2XL MEGA ROLL SANITIZING WIPES 2300 COUNT MEGAROLL- benzalkonium chloride cloth 2XL Corporation

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.

2xl Mega Roll sanitizing wipes 2300 count megaroll

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

- to sanitize hands without requiring water or a rinse
- kills 99.9% of most common germs

Warnings

For external use only

When using this product

- do not use in or near eyes
- discontinue use if irritation and redness develop

Keep out of reach of children.

In case of accidental ingestion, seek medical attention or contact a poison control center immediately.

Directions

- wet hands thoroughly with wipe
- allow to dry without rinsing

Other information

• store at room temperature

Inactive ingredients

Water, Phenoxyethanol, Decyl Glucoside, Potassium Sorbate, Sodium Benzoate, Disodium EDTA, Citric Acid, Aloe Barbadensis (Aloe) Leaf Extract, Fragrance.

Questions? Comments?

Call (888) 977-3726

Package Labeling:

Kills 99.9% of most common germs that may cause illness



- Quickly removes soil and kills most common germs that may cause illness
- Textured for superior cleaning
- Sanitizes hands while cleaning effectively
- Convenient, easy to use and ideal for placement in offices, restaurants, health clubs or anywhere else germs may be

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NET CONTENTS 2,300 Wipes per Roll (8" x 5")

Net weight: 5460 g/roll

Distributed by: 2XL Corporation 7550 Industrial Drive Forest Park, IL 60130 888-977-3726 www.2xlcorp.com

NDC 71995-020-01

2XL-422 Refill

2XL MEGA ROLL SANITIZING WIPES 2300 COUNT MEGAROLL

benzalkonium chloride cloth

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HUMAN OTC DRUG NDC:71995-002 Item Code (Source) **Product Type TOPICAL Route of Administration**

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZ ALKONIUM CHLORIDE	1.3 mg in 1 mL			

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
PHENOXYETHANOL (UNII: HIE492ZZ3T)		
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)		
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		

F	Packaging						
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:71995- 002-00	2300 in 1 POUCH	01/21/2014				
1		5 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)					

Marketing Information					
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
part333E	01/21/2014				
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Labeler - 2XL Corporation (148004059)

Revised: 11/2022 2XL Corporation