BIOXYGEN ANTISEPTIC HAND SANITIZER- isopropyl alcohol liquid
U S Fluids Inc

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.

BiOxygen Antiseptic Hand Sanitizer

Drug Facts

Active ingredient

Isoprpoyl alcohol 70% v/v

Purpose

Antiseptic

Uses

- hand sanitizer to decrease bacteria on the skin
- recommended for repeated use
- for use when soap and water are not available

Warnings

Flammable, keep away from fire/flame.

For external use only.

Do not use ▪ on children less than 2 months of age
▪ on open skin wounds

When using this product ▪ do not get into eyes. In case of contact, rinse eyes thoroughly with water

Stop use and ask a doctor if ▪ irritation and redness develop.
▪ condition persists more than 72 hours

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

Directions

- wet hands thoroughly with product and allow to dry without wiping.
- supervise children under 6 years of age when using this product to avoid swallowing.

Other information

▪ store between 15-30°C (59-83°F) ▪ avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients glycerin, hydrogen peroxide, hydroxyethylcellulose, water

***Questions?* +1-212-574-0784**

You may also report serious side effects to this phone number.

Mon-Fri 9.00 AM- 5.00 PM

Manufactured for Ener.co, 747 Third Avenue, New York, NY, 10017

Packaging

BIOXYGEN ANTISEPTIC HAND SANITIZER

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75189-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	0.7 L in 1 L

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75189-100-02	0.06 L in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
2	NDC:75189-100-04	0.12 L in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
3	NDC:75189-100-08	0.24 L in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
4	NDC:75189-100-10	0.3 L in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
5	NDC:75189-100-01	3.79 L in 1 JUG; Type 0: Not a Combination Product	05/01/2020	
6	NDC:75189-100-05	18.93 L in 1 JUG; Type 0: Not a Combination Product	05/01/2020	
7	NDC:75189-100-30	113.56 L in 1 DRUM; Type 0: Not a Combination Product	05/01/2020	
8	NDC:75189-100-55	208.20 L in 1 DRUM; Type 0: Not a Combination Product	05/01/2020	
9	NDC:75189-100-75	1041 L in 1 CONTAINER; 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 - US Fluids Inc (139405646)

Establishment

Name	Address	ID/FEI	Business Operations
US Fluids Inc		139405646	manufacture(75189-100)

Revised: 5/2020 US Fluids Inc