

MEDICHOICE CLEANSING- benzalkonium chloride cloth
Owens & Minor Distribution, Inc.

MediChoice Cleansing Towelette

Drug Facts

Active ingredient

Benzalkonium Chloride, 0.13% w/w

Purpose

First Aid Antiseptic

Use

First aid antiseptic product to apply topically to the skin to help prevent infection in minor cuts, scrapes, and burns.

Warnings

For external use only on mucouse membranes or broken or irritated skin.

Do not use

- in the eyes or apply over large areas of the body.
- longer than 1 week unless directed by a doctor.
- no infants. **Consult a doctor**
- in case of deep or punture wounds, animal bites, or serious burns.

Stop use and consult a doctor

If the condition persists or gets worse.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison control Center right away.

Directions

- Clean the affected area 1 to 3 times daily or as directed by a doctor. May be covered with a sterile bandage when dry.

Other information

- Store at room temperature.
- Allow pad to air dry before disposal.

Inactive ingredient

Isopropyl alcohol, Purified Water

NDC 39892-0305-1

MEDICHOICE®

Cleansing Towelette

Benzalkonium Chloride First Aid Antiseptic

1 Each
Reorder: **CLT100**

NOT MADE WITH natural rubber latex

sterile[®]

WARNING: For External use only.
CAUTION: Sterility of contents guaranteed in unopened, undamaged package.

LOT stamped position

Distributed by Owens & Minor
9120 Lockwood Boulevard
Mechanicsville, VA 23116
Made in China

Rev. E

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MEDICHOICE CLEANSING

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:39892-0305
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:39892-0305-1	10 in 1 CASE	08/29/2017	
1		100 in 1 BOX		
1		1 in 1 PACKET		
1		0.75 g in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/29/2017	

Labeler - Owens & Minor Distribution, Inc. (847412269)

Registrant - Owens & Minor Distribution, Inc. (847412269)

Revised: 12/2023

Owens & Minor Distribution, Inc.