

**WIPE OUT ANTIBACTERIAL LIQUID HAND LEMON SCENT- benzalkonium chloride soap  
TZUMI INNOVATIONS LLC**

*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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**77878-030, Style: WP0024 LEM**

**Active Ingredient(s)**

Benzalkonium Chloride 0.13%

**Purpose**

Antibacterial

**Use**

For hand washing to decrease bacteria on the skin

**Warnings**

For external use only

When using this product do not get into eyes. If contact occurs, rinse eyes thoroughly with water

Stop use and ask a doctor if irritation and redness develop, condition persists for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

- Wet hands
- Apply palmful to hands
- Scrub thoroughly for at least 15 seconds
- Rinse thoroughly and dry

**Inactive ingredients**

Water, Cocamidopropyl dimethylamine oxide, Citric Acid, Fragrance, Kathon, Glycerin, Fatty acyl diethanol amine, Disodium EDTA, Hydroxyethyl Cellulose (250HHR), Red 7 (CI 16255), Yellow 5 (CI 19140)

LABEL - FRONT



LABEL - BACK



**WIPE OUT ANTIBACTERIAL LIQUID HAND LEMON SCENT**

benzalkonium chloride soap

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:778 78-030
<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 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

DISODIUM HEDTA (UNII: KME849MC7A)	
HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
D&C RED NO. 7 (UNII: ECW0LZ41X8)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77878-030-01	221 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/23/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/23/2020	

**Labeler** - TZUMI INNOVATIONS LLC (117426322)

### Establishment

Name	Address	ID/FEI	Business Operations
Aogrand International Trade Corporation		421353092	manufacture(77878-030)

Revised: 12/2020

TZUMI INNOVATIONS LLC