

**XTRA- benzalkonium chloride spray**  
**MY IMPORTS USA LLC**

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**XTRA**

**Active ingredient**

Benzalkonium Chloride 0.13%

**Purpose**

Antibacterial

**Uses**

for handwashing, to reduce bacteria on skin

**Warnings**

For external use only.

**Keep out of reach of children**

Keep out of reach of children

**Do not use**

Do not use with chlorine bleach to avoid irritating fumes.

**Avoid contact with eyes.**

If eye contact occurs flush thoroughly with water.

**Do not ingest**

Do not ingest. If ingested, drink plenty of water, contact a physician.

Store in a cool and dry place, and avoid direct sunlight.

**Directions**

Spray to wet hand. Scrub and rinse thoroughly.

**Inactive ingredients**

Water (Aqua)

Sodium Laureth Sulfate

Decyl Glucoside  
Citric Acid  
Methylisothiazolone  
Phenoxyethanol  
Sodium Benzoate  
Fragrance

66.9\*104



**XTRA**

benzalkonium chloride spray

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:51628-4439
<b>Route of Administration</b>	SUPRACHOROIDAL		

**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 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>SODIUM LAURETH SULFATE</b> (UNII: BPV390UAP0)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>DECYL GLUCOSIDE</b> (UNII: Z17H97EA6Y)	
<b>CITRIC ACID</b> (UNII: 2968PHW8QP)	
<b>METHYLISOTHIAZOLINONE</b> (UNII: 229D0E1QFA)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>WATER</b> (UNII: 059QF0KO0R)	

**Product Characteristics**

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	ORANGE	<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51628-4439-1	399 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/27/2024	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	11/27/2024	

**Labeler - MY IMPORTS USA LLC (195767988)**

**Establishment**

Name	Address	ID/FEI	Business Operations
Zhejiang Jifu Daily Chemical Co., Ltd.		710111693	manufacture(51628-4439) , label(51628-4439)

