

**ADRIANA 8OZ HAND SANITIZER -ORIGINAL- alcohol gel
OSTL, 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.

Active ingredient

Ethyl Alcohol 62.0%

Purpose

Antimicrobial

Uses

hand sanitizer to help reduce bacteria on the skin.

Warnings

For external use only.

Flammable. Keep away from heat and flame.

When using this product

avoid contact with face, eyes, and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

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.

Directions

- wet hands thoroughly with product and rub into skin until dry.
- Children under 6 years of age should be supervised by an adult when using this product.

Inactive ingredients

Water (Aqua), Aloe Barbadensis Leaf Juice, Carbomer, Fragrance, Glycerin, Propylene Glycol, Tocopheryl Acetate (Vitamin E), Triethanolamine

*Kills 99.9% of many common germs.

Imported by: **O.S.T.L. Inc**

Made in P.R.C.

Distributed by: **VERNON SALES. INC**

Drug Facts



KILLS 99.9% OF GERMS*

Hand Sanitizer

ANTIBACTERIAL

with Moisturizers & Vitamin E

8 FL OZ (236 ML)

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ADRIANA 8OZ HAND SANITIZER -ORIGINAL

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69950-021
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69950-021-01	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/31/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/31/2018	

Labeler - OSTL, INC. (020117798)

Registrant - OSTL, INC. (020117798)

Revised: 2/2018

OSTL, INC.