ANTISEPTIC TOWELETTE- benzalkonium chloride liquid
Med-Nap LLC

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.

Antiseptic Towelette 750

Active Ingredients

Benzalkonium Chloride 0.133%

Purpose

First Aid Antiseptic

Keep out of reach of children.

Caution Keep Out of Reach of Children. If swallowed, get medical help or contact a Poison Control Center right away.

Use

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

WARNINGS:

For external use only. Do not use in or around the eyes.

Directions

Tear open packet, unfold and use as a washcloth.

Stop Use

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

Inactive Ingredients

Kathon CG, Water

Antiseptic Towelette Label

WET NAP bulk product Label

MAX

PACKAGING

NDC 59647-750-01

Antiseptic Towelette

Contains Benzalkoniun Chloride

For External use Only

1 Towelette

MAX PACKAGING

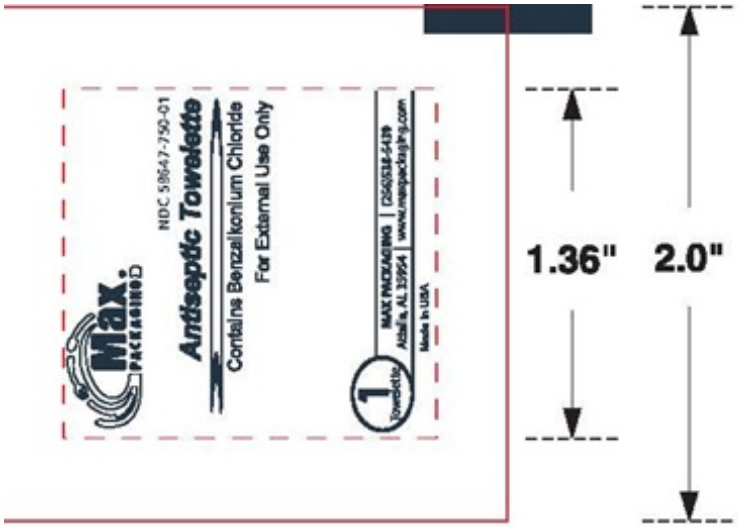
Attalla, AL

(256) 538-543www.maxpackaging.com

Made in USA

EXP: 03/2018

LOT MN06916 ABC



WET NAP

60,000

MAX

ANTISEPTIC

LOT #15048 MAX-BZK

WET NAP

60,000

MAX

ANTISEPTIC

LOT #15048 MAX-BZK

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59647-750
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.1 mg in 1.4 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
MAGNESIUM CHLORIDE ANHYDROUS (UNII: 59XN63C8VM)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59647-750-00	60000 in 1 CONTAINER	02/15/2016	
1	NDC:59647-750-01	1.4 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	part333A	02/15/2016	

Labeler - Med-Nap LLC (079086400)

Registrant - Med-Nap LLC (079086400)

Establishment

Name	Address	ID/FEI	Business Operations
Med-Nap LLC		079086400	manufacture(59647-750)

Revised: 2/2016 Med-Nap LLC