

**CIRCLE K HAND SANITIZER- alcohol gel
Lil' Drug Store Products, Inc.**

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.

Circle K ® Hand Sanitizer, 1.25oz

Drug Facts

Active ingredient

Ethyl alcohol 62%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only: hands

Flammable. Keep away from heat and flame.

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation or redness develops
- condition persists for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, glycerin, propylene glycol, acrylates/C10-C30 alkyl acrylate crosspolymer, triethanolamine, aloe barbadensis leaf juice, maltodextrin

Questions?

1-877-507-6516 (M-F 8AM-4:30PM CST)

Proudly distributed by Circle K Stores Inc

PRINCIPAL DISPLAY PANEL - 37 mL Bottle Label

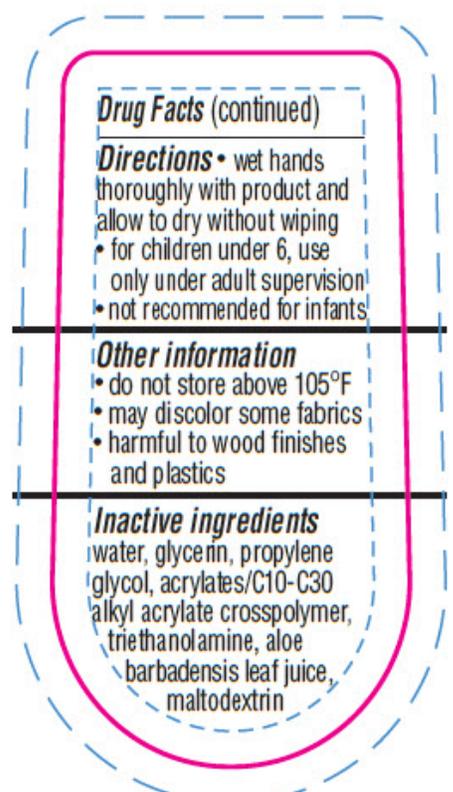
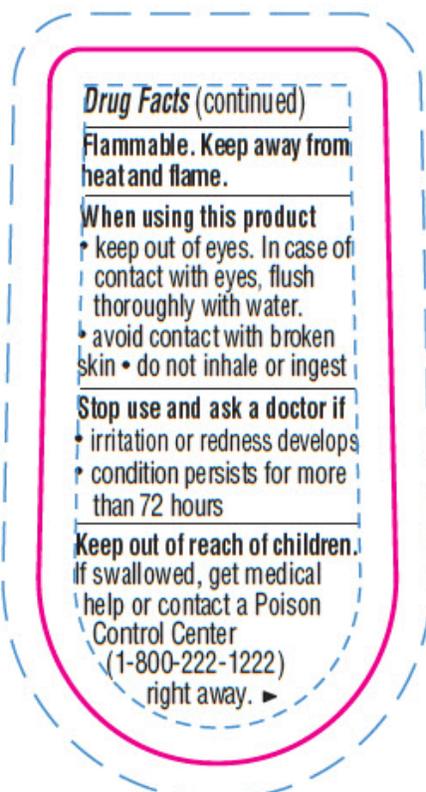
CIRCLE K ®

Hand
Sanitizer

Kills 99.99%
of Germs*

*Effective at eliminating 99.99% of
many common harmful germs and
bacteria in as little as 15 seconds.

1.25 FL OZ
(37 mL)



CIRCLE K HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66715-5863	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	62 mg in 100 mL	
Inactive Ingredients				
Ingredient Name				Strength
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
TROLAMINE (UNII: 9O3K93S3TK)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66715-5863-2	37 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/06/2017	01/28/2025
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part333E		06/06/2017	01/28/2025

Labeler - Lil' Drug Store Products, Inc. (093103646)

Revised: 8/2022

Lil' Drug Store Products, Inc.