

CLOROX HAND SANITIZER- alcohol solution
Brand Buzz 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.

Clorox Hand Sanitizer (Bleach-Free)

Active Ingredient

Alcohol 71% v/v

Purpose

Antiseptic

Use

Hand sanitizer to help decrease bacteria on skin that can potentially cause disease.

Warnings

Flammable. Keep away from fire or flame.

For external use only.

Do not use in the eyes.

Stop use and ask a doctor if

- irritation and redness develop
- condition persists for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact a poison control center immediately.

Directions

- Spray on clean, dry hands and nails
- Wet hands thoroughly with product
- Rub hands together for no less than 15 seconds, paying particular attention to interdigital spaces, fingernails and cuticles
- Allow to dry without wiping
- Rub hands together briskly until dry

Other Information

- Store in a cool, dry place below 104 deg F (40 C)

Inactive Ingredients

Water, Isopropyl Alcohol, Glycerin, Glyceryl Laurate

Questions or Comments?

Visit us at www.cloroxprofessional.com or call **1-888-508-4750**.

Package Label: 59ml/500ml

Clorox Commercial Solutions®

CLOROX

BLEACH-FREE

Hand Sanitizer

Kills greater than 99.999%* of germs on contact

Contains hand moisturizers

Alcohol-based

*of FDA organism in the TFM for Topical Antimicrobial Drug Products



CLOROX HAND SANITIZER

alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69540-0028
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	71 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERYL LAURATE (UNII: Y98611C087)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69540-0028-1	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/20/2021	
2	NDC:69540-0028-4	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/20/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/20/2021	

Labeler - Brand Buzz LLC (079266204)

Revised: 2/2023

Brand Buzz LLC