

**BENZALKONIUM CHLORIDE- benzalkonium chloride swab**  
**Professional Disposables International, Inc.**

*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.*

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**HYGEA BENZALKONIUM CHLORIDE ANTISEPTIC TOWELETTE 10819-3740**

**USE**

For hand washing to decrease bacteria on the skin.

**PURPOSE**

Antiseptic handwash

**WARNINGS**

- **For external use only.**
- If swallowed, get medical help or contact a Poison Control Center right away.
- Discontinue use if irritation and redness develop. If condition persists for more than 72 hours consult a doctor.
- Do not use in the eyes.

**Keep out of reach of children.**

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**DIRECTIONS**

- Tear open packet.
- Unfold and use; discard in trash receptacle after single use.
- Wet hands thoroughly with product and allow to dry without wiping.

**OTHER INFORMATION**

Lot number and expiration date can be found on back panel of packet.

**INACTIVE INGREDIENTS**

Purified Water, Potassium Sorbate, PPG-2 Hydroxyethyl Cocamide, Disodium EDTA, Aloe Barbadensis Leaf Juice

**SPL UNCLASSIFIED SECTION**

**Professional Disposables International, Inc.**

Orangeburg, NY 10962-1376 - USA - 1-800-999-6423

Made in USA

**ACTIVE INGREDIENT**

Benzalkonium Chloride, 0.13% w/w

**DOSAGE**




Wet hands thoroughly with product and allow to dry without wiping.

**PRINCIPAL DISPLAY PANEL-100 COUNT**

**Hygea**  
**Benzalkonium Chloride**  
**Antiseptic Towelette**

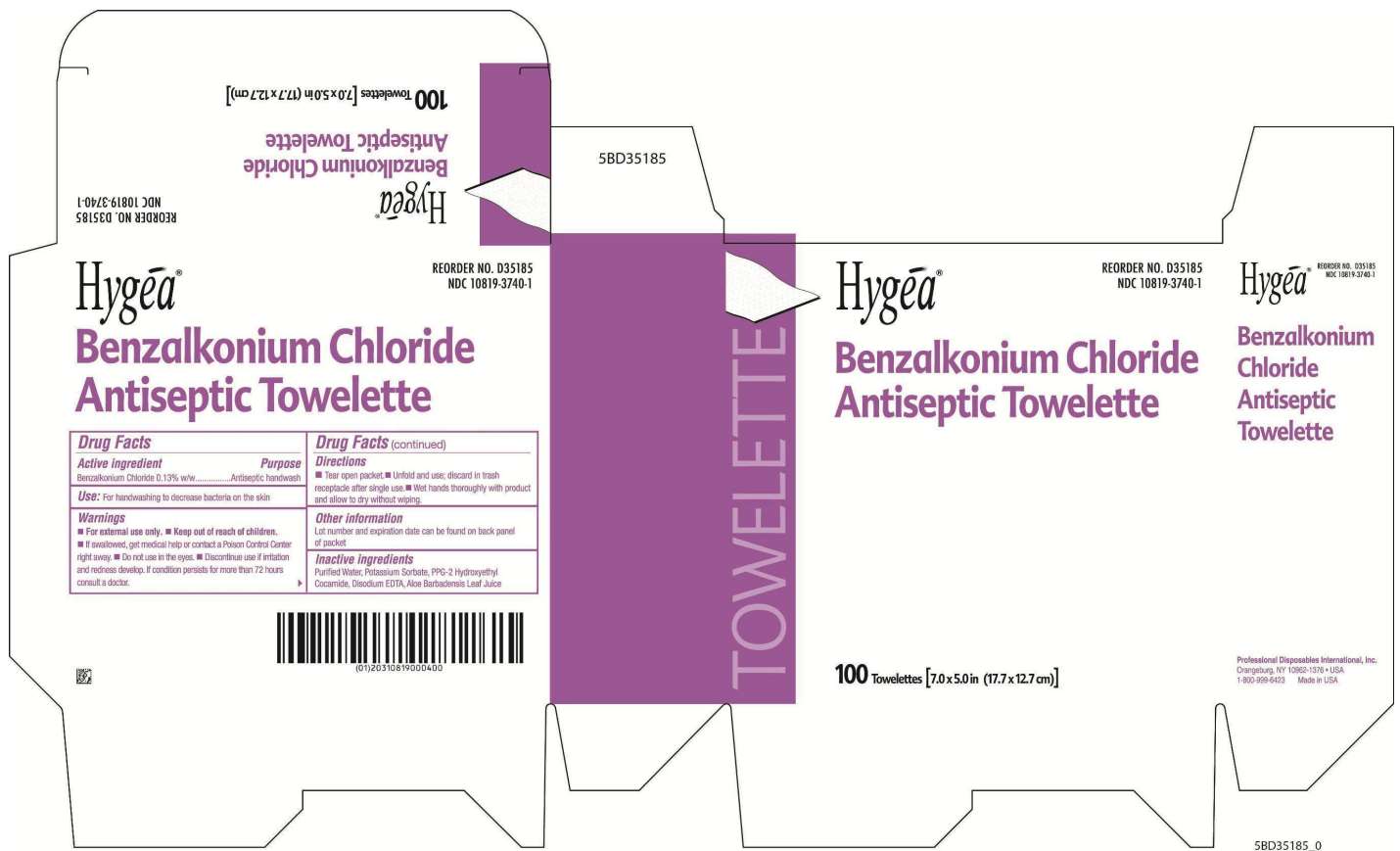
1 Towelette [7.0 x 5.0 in (17.7 x 12.7 cm)] 100 Towelettes per Box

Packet Label

	<p>REORDER NO. D351 NDC 10819-3740-1 <b>Benzalkonium Chloride</b> <b>Antiseptic Towelette</b></p>	 <small>(01)10310819000403</small>	<p><b>Drug Facts</b> (continued)</p>	<p>TEAR HERE</p>									
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Box Label



## BENZALKONIUM CHLORIDE

benzalkonium chloride swab

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10819-3740
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.0013 g in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:10819-3740-1	3 mL in 1 PACKET; Type 1: Convenience Kit of Co-Package	01/01/2012	
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>		<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph not final	part333A		01/01/2012	

**Labeler** - Professional Disposables International, Inc. (800777117)

**Establishment**

Name	Address	ID/FEI	Business Operations
Professional Disposables International, Inc.		800777117	manufacture(10819-3740)

Revised: 10/2019

Professional Disposables International, Inc.