MED NAP BENZALKONIUM CHLORIDE ANTISEPTIC- benzalkonium chloride liquid Acme United Corporation

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.

Med Nap Benzalkonium Chloride Antiseptic Towelette

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

First Aid Antiseptic

Uses

First Aid antiseptic to help prevent infection in minor

- cuts
- scrapes
- burns

Warnings

For External Use Only.

Do not use

- in the eyes
- over large areas of the body
- •longer than 1 week

Ask a doctor before use if you have •Deep or puncture wounds •Animal bites

Serious burns

Stop use

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

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Keep out of reach of children.

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

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

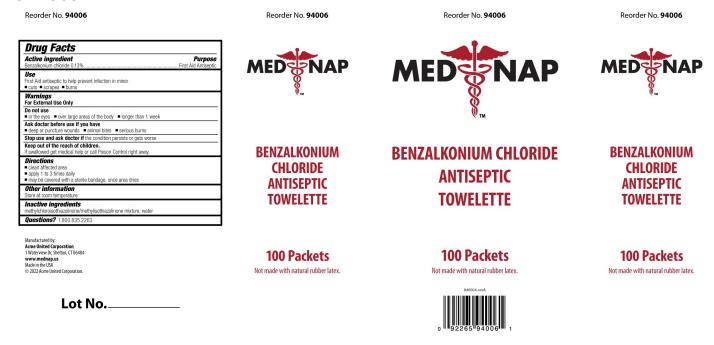
Other Information

Store at room temperature

Inactive Ingredients

methylchloroisothiazolinone/methylisothiazolinone mixture, water 1.800.835.2263

Box Label



MED NAP BENZALKONIUM CHLORIDE ANTISEPTIC benzalkonium chloride liquid Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:0924-0246 Route of Administration TOPICAL

ı	Active Ingredient/Active Moiety		
	Ingredient Name	Basis of Strength	Strength
		BENZALKONIUM CHLORIDE	0.0018 mg in 1.35 mL

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)			
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:0924-0246- 00	1.35 mL in 1 PACKET; Type 0: Not a Combination Product	02/11/2022			
2	NDC:0924-0246- 01	100 in 1 BOX	02/11/2022			
2		1.35 mL in 1 PACKET; Type 0: Not a Combination Product				
3	NDC:0924-0246- 02	1000 in 1 BOX	02/11/2022			
3		1.35 mL in 1 PACKET; Type 0: Not a Combination Product				
4	NDC:0924-0246- 03	12000 in 1 BOX	02/11/2022			
4		1.35 mL in 1 PACKET; Type 0: Not a Combination Product				
5	NDC:0924-0246- 04	20 in 1 BOX	02/28/2023			
5		1.35 mL in 1 PACKET; Type 0: Not a Combination Product				
6	NDC:0924-0246- 05	25 in 1 BOX	02/28/2023			
6		1.35 mL in 1 PACKET; Type 0: Not a Combination Product				
7	NDC:0924-0246- 06	60000 in 1 CONTAINER	02/28/2023			
7		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 not final	part333E	02/11/2022		

Establishment					
Name	Address	ID/FEI	Business Operations		
Acme United Corporation		117825595	manufacture(0924-0246)		

Revised: 2/2023 Acme United Corporation