

**SANELL HAND SANITIZER- alcohol spray**  
**OraLabs**

*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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**Drug Facts**

**Active ingredient**

Ethyl Alcohol 63.00%

**Purpose**

Antiseptic

**Keep Out of Reach of Children**

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

**Uses**

For handwashing to decrease bacteria on the skin.

**Warnings**

For external use only: Hands. Flammable. Keep away from fire or flame. Do not use in the eyes. In case of contact, rinse eyes thoroughly with water.. Stop use and ask a doctor: if irritation and redness develop and persists for more than 72 hours.

**Directions**

Wet hands thoroughly with product and allow to dry without wiping. Supervise children in the use of this product.

**Inactive Ingredients**

Dimethicone, Disodium EDTA, DMDM Hydantoin, Fragrance, Polysorbate 20, Propanediol, Water.

**Package/Label Principal Display Panel**



**Kills 99.9% of Germs**  
Plus **MOISTURIZERS**



ASI 66712 PPAI 255799  
SAGE 66841

**HAND SANITIZER**  
NET WT. 0.25 fl oz (7.5mL)

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**Active ingredients**    **Purpose**  
 Alcohol Denat. 63% .....Antiseptic  
**Uses** • For hand washing to decrease bacteria on the skin.  
**Warnings** • For external use only.  
**Flammable. Keep away from fire or flame. Do not use** in the eyes. In case of contact, rinse eyes thoroughly with water.  
**Stop use and ask a doctor if** redness or irritation develops and persist for more than 72 hours.  
**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center.  
**Directions** • Wet hands thoroughly with product and allow to dry without wiping • supervise children in the use of this product.  
**Inactive ingredients:** Dimethicone, Disodium EDTA, DMDM Hydantoin, Fragrance, Polysorbate 20, Propanediol, Water.  
**Questions? 1-800-290-0557**

LAB-PPD-32997

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## SANELL HAND SANITIZER

alcohol spray

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63645-177
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	4.275 mL in 7.5 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	0.003 mL in 7.5 mL
<b>WATER</b> (UNII: 059QF0KO0R)	2.6857 mL in 7.5 mL

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63645-177-01	7.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/07/2018	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/07/2018	

**Labeler** - OraLabs (801824756)

**Registrant** - OraLabs (801824756)

## Establishment

Name	Address	ID/FEI	Business Operations
OraLabs		801824756	MANUFACTURE(63645-177) , LABEL(63645-177)

Revised: 11/2022

OraLabs