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:

2XL MEGA ROLL SANITIZING WIPES 2300 COUNT MEGAROLL

benzalkonium chloride cloth

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

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM 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)	

Packaging				
#	**Item 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

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/21/2014	

Labeler - 2XL Corporation (148004059)

Revised: 11/2022 2XL Corporation