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

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

•**i**n 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

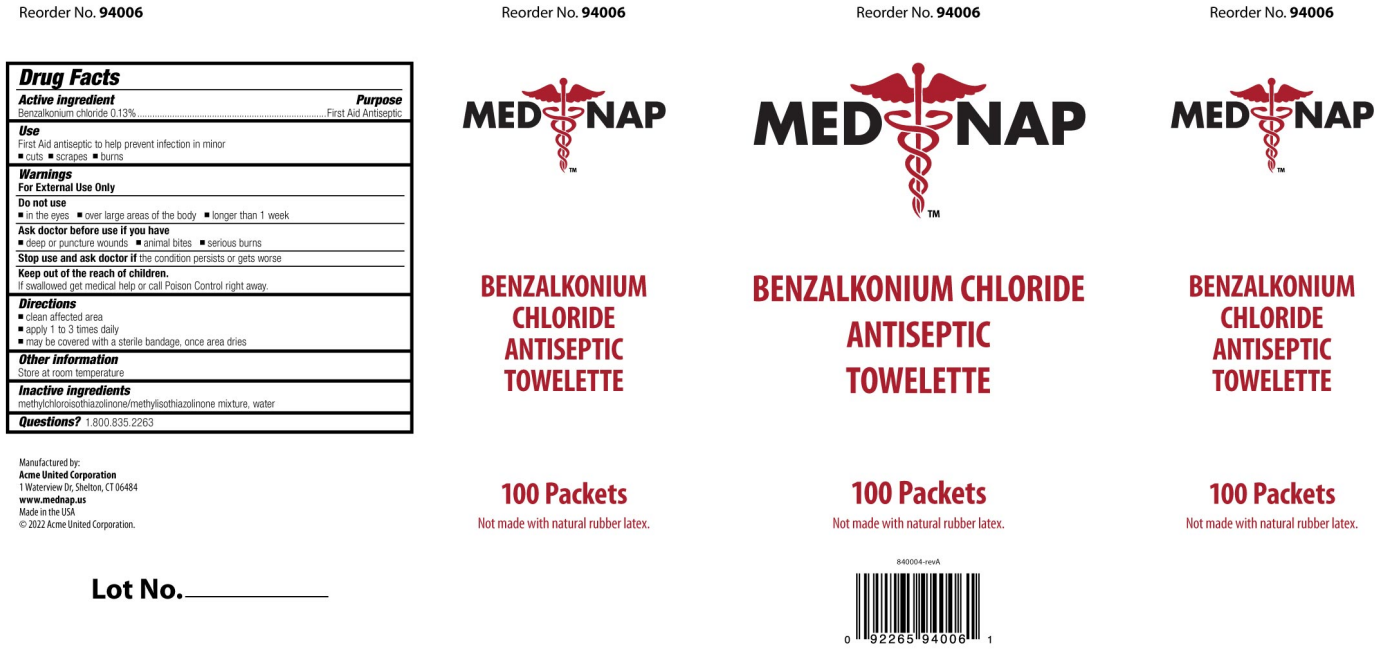
Store at room temperature

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		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM	0.0018 mg

UNII:7N6JUD5X6Y)	CHLORIDE	in 1.35 mL

Inactive Ingredients

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

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		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	02/11/2022	

Labeler - Acme United Corporation (001180207)

Establishment

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

Revised: 11/2024 Acme United Corporation