### INOCULEX HAND SANITIZER- ethyl alcohol liquid Private Label Supplements

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.

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#### Inoculex<sup>TM</sup> Hand Sanitizer

#### **Drug Facts**

#### **Active Ingredient**

Ethyl Alcohol 70%

#### **Purpose**

Antiseptic

**Use** as hand sanitizer to help reduce bacteria on the skin

#### **Warnings**

Flammable, keep away from fire or flame.

For external use only.

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

**Stop use and ask a doctor** if irritation or redness develops.

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

**Directions** • apply palmful to hands • rub hands together vigorously until dry • supervise children in the use of this product.

**Other Information** store below 110°F (43°C) • may discolor certain fabrics or surfaces.

*Inactive ingredients:* Fragrance, Glycerol, Hydrogen Peroxide, \*Hydroxypropyl Cellulose, \*Hydroxypropyl Methycellulose, Water

\*May or may not include

#### www.inoculex.com

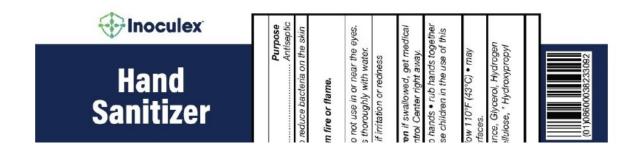
Distributed by: Inoculex, LLC

16 S Pennsylvania Ave.,

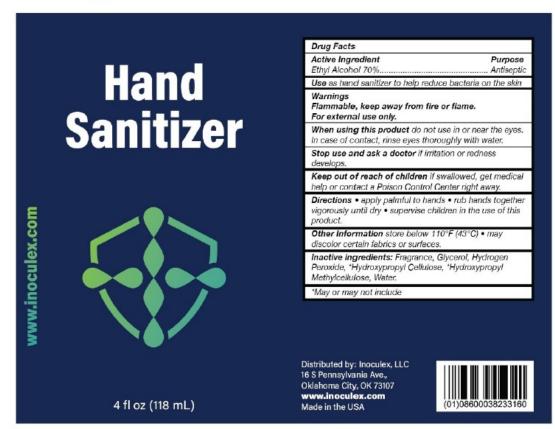
Oklahoma City, OK 73107

Made in the USA

#### **Packaging**













discolor certain fabrics or surfaces.

Inactive ingredients: Fragrance, Glycerol, Hydrogen Peroxide, "Hydroxypropyl Cellulose, "Hydroxypropyl Methylcellulose, Water.

\*May or may not include

8 fl oz (236 mL)

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# Hand Sanitizer

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16 fl oz (473 mL)

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## Hand **Sanitizer**

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32 fl oz (946 mL)

Drug Facts

Active Ingredient Ethyl Alcohol 70%.

Antiseptic

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Warnings

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## Hand **Sanitizer**



1 GAL (128 fl oz) 3.78L

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Inactive ingredients: Fragrance, Glycerol, Hydroger Peroxide, "Hydroxypropyl Cellulose, "Hydroxypropyl Methylcellulose, Water.

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### INOCULEX HAND SANITIZER

ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	

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		

NDC:77017-015

Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6A3C0OX)			
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)			
HYDRO XYPRO PYL CELLULO SE, UNSPECIFIED (UNII: 9 XZ8 H6 N6 OH)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
WATER (UNII: 059QF0KO0R)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:77017-015- 02	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/13/2020			
2	NDC:77017-015- 04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/13/2020			
3	NDC:77017-015- 08	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/13/2020			
4	NDC:77017-015- 16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/13/2020			
5	NDC:77017-015- 32	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/13/2020			
6	NDC:77017-015- 28	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/13/2020			

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not final	part333A	05/13/2020					

Revised: 5/2020 Private Label Supplements