MAGICARE PREMIUM HAND SANITIZING WIPES- alcohol cloth
Chaozhou Cecilia Technology Co., Ltd.

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.

MagiCare Premium Hand Sanitizing Wipes

DRUG FACTS

Active Ingredient

Ethyl Alcohol 75%

Purpose

Antimicrobial

Warnings:

Flammable. Keep away from fire or flame.

• For external use only

• Do not use in or near the eyes.

In case of contact, rinse eyes thoroughly with water.

• Stop use and consult with a physician if irritation develops and lasts for more than 72 hours.

• Keep out of reach of children.

• If swallowed, get medical help or contact a Poison Control Center Immediately.

Uses:

Hand sanitizer to help reduce bacteria on the skin.

Directions:

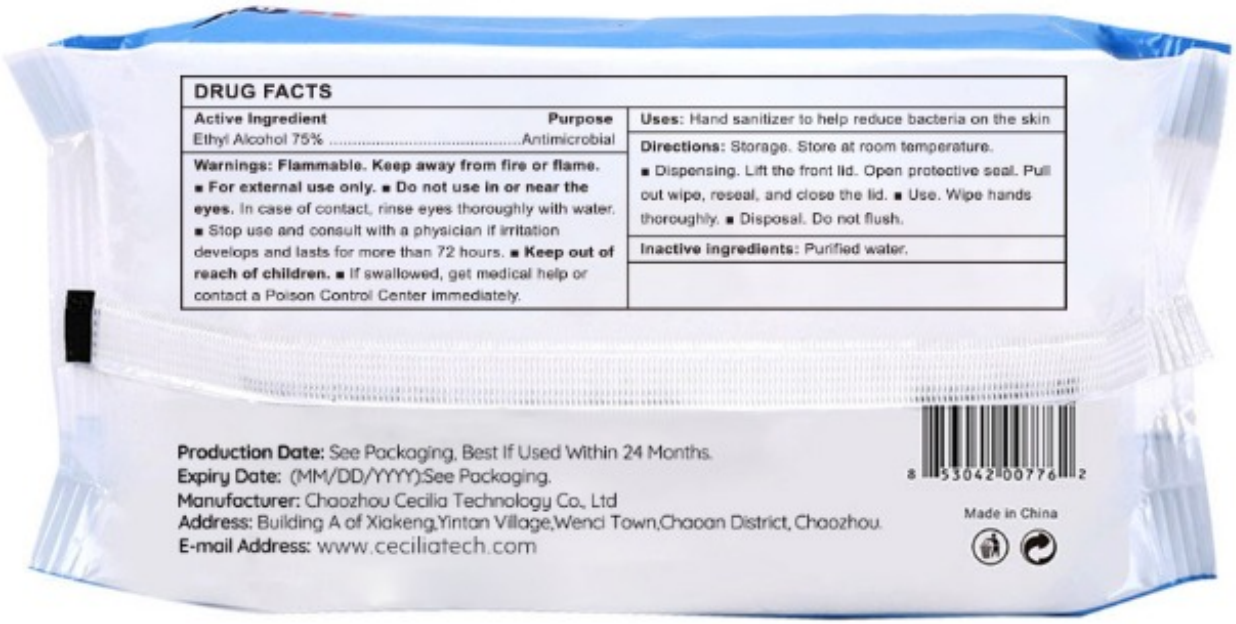
Storage. Store at room temperature.

• Dispensing. Lift the front lid. Open protective seal. Pull out wipe, reseal, and close the lid. • Use. Wipe hands thoroughly. • Disposal. Do not flush.

Inactive ingredients:

Purified water.

Package Labeling: 80294-301-04



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Uses: Hand sanitizer to help reduce bacteria on the skin

Directions: Storage. Store at room temperature. ■ Dispensing. Lift the front lid. Open protective seal. Pull out wipe, reseal, and close the lid. ■ Use. Wipe hands thoroughly. ■ Disposal. Do not flush.

Inactive ingredients: Purified water.

Production Date: See Packaging, Best If Used Within 24 Months.
Expiry Date: (MM/DD/YYYY):See Packaging.
Manufacturer: Chaozhou Cecilia Technology Co., Ltd
Address: Building A of Xiakeng,Yintan Village,Wenci Town,Chaoan District, Chaozhou.
E-mail Address: www.ceciliatech.com

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Made in China


MagiCare
PREMIUM
HAND SANITIZING WIPES
75%
ALCOHOL
Kills 99% of Germs
Kills 99% of Germs


Kills 99% of Germs
75%
Kills 99% of Germs


80
WIPES
Net Contents
Wipes 6 x 8 in.
Kills 99% of Germs
75%
Kills 99% of Germs

Package Labeling: 80294-301-05

MAGICARE PREMIUM HAND SANITIZING WIPES

alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80294-301
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80294-301-04	80 in 1 POUCH	06/01/2020	
1		240 g in 1 POUCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		
2	NDC:80294-301-05	100 in 1 CANISTER	06/01/2020	
2		300 g in 1 CANISTER; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/01/2020	

Labeler - Chaozhou Cecilia Technology Co., Ltd. (554536556)

Revised: 3/2023 Chaozhou Cecilia Technology Co., Ltd.