MED NAP - 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.

Benzalkonium Chloride Towelette - 245

Drug Facts
Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

First Aid Antiseptic

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.

Warnings

For External Use Only.

Do not use in or around the eyes.

Do not apply over large area of the body.

Stop use

if irritation, redness or other symptoms develop.

Consult a doctor if the condition persists or gets worse.

Directions

Tear open packet, unfold towelette and use to cleanse desired skin area. Discard towelette appropriately after a single use.

Inactive Ingredients

Water, Kathon CG

MED NAP BENZALKONIUM CHLORIDE TOWELETTE

MED NAP

Record No. 3303

NDC # 59647-245-01

BZK
ANTISEPTIC
TOWELETTE

LatexFree

1 Pouch

Made by:
MED-NAP LLC,
Brooksville, FL 34601

www.mednap.us

Recorder No. 3303

Made in the USA

* See box for full Drug Facts information

Lot: 16325 E F G EXP: 10/2018

Made In USA 1 REV 3


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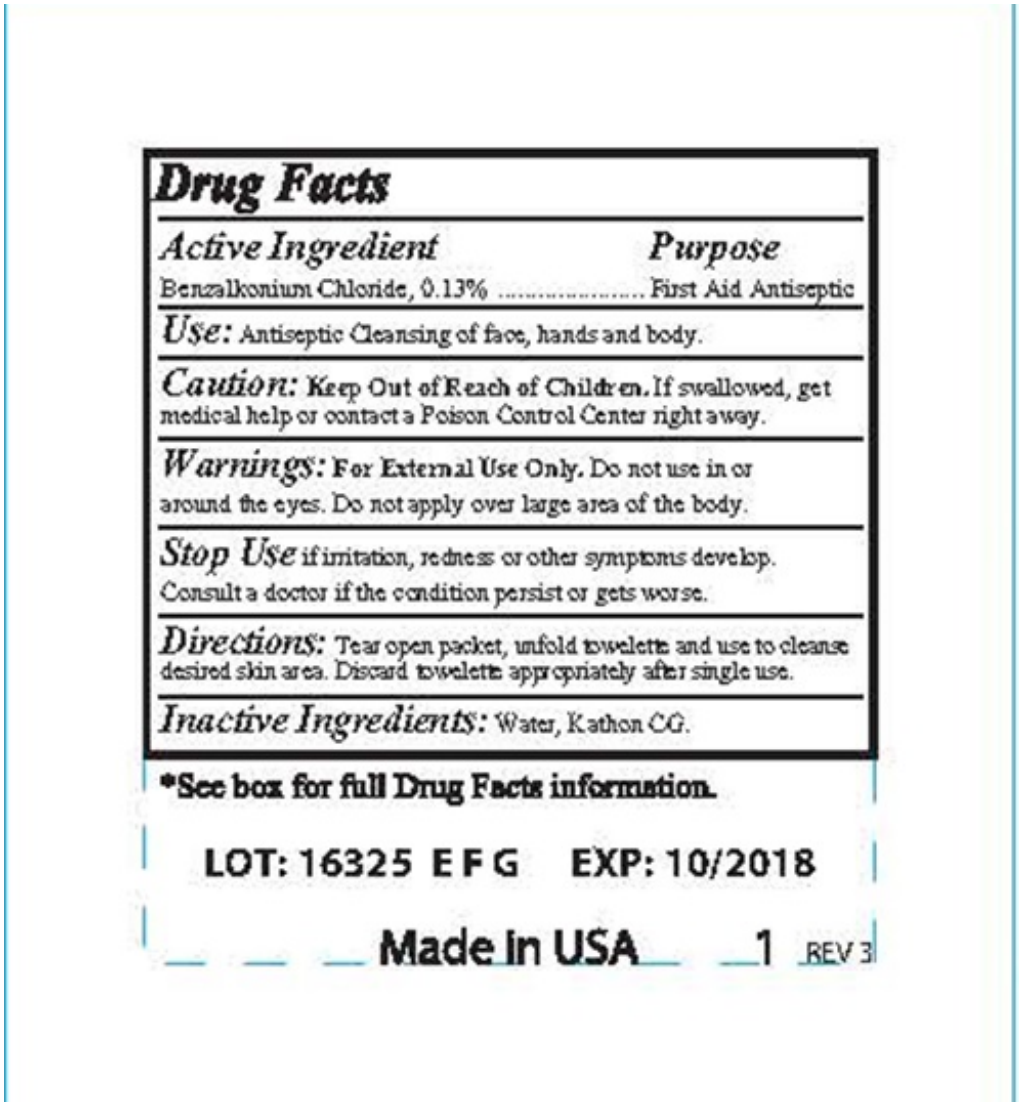

MED NAP

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59647-245
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1.7 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:59647-245-01	100 in 1 BOX	09/09/2013	
1	NDC:59647-245-00	1.7 mL in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:59647-245-02	1000 in 1 BOX	09/09/2013	
2	NDC:59647-245-00	1.7 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	09/09/2013	

Labeler - Med-Nap LLC (079086400)

Registrant - Med-Nap LLC (079086400)

Establishment

Name	Address	ID/FEI	Business Operations
Med-Nap LLC		079086400	manufacture(59647-245)

Revised: 11/2016 Med-Nap LLC