MED NAP OBSTETRICAL 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 Obstetrical Towelette

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

First Aid antiseptic

Use:

Antiseptic cleansing of face, hands, and body without soap and water. Air dries in seconds.

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

Stop use

Stop use if irritation, redness, or other symptoms develop. Consult a doctor if the condition persists or gets worse.

Directions

Tear open packet, unfold, and use as a washcloth.

Other Information

Store at room temperature

Inactive Ingredients

chlorothymol, isopropyl alcohol, methylchloroisothiazolinone/methylisothiazolinone mixture, water

Questions? 1.800.835.2263

Box Image

Reorder No. 94005

Manufactured by:
Acme United Corporation

1 Waterview Dr, Shelton, CT 06484
www.mednap.us
Made in the USA
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Lot No.

MEDWNAP

Reorder No. 94005

OBSTETRICAL TOWELETTE Benzalkonium Chloride Antiseptic

100 Packets

Not made with natural rubber latex.

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Benzalkonium Chloride Antiseptic

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840003-revA

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MED NAP OBSTETRICAL TOWELETTE

benzalkonium chloride liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0924-0244

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZ ALKONIUM	0.0018 mg

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

CHLOROTHYMOL (UNII: LJ25TI0CVT)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-0244- 00	1.35 mL in 1 PACKET; Type 0: Not a Combination Product	02/25/2022	
2	NDC:0924-0244- 01	100 in 1 BOX	02/25/2022	
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	02/25/2022	
	partosse	02/25/2022	

Labeler - Acme United Corporation (001180207)

Establishment				
Na me	Address	ID/FEI	Business Operations	
Acme United Corporation		117825595	manufacture(0924-0244)	

Revised: 2/2022 Acme United Corporation