CLOROX HAND SANITIZER- alcohol solution The Clorox Company

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

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.

Discontinue 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

Ouestions or Comments?

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

Clorox

Commercial

Solutions

CLOROX

BLEACH-FREE

Hand Sanitizer

Kills greater than

99.999%* of

germs on contact

Contains hand moisturizers

Alcohol-based

*of FDA organisms in the TFM for

Topical Antimicrobial Drug

Products

16.9 FL OZ I 500 mL



CLOROX HAND SANITIZER

alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:26509-0106
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:26509- 0106-1	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/06/2006	
2	NDC:26509- 0106-4	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/06/2006	
3	NDC:26509- 0106-5	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/06/2006	
4	NDC:26509- 0106-6	1000 mL in 1 BAG; Type 0: Not a Combination Product	11/06/2006	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/06/2006	

Labeler - The Clorox Company (009138033)

Establishment				
Name	Address	ID/FEI	Business Operations	
Best Sanitizers, Inc		627278224	manufacture(26509-0106)	

Revised: 4/2021 The Clorox Company