MED NAP CLEANSING TOWELETTE- 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 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

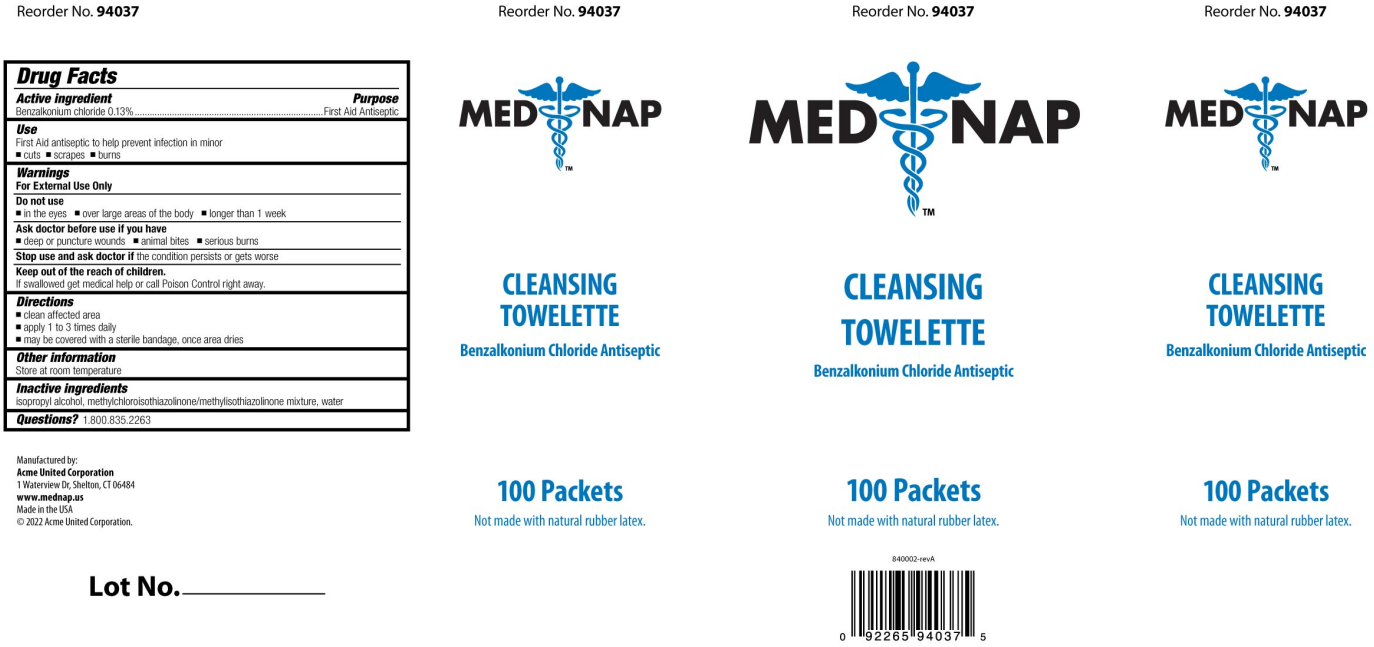
Other Information

Store at room temperature

Inactive Ingredients

isopropyl alcohol, methylchloroisothiazolinone/methylisothiazolinone mixture, water

Questions? 1.800.835.2263



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: 059QF0KO0R)	
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 not final	part333E	03/02/2022	

Labeler - Acme United Corporation (001180207)

Establishment

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

Revised: 3/2022 Acme United Corporation