

SANELL HAND SANITIZER- alcohol spray
OraLabs

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)

Drug Facts
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

SPY SAN ORI

SANELL HAND SANITIZER

alcohol spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63645-177
Route of Administration	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
WATER (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 Drug	M003	03/07/2018	

Labeler - OraLabs (801824756)

Registrant - OraLabs (801824756)

Establishment

Name	Address	ID/FEI	Business Operations
OraLabs		801824756	manufacture(63645-177) , label(63645-177) , analysis(63645-177)

Revised: 4/2025

OraLabs