# G AND Y ANTIBACTERIAL WET WIPES- benzalkonium chloride cloth ERUSLU SAGLIK URUNLERI SANAYI VE TICARET ANONIM SIRKETI

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.

-----

#### **G&Y Antibacterial Wet Wipes**

#### **Drug Facts**

#### Active ingredients

Benzalkonium Chloride 0.13% w/w

#### **Purpose**

**Antibacterial** 

#### Uses

For handwashing to decrease bacteria on the skin

### Warning

# For external use only

#### Do not use

- in the eyes.
- if you are allergic to any of the ingredients.

When using this product if eye contact occurs, rinse eyes thoroughly with water.

**Stop use and ask a doctor** if irritation and redness develop and persist 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**

- Storage. Store at room temperature.
- **Dispensing**. Remove seal band. Lift tab at front of lid. Pull out wipe and reseal lid. Close lid to retain moisture.
- Use. Apply wipe thoroughly to hands as desired. Allow to dry without wiping.
- Disposal. Dispose of used wipes in trash receptacle after use. Do not flush.

#### Other information

Production Date, Expiry Date and Lot Number on side

#### Inactive ingredients

Benzoic Acid, C12-15 Pareth-12, Dehydroacetic Acid, Fragrance, Glycerin, Phenoxyethanol, Purified Water, Tetrasodium Glutamate Diacetate.

Kills 99.9% of Germs that may cause illness.

6.3 IN x 7.1 IN (16 cm x 18 cm)

Questions? +1 (862) 257-3339

You may also report serious side effects to this phone number.

Mon-Fri 9:00 AM - 5:00 PM

Distributed by: G&Y Products, Inc.

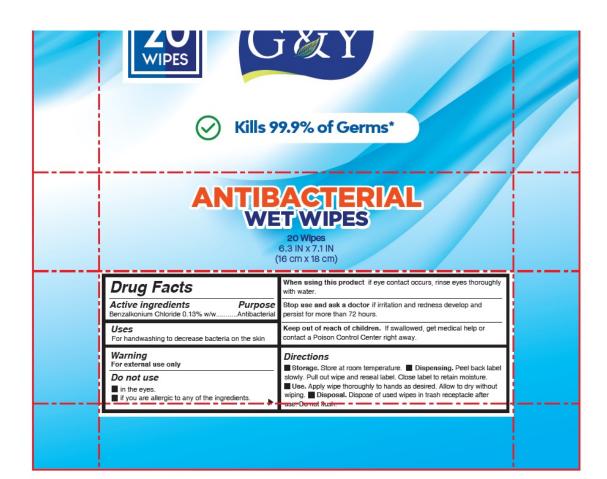
25 Shady St, Paterson, NJ 07524

info@gyproduct.com

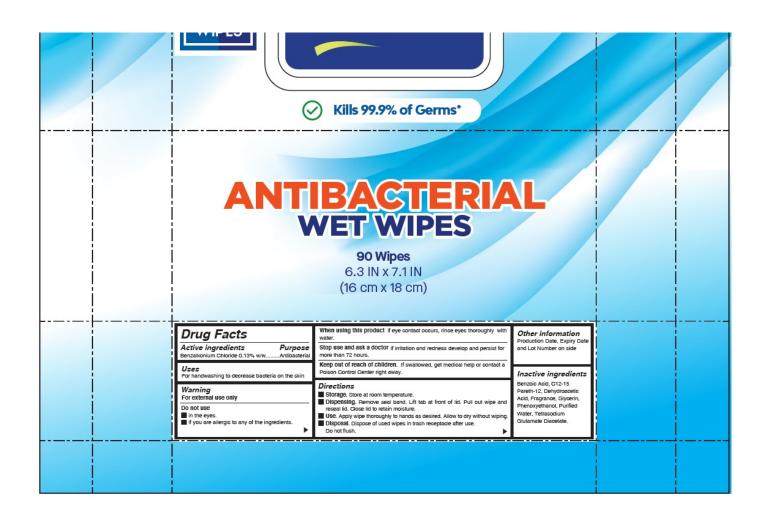
**Country of origin: Turkey** 

## **Packaging**









## **G AND Y ANTIBACTERIAL WET WIPES**

benzalkonium chloride cloth

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:77613-011

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name	<b>Basis of Strength</b>	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

# Inactive Ingredients Ingredient Name Strength BENZOIC ACID (UNII: 8SKN0B0MIM) C12-15 PARETH-12 (UNII: 131316X18L) DEHYDROACETIC ACID (UNII: 2KAG279R6R) GLYCERIN (UNII: PDC6A3C0OX) PHENOXYETHANOL (UNII: HIE492ZZ3T) WATER (UNII: 059QF0K00R) TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:77613-011- 20	20 in 1 PACKAGE	05/14/2020				
1		3.5 mL in 1 PACKET; Type 0: Not a Combination Product					
2	NDC:77613-011- 90	90 in 1 PACKAGE	05/14/2020				
2		3.5 mL in 1 PACKET; Type 0: Not a Combination Product					

eting Start Marketing End Date Date
220

# **Labeler -** ERUSLU SAGLIK URUNLERI SANAYI VE TICARET ANONIM SIRKETI (565415460)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
ERUSLU SAGLIK URUNLERI SANAYI VE TICARET ANONIM SIRKETI		565415460	manufacture(77613-011)				

Revised: 2/2022 ERUSLU SAGLIK URUNLERI SANAYI VE TICARET ANONIM SIRKETI