REDIWIPES ANTISEPTIC CLEANING TOWELETTES- bzk antiseptic wipes swab
Redicare 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.

BZK Wipes

Active Ingredients.....Uses

Benzalkonium Chloride 0.13%.....First Aid Antiseptic

Uses:

Antiseptic cleansing of face, hands and body to decrease bacteria on skin without soap and water. Antiseptic to help prevent further infection in minor cuts, scrapes, and burns.

Warnings

For external use only.

Do not use:
- In the eyes. If this happens, rinse thoroughly with water.
- Over large areas of the body
- Longer than 1 week unless directed by a doctor

Stop use and ask doctor if:
- Irritation of redness develop and persists for more than 72 hours.
- the condition persists or gets worse

Keep out of reach of children:

If swallowed, get medical help or contact a Poison Control Center right away.

Ask a doctor before use if you have:
- Deep or puncture wounds
- Animal bites
- Serious burns

Directions:

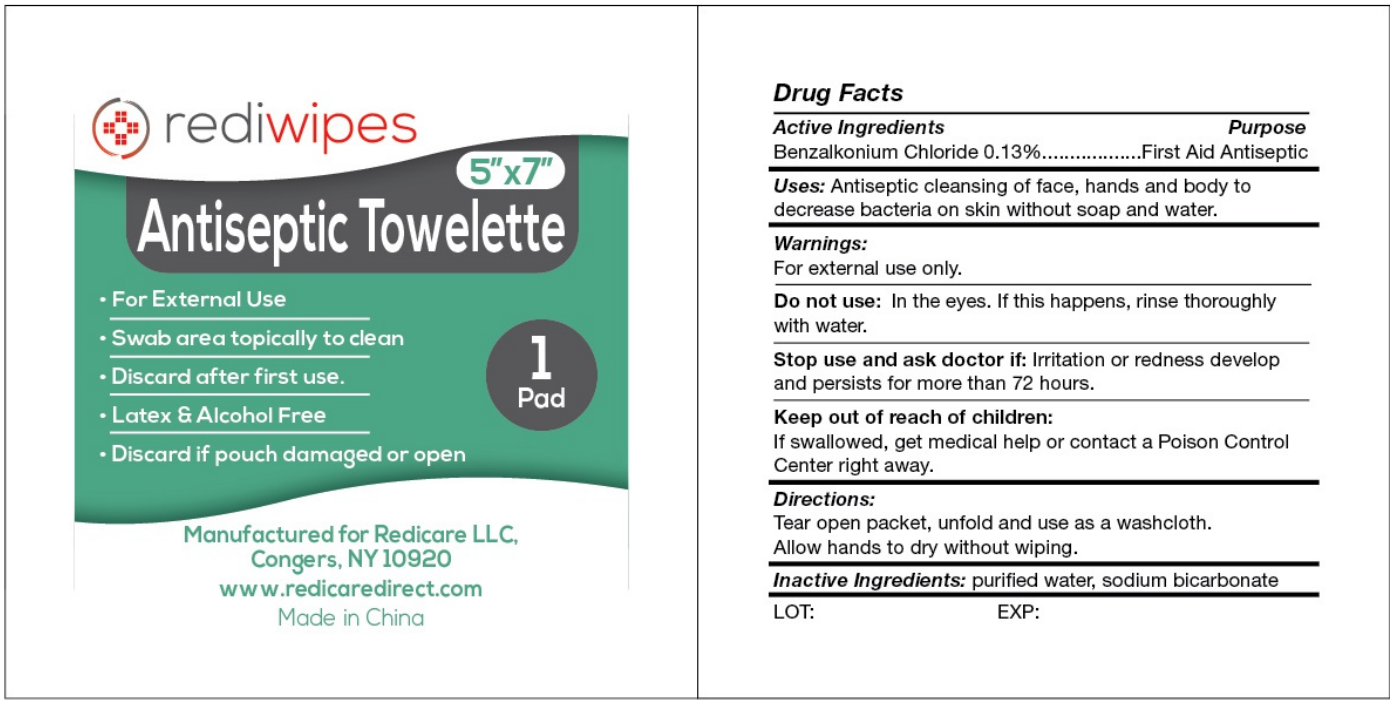
Tear open packet, unfold and use as a washcloth. Allow hands to dry without wiping.

Inactive Ingredients:

purified water, sodium bicarbonate

Other information:

- For external use only.
- Store at room temperature
- Do not use if packet is torn or opened



REDIWIPES ANTISEPTIC CLEANING TOWELETTES

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Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71105-323
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71105-323-03	1 in 1 BOX	03/31/2017	
1	NDC:71105-323-02	25 in 1 PACKET		
1	NDC:71105-323-01	1.5 g in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/31/2017	

Labeler - Redicare LLC (800149346)

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