyacrylic Acid

d. Hydrogen Peroxide

e. PEG 600

f. Glycerin

e.Triisopropanolamine

Drug Facts

Active Ingredients Purpose

Alcohol (Ethyl Alcohol) 70% v/v..... Antiseptic

Use(s)

Health care personnel wipe to help reduce bacteria that potentially can cause disease.

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.

Directions

- Unfold and apply wipe thoroughly onto hands or surface. Discard after single use. Do not flush.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients: Purified Water USP, Polyacrylic Acid, PEG, Hydrogen Peroxide, Glycerin, Neutralizer.

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

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

Warnings

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Package Label - Principal Display Panel

500 mL NDC: 74135-001-11

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HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74135-001
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**
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
POLYACRYLIC ACID (8000 MW) (UNII: 73861X4K5F)	
WATER (UNII: 059QF0KO0R)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
GLYCERIN (UNII: PDC6A3C0OX)	
TRIISOPROPANOLAMINE (UNII: W9EN9DLM98)	

Product Characteristics			
Color		**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:74135-001-01	100 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/15/2020	

Drug Facts

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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/15/2020	

Labeler - Urban Electric Power (078522589)

Establishment

Name	Address	ID/FEI	Business Operations
Urban Electric Power		078522589	manufacture(74135-001)