

**ANTISEPTIC TOWELETTE- benzalkonium chloride swab**  
**GFA Production (Xiamen) Co., Ltd.**

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**DRUG FACTS**

**Active Ingredient:**

Benzalkonium Chloride 0.13%

**Purpose:**

First Aid Antiseptic

**Use:**

For Professional and Hospital use. Helps prevent infection. Antiseptic cleansing of face, hands and body without soap and water.

**Warnings:**

For external use only.

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away. If unusual redness, swelling or other symptoms occur, consult a physician immediately.

**Do not use:**

In the eyes, or over large areas of the body.

**Directions:**

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

**Inactive ingredient:**

Purified water.

**Package Labeling:**

**DRUG FACTS - Antiseptic Towelette****Active Ingredient:** Benzalkonium Chloride 0.13%.. **Purpose:** First Aid Antiseptic**Use:** For Professional and Hospital use. Helps prevent infection. Antiseptic cleansing of face, hands and body without soap and water.**Warnings:** For external use only. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. If unusual redness, swelling or other symptoms occur, consult a physician immediately.**Do not use:** In the eyes, or over large areas of the body.**Directions:** Tear open packet, unfold towelette and use to cleanse desired skin area. Discard towelette appropriately after single use.**Inactive ingredient:** Purified water.

LOT/EXP:

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REORDER AST-001

**Antiseptic  
Towelette**

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**Toallitas  
Antisepticas**

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Suite B, 29 Harley Street  
LONDON W1G 9QR, England, United Kingdom**ANTISEPTIC TOWELETTE**

benzalkonium chloride swab

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50814-002
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50814-002-01	1 in 1 BOX	03/22/2016	
1		0.45 g in 1 POUCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
2	NDC:50814-002-02	1 in 1 BOX	01/01/2019	
2		0.25 g in 1 POUCH; 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 Drug	M003	03/22/2016	

**Labeler** - GFA Production (Xiamen) Co., Ltd. (421256261)

Revised: 10/2023

GFA Production (Xiamen) Co., Ltd.