G AND Y DISINFECTING WET WIPES- g and y disinfecting wet wipes cloth G&Y PRODUCTS INC.

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.

Active Ingredient(s)

Benzalkonium Chloride 0.2% w/w

Purpose

Antimicrobial

Uses

For handwashing to decrease bacteria on the skin.

Warnings

For external use only.

Do not use

Do not use

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

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

Stop use and ask a doctor if irritation and redness develop and persist more than 72 hours.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Cloth

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

Inactive ingredients

Benzoic Acid, C12-14-Alkyldimethyl (Ethylbenzyl) Ammonium Chloride, C12-15 Pareth-12, Dehydroacetic Acid, Didecyl Dimethyl Ammonium Chloride, Ethyl Alcohol, Fragrance, Glycerin, Phenoxyethanol, Purified Water, Tetrasodium Glutamate Diacetate,

Other information

Production Date, Expiry Date and Lot Number on side.

Package Label

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

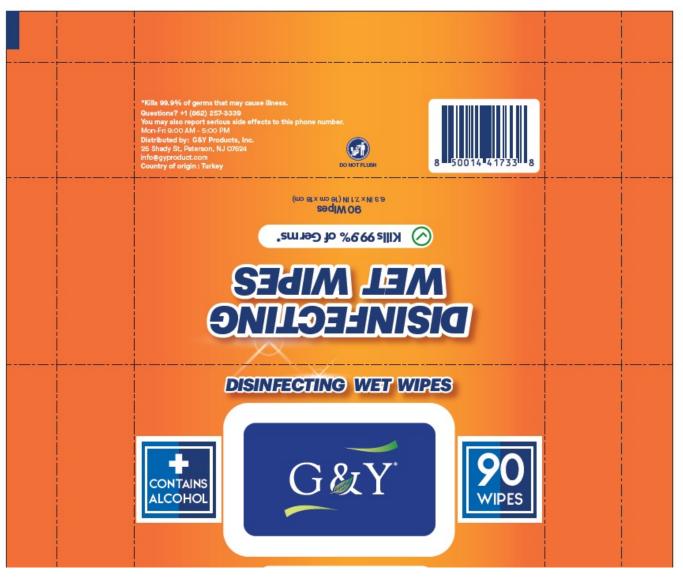
Distributed by: G&Y Products, Inc.

25 Shady St, Paterson, NJ 07524

info@gyproduct.com

Country of origin: Turkey

Packaging



Drug Facts

Drug Facts	When using this product if eye contact occurs, rinse eyes thoroughly with water.	Other information Production Date, Expiry Date	
Active ingredients Purpose Benzalkonium Chloride 0.2% w/wAntimicrobial	Stop use and ask a doctor if irritation and redness develop and persist for more than 72 hours.	and Lot Number on side	
Uses For handwashing to decrease bacteria on the skin	Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	Inactive ingredients Benzoic Acid, C12-14-Alky- Idimethyl(Ethylbenzyl)Ammoni- um Chloride, C12-15 Pareth-12, Dehydroacetic Acid, Didecyl Dimethyl Ammonium Chloride,	
Warning	Directions Storage, Store at room temperature.		
For external use only	Dispensing, Remove seal band. Lift tab at front of lid. Pull out wipe and	Dimethyl Ammonium Chloride	

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Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:80		0903-012		
Route of Administration	TOPICAL					
Active Ingredient/Active Moiety						
Ingredient Name			Basis of Strength	Strength		
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)			BENZALKONIUM CHLORIDE	0.2 g in 3.8 mL		

Inactive Ingredie	nts		
	Ingredient Name		Strength
ALCOHOL (UNII: 3K9	958V90M)		
GLYCERIN (UNII: PDC	C6A3C0OX)		
PHENO XYETHANO L	(UNII: HIE492ZZ3T)		
TETRASODIUM GLU	TAMATE DIACETATE (UNII: 5EHL5014MY)		
N-ALKYL ETHYLBEN	NZYL DIMETHYL AMMONIUM CHLORIDE (C12	-C14) (UNII: G258TFN61X)	
DIDECYL DIMO NIUM	CHLORIDE (UNII: JXN40O9Y9B)		
WATER (UNII: 059QF)KO0R)		
BENZOIC ACID (UNII	8SKN0B0MIM)		
	JNII: 131316X18L)		
C12-15 PARETH-12 (U	JNII: 131316 X18 L) C ID (UNII: 2KAG279 R6 R)		
C12-15 PARETH-12 (U	,		
C12-15 PARETH-12 (U DEHYDRO ACETIC AG Packaging	,	Marketing Start Date	• Marketing End Dat
C12-15 PARETH-12 (T DEHYDROACETIC A Packaging # Item Code	CID (UNII: 2KAG279R6R) Package Description	Marketing Start Date	• Marketing End Dat
C12-15 PARETH-12 (C DEHYDROACETIC AC Packaging # Item Code 1 NDC:80903-012-90	CID (UNII: 2KAG279R6R) Package Description	07/03/2020	e Marketing End Dat
C12-15 PARETH-12 ((DEHYDROACETIC AC Packaging	CID (UNII: 2KAG279R6R) Package Description 90 in 1 PACKAGE	07/03/2020	e Marketing End Dat
C12-15 PARETH-12 (U DEHYDROACETIC AC Packaging # Item Code 1 NDC:80903-012-90	CID (UNII: 2KAG279R6R) Package Description 90 in 1 PACKAGE	07/03/2020	• Marketing End Dat
C12-15 PARETH-12 (U DEHYDROACETIC AC Packaging # Item Code 1 NDC:80903-012-90 1	CID (UNII: 2KAG279R6R) Package Description 90 in 1 PACKAGE 3.8 mL in 1 PACKAGE; Type 0: Not a Combination	07/03/2020	e Marketing End Dat
C12-15 PARETH-12 (U DEHYDROACETIC AC Packaging # Item Code 1 NDC:80903-012-90	CID (UNII: 2KAG279R6R) Package Description 90 in 1 PACKAGE 3.8 mL in 1 PACKAGE; Type 0: Not a Combination ormation	07/03/2020 n Product	Marketing End Dat

Labeler - G&Y PRODUCTS INC. (081236664)

Registrant - G AND Y PRODUCTS INC. (081236664)

Establishment

Name	Address	ID/FEI	Business Operations
G&Y PRODUCTS INC.		081236664	manufacture(80903-012), label(80903-012)

Revised: 12/2020

G&Y PRODUCTS INC.