

ANTISEPTIC WIPE- bzk antiseptic wipe liquid

Total Resources

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.

TR-1331 NDC 55550-215-70

Antiseptic Wipe

Active ingredient

Benzalkonium Chloride, 0.13%

Purpose

First Aid Antiseptic

Use

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

Warnings

Do not use in the eyes or apply over large areas of the body.

Stop use if

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

If swallowed, seek medical help or contact a Poison Control Center immediately.

Directions

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

Other information

Store at room temperature 59° - 86°F (15° - 30°C)

Inactive ingredient

water

Questions?

Call 1-888-771-7824

Label

NDC# 55550-210-70



ANTISEPTIC WIPE

Drug Facts

Active ingredient	Purpose
Benzalkonium Chloride, 0.13%	First Aid Antiseptic

Manufactured for:
TOTAL RESOURCES INT'L • WALNUT, CA 91789
00-TOW-94185 Rev.00

Drug Facts continued

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

Warnings

Do not use in the eyes or apply over large areas of the body.

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

Keep out of reach of children. If swallowed, seek medical help or contact a Poison Control Center immediately.

Directions

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

Other information

Store at room temperature 59° - 86°F (15° - 30°C)

Inactive ingredient water

Questions? Call 1-888-771-7824

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EXP:

ANTISEPTIC WIPE

bzk antiseptic wipe liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55550-215
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 mg in 1 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	1 mg in 1 mg

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55550-215-70	1000 in 1 CASE	06/19/2018	
1		1 mg 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	06/19/2018	

Labeler - Total Resources (790160535)

Revised: 1/2020

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