GOODDAY CLEAN WET WIPES- benzalkonium chloride cloth Maxkorea Co., Ltd.

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

Benzalkonium Chloride 0.2%

INACTIVE INGREDIENTS

Water, Sodium hydroxide, Glycerin, Yellow 4, Blue 1

PURPOSE

Antiseptic

WARNINGS

For external use only

Do not use

- on infants
- on open skin wounds

When using this product

- avoid getting into the eyes
- In case of eye contact immediately flush eyes thoroughly with water

Stop use and ask a doctor

- if irritation or redness develops
- conditions persist for more than 72 hours
- redness is present

KEEP OUT OF REACH OF CHILDREN

■ in case of accidental ingestion, contact a doctor or Poison Center immediately

Uses

■ hand sanitizing wipes to help reduce bacteria on the skin. For use when soap and water not available

Directions

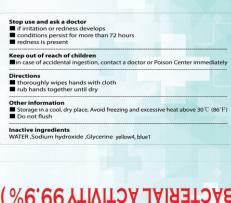
- thoroughly wipes hands with cloth
- rub hands together until dry

Other information

■ Storage in a cool, dry place, Avoid freezing and excessive heat above 30 [1] (86 [1])

■ Do not flush

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL





ANTIBACTERIAL WIPES (ANTIBACTERIAL ACTIVITY 99.9%) Sterilization, deodorization and cleaning at the same time!!

STERILIZATION AND THE WET TISSUE FOR DISINFECTION







Sterilizing Antiseptic

20 pcs



Sterilization, deodorization and cleaning at the same time!! 3 Protect your health from harmful bacteria

VIRUS OUTBREAK WET WIPES















Drug Facts Active ingredients Benzalkonium Chloride 0.2% Purpose Antiseptic Uses

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Drug facts continued







GOODDAY CLEAN WET WIPES

benzalkonium chloride cloth

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:79907-010

Route of Administration TOPICAL

Active Ingredient/Active Moiety

1	, 		
	Ingredient Name	Basis of Strength	Strength
	Benzalkonium Chloride (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	0.2 in 100

Inactive Ingredients

8			
Ingredient Name	Strength		
Water (UNII: 059QF0KO0R)			
SO DIUM HYDRO XIDE (UNII: 55X0 4QC32I)			
Glycerin (UNII: PDC6 A3C0 O X)			

Glycerin (UNII: PDC6A3C0OX)

Packaging

	# Item Code		Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:79907-010-01	20 in 1 PACKET; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/01/2020	

Labeler - Maxkorea Co., Ltd. (694487095)

Registrant - Maxkorea Co., Ltd. (694487095)

Establishment			
Name	Address	ID/FEI	Business Operations
Maxkorea Co., Ltd.		694487095	manufacture(79907-010)

Revised: 7/2020 Maxkorea Co., Ltd.