

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

-----  
**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>   | TOPICAL   |  |                             |                           |
| <b>Active Ingredient/Active Moiety</b>                                     |   |  |                             |                           |
| <b>Ingredient Name</b>   |   | <b>Basis of Strength</b>                                     | <b>Strength</b>             |                           |
| BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y) |   | BENZALKONIUM CHLORIDE  | 0.13 g in 100 mL            |                           |
| <b>Inactive Ingredients</b>  |   |  |                             |                           |
| <b>Ingredient Name</b>   |   |  | <b>Strength</b>             |                           |
| SODIUM LAURETH SULFATE (UNII: BPV390UAP0)                                  |   |  |                             |                           |
| PHENOXYETHANOL (UNII: HIE492ZZ3T)  |   |  |                             |                           |
| DECYL GLUCOSIDE (UNII: Z17H97EA6Y)   |   |  |                             |                           |
| CITRIC ACID (UNII: 2968PHW8QP)   |   |  |                             |                           |
| METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)                                   |   |  |                             |                           |
| SODIUM BENZOATE (UNII: OJ245FE5EU)   |   |  |                             |                           |
| WATER (UNII: 059QF0KO0R)   |   |  |                             |                           |
| <b>Product Characteristics</b>   |   |  |                             |                           |
| <b>Color</b>   |   | <b>Score</b>   |                             |                           |
| <b>Shape</b>   |   | <b>Size</b>  |                             |                           |
| <b>Flavor</b>  | ORANGE  | <b>Imprint Code</b>  |                             |                           |
| <b>Contains</b>  |   |  |                             |                           |
| <b>Packaging</b>   |   |  |                             |                           |
| <b>#</b>   | <b>Item Code</b>                                | <b>Package Description</b>                                   | <b>Marketing Start Date</b> | <b>Marketing End Date</b> |
| 1  | NDC:51628-4439-1                                | 399 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | 11/27/2024                  |                           |
| <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 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) |

