

MED NAP CLEANSING TOWELETTE- benzalkonium chloride liquid
Acme United Corporation

Med Nap Cleansing Towelette

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

First Aid antiseptic

Use:

First Aid antiseptic to help prevent infection in minor

- cuts
- scrapes
- burns

Caution:

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control center right away.

Warnings

For external use only

Do not use ■in the eyes ■over large areas of the body •longer than 1 week

Stop use

Stop use and ask doctor if the condition persists or gets worse

Ask doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Directions

To open: Tear packet open at notch, remove towelette, use it only once

- Clean affected area
- Apply 1 to 3 times daily
- May be covered with a sterile bandage, once area dries

Other Information

Store at room temperature

Inactive Ingredients

isopropyl alcohol, methylchloroithiazolinone/methylisothiazolinone mixture, water

Questions? 1.800.835.2263

Reorder No. **94037**

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Manufactured by:
Acme United Corporation
1 Waterview Dr, Shelton, CT 06484
www.mednap.us
Made in the USA
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Lot No. _____

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**CLEANSING
TOWELETTE**
Benzalkonium Chloride Antiseptic

100 Packets

Not made with natural rubber latex.

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Box Image

MED NAP CLEANSING TOWELETTE			
benzalkonium chloride liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-0243
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.0018 mg in 1.35 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-0243-00	1.35 mL in 1 PACKET; Type 0: Not a Combination Product	03/02/2022	
2	NDC:0924-0243-01	100 in 1 BOX	03/02/2022	
2		1.35 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/02/2022	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		117825595	manufacture(0924-0243)

Revised: 11/2024

Acme United Corporation