G AND Y ANTIBACTERIAL HAND 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 Hand 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

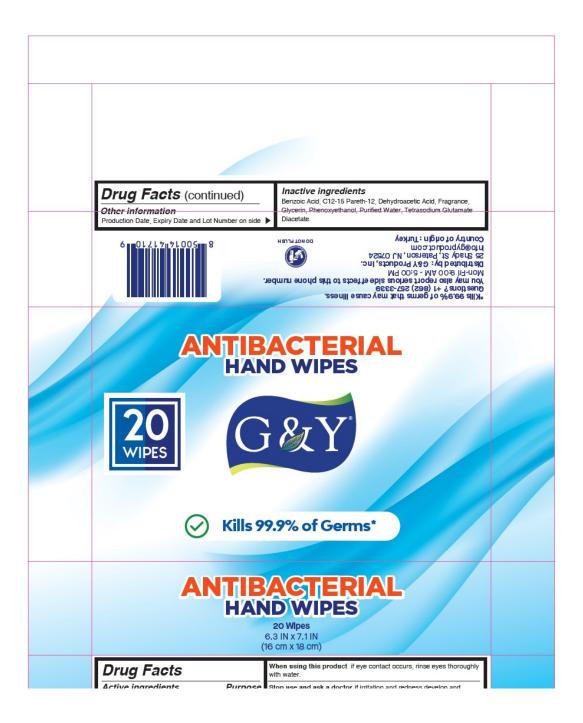
info@gyproduct.com

Country of origin : Turkey

Packaging



	90 Wipes 6.3 IN x 7.1 IN (16 cm x 18 cm)		
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G AND Y ANTIBACTERIAL HAND WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77613-010
Route of Administration	TOPICAL		

6.6	
of Strength	Strength
	0.13 g in 100 mL

Inactive Ingredients

		Ingredient Name		Strength
BE	NZOIC ACID (UNII:	8 S K N0 B 0 MIM)		
C12	2-15 PARETH-12 (U	INII: 131316X18L)		
DE	HYDRO ACETIC AC	CID (UNII: 2KAG279R6R)		
GL	YCERIN (UNII: PDC	6A3C0OX)		
PH	ENO XYETHANO L	(UNII: HIE492ZZ3T)		
WA	ATER (UNII: 059QF0	KO0R)		
TE	TRASODIUM GLU	FAMATE DIACETATE (UNII: 5EHL5014MY)		
Pa	ckaging			
#	Item Code	Package Description	Marketing Start Date	Markating End Dat

I NDC://613-010-20	20 in 1 PACKAGE	05/14/2020			
1	3.5 mL in 1 PACKET; Type 0: Not a Combination Product				
2 NDC:77613-010-90	90 in 1 PACKAGE	05/14/2020			
2	3.5 mL in 1 PACKET; Type 0: Not a Combination Product				
Marketing Information					
Marketing Inf	ormation				
Marketing Inf Marketing Catego		Marketing Start Date	Marketing End Date		
	y Application Number or Monograph Citation	Marketing Start Date 05/14/2020	Marketing End Date		

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

Establishment

Name	Address	ID/FEI	Business Operations
ERUSLU SAGLIK URUNLERI SANAYI VE TICARET ANONIM SIRKETI		565415460	manufacture(77613-010)

Revised: 5/2020

ERUSLU SAGLIK URUNLERI SANAYI VE TICARET ANONIM SIRKETI