NORTHSIDE HOSPITAL HAND SANITIZING- 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.

Northside Hospital (hand san wipe) - 748

Drug Facts
Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

First Aid Antiseptic

Use:

Antiseptic Cleansing of face, hands and body.

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 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 appopriately after single use.

Inactive Ingredients:

Water, Methylchloroisothiazolinone, Methylisothiazolinone

NORTHSIDE HOSPITAL - Instant hand sanitizing wipes - product label

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NORTHSIDE HOSPITAL

Instant hands sanitizing wipes
Hand washing reduces Infections

Latex Free

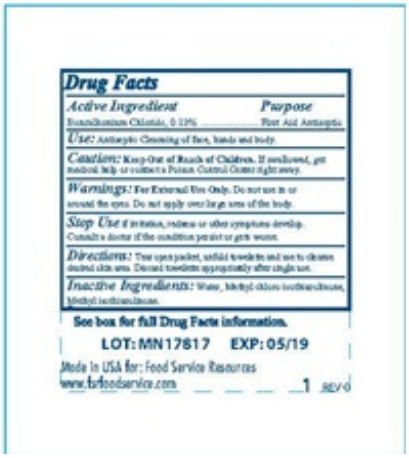
1 Pouch

NDC# 59647-748-01

See box for full Drug Facts information

LOT: MN17817 EXP: 05/19

Made in the USA for: Food Service Resources
www.fsrfoodservice.com 1 REV 0



NORTHSIDE HOSPITAL HAND SANITIZING

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59647-748
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)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59647-748-00	1000 in 1 CONTAINER	07/05/2017	
1	NDC:59647-748-01	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	part333A	07/05/2017	

Labeler - Med-Nap LLC (079086400)

Registrant - Med-Nap LLC (079086400)

Establishment

Name	Address	ID/FEI	Business Operations
Med-Nap LLC		079086400	manufacture(59647-748)

Revised: 7/2017 Med-Nap LLC