CLOROX ANTIMICROBIAL HAND SANITIZER- ethyl alcohol gel
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.

Drug Facts

Ethyl Alcohol 62% w/w

Antimicrobial

Uses

• To sanitize hands without requiring water or a rinse

Warnings

FLAMMABLE

For external use only

When using this product

• do not use near heat or flame

• do not use in or near eyes

• discontinue use if irritation and redness develop, or if condition persists for more than 72 hours

KEEP OUT OF REACH OF CHILDREN. In case of accidental ingestion, seek medical attention or contact a poison control center immediately.

Directions

• Wet hands thoroughly with product and allow to dry without wiping

Other Information

• Store at room temperature

Inactive ingredients

- Aloe Barbadensis Leaf Juice
- Carbomer
- Diisopropylamine
- FD&C Blue #1
- Fragrance
- Glycerin
- Isopropyl Myristate
- Phenoxyethanol
- Tocopherol Acetate
- Water

QUESTIONS OR COMMENTS?

Call **1-800-638-2625** or visit www.hlk.cc

Clorox Antimicrobial Hand Sanitizer

18 FL OZ (532 mL)

CLOROX ANTIMICROBIAL HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:26509-0010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	620 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
DIISOPROPANOLAMINE (UNII: 0W44HYL8T5)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:26509-0010-8	532 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/25/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/25/2016	

Labeler - The Clorox Company (009138033)

Registrant - Carroll Company (007372329)

Establishment

Name	Address	ID/FEI	Business Operations
Carroll Company		007372329	manufacture(26509-0010)

Revised: 1/2016 The Clorox Company