GERM OUT ANTIBACTERIAL WET WIPES- benzalkonium chloride cloth
Flex Beauty Labs LLC

Germ Out Antibacterial Wet Wipes

DRUG FACTS

Active Ingredients

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Use

To reduce bacteria on the skin.

Warnings

For external use only.

DO NOT USE

if you are allergic to any of the ingredients.

When using this product

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or redness develops.

Keep out of reach of children.

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

Directions

Adults and children over 2 years of age: Apply to hands, Allow to dry without wiping.
Children under 2 years of age: ask doctor prior to use.

Other information:

Store below 110°F(43°C). May discolor certain fabrics or surfaces.

Inactive Ingredients:

Water (Aqua), Glycerin, Phenoxyethanol, Polysorbate 20, Sodium PCA, Fragrance (Parfum), Ethylhexylglycerin, Tocopheryl Acetate (Vitamin E), Aloe Barbadensis Leaf Extract, Tetrasodium EDTA, Citric Acid.

Package Labeling:20ct

Package Labeling:60ct

GERM OUT ANTIBACTERIAL WET WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72308-025
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 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72308-025-20	20 in 1 PACKET	08/15/2020	
1		3.2 g in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
2	NDC:72308-025-60	60 in 1 PACKET	08/15/2020	
2		3.2 g 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 Drug	505G(a)(3)	08/15/2020	

Labeler - Flex Beauty Labs LLC (080858917)

Revised: 12/2023 Flex Beauty Labs LLC