# SIRETIZER HAND SANITIZER- alcohol liquid South Tenth Development, LLC

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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#### SIREtizer Hand Sanitizer

# **Drug Facts**

# **Active Ingredient**

Ethyl Alcohol 80%

# **Purpose**

Antiseptic

#### Uses

- -Rcducebacteria on the skin
- -Recommended for repeted use

#### **Warnings**

For external use only.

Flammable Keep away from fire, flame, or sparks

#### When using this product

do not use in or near eyes. In case of contact, rinse eyes thoroughly with water

#### Stop use & ask a doctor

If irritation or rash appears on the skin.

#### Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

- Place enough to cover both hands in the palm, and rub hands together until dry.
- Children under 6 years of age should be supervised by an adult when applying this product.

#### Other Information

- Store below 105°F.
- May discolor certain fabrics or surfaces.

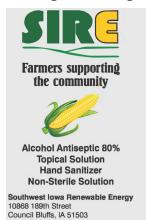
#### **Inactive Ingredients**

Distilled water, glycerol, hydrogen peroxide, denatonium benzoate.

## **Questions or Comments?**

Log onto our website www.sireethanol.com

# **Package Labeling:**







## SIRETIZER HAND SANITIZER

alcohol liquid

For SDS Information visit: www.SIREethanol.com

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78705-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.8 mL in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)		
<b>DENATO NIUM BENZO ATE</b> (UNII: 4YK5Z54AT2)		

l	Packaging			
l	# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
l	1 NDC:78705-002-00	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph not final	part333E	06/01/2020	

# Labeler - South Tenth Development, LLC (117521920)

Revised: 6/2020 South Tenth Development, LLC