

**ZIVA ANTIBACTERIAL MULTIPURPOSE WIPES- benzalkonium chloride cloth**  
**Ziva Wetwipes FZCo**

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**Drug Facts**

**Active Ingredient**

Benzalkonium Chloride 0.13%

**Direction of use:**

Open the flip top / peel back the front label slowly. Pull wipes as needed. Reseal pouch by firmly running thumb over label / close the flip top to prevent moisture loss.

**Precautions:**

Do not apply to irritated or damaged skin. In case of gluteal erythema (redness of skin), contact your doctor or pharmacist. Keep in a cool place away from direct sunlight. Prevent contact with eyes. In case of irritation wash away with water immediately.

**Ingredients:**

Aqua, Pentylene Glycol and Glyceryl Caprylate / Caprate, Dimethicone copolyol, PEG 40 Hydrogenated Castor Oil, Tocopheryl Acetate (Vitamin E), Glycerine, Sodium cocoamphodiacetate, Cocamidopropyl Betaine, Citric Acid, Disodium EDTA, Aloe Barbadosensis Leaf Juice.

**Product label**

<b>ZIVA ANTIBACTERIAL MULTIPURPOSE WIPES</b>			
benzalkonium chloride cloth			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:81450-002
<b>Route of Administration</b>	TOPICAL		
<b>Active Ingredient/Active Moiety</b>			
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
	<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

**Inactive Ingredients**

Ingredient Name	Strength
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)	
PEG/PPG-18/18 DIMETHICONE (UNII: 9H0A07T794)	
SODIUM COCOAMPHOACETATE (UNII: W7Q5E87674)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
WATER (UNII: 059QF0KO0R)	
HYDROGENATED CASTOR OIL (UNII: ZF94AP8MEY)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C00X)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81450-002-01	80 in 1 PACKAGE	12/24/2021	
1		4.59 mL in 1 POUCH; Type 0: Not a Combination Product		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	12/24/2021	

**Labeler** - Ziva Wetwipes FZCo (559186188)**Registrant** - Dirra USA (117678868)**Establishment**

Name	Address	ID/FEI	Business Operations
Ziva Wetwipes FZCO		559186188	manufacture(81450-002)

Revised: 10/2023

Ziva Wetwipes  
FZCo