

**EVERLIT SURVIVAL ANTISEPTIC WIPE- antiseptic wipe cloth
EVERLIT GLOBAL INC.**

83807-024, Everlit Survival Antiseptic Wipe

Active Ingredient

Benzalkonium Chloride, 0.13%

Purpose

To disinfect skin surfaces

Uses

Sanitize Hands/ Clean Wounds

Warnings

Avoid contact with eyes, if happens, rinse thoroughly with water

For External Use Only

If pain, irritation, swelling, or infection develops, discontinue use and consult a physician

Keep out of reach of children

In case of accidental swallowing, seek medical help or contact a Poison Control Center immediately

Directions

Tear open packet, unfold towelette, use to cleanse desired skin area. Discard towelette appropriately after use.

Other Information

Store at room temperature

Inactive Ingredients

Purified Water, Sodium Bicarbonate



EVERLIT CARE® PRODUCTS

ANTISEPTIC WIPE

FIRST AID ANTISEPTIC
SINGLE-USE | EXTERNAL USE ONLY

NDC: 83807-024-01

LOT 251210
2026-01-15
2031-01-14

PROUDLY OWNED BY VETERANS

DRUG FACTS

Active Ingredients Benzalkonium Chloride 0.13%

Inactive Ingredients Purified water, Sodium Bicarbonate

Purpose To disinfect surfaces and skin

Use Sanitize hands | Clean wounds | Disinfect surfaces

Directions
Tear open packet, unfold towelette, use to cleanse desired skin area. Discard towelette appropriately after use.

Storage Store at room temperature

Warnings
Avoid contact with eyes. If happens, rinse thoroughly with water. For External Use Only. If pain, irritation, redness, swelling, or infection develops, discontinue use and consult physician.

Keep Out Of Reach Of Children
In case of accidental swallowing, seek medical help and contact poison control immediately.

MANUFACTURED FOR EVERLIT GLOBAL INC. ONTARIO, CALIFORNIA MADE IN P.R.C.

PROUDLY OWNED BY VETERANS

TEAR HERE

EVERLIT SURVIVAL ANTISEPTIC WIPE				
antiseptic wipe cloth				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83807-024	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)		BENZALKONIUM CHLORIDE	1.22 g	
Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
SODIUM BICARBONATE (UNII: 8MDF5V39QO)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83807-024-01	1 in 1 PACKET; Type 0: Not a Combination Product	12/26/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	12/26/2023	

Labeler - EVERLIT GLOBAL INC. (122311450)

Registrant - EVERLIT GLOBAL INC. (122311450)

Establishment

Name	Address	ID/FEI	Business Operations
Changzhou Maokang Medical Products Co., Ltd		421317073	manufacture(83807-024)

Revised: 1/2026

EVERLIT GLOBAL INC.