SANITIZING HAND WIPES MOISTURIZE WITH ALOE EXTRACT- benzalkonium chloride cloth
COMMONWEALTH WHOLESALE CORP

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.

CWC Wet Wipes Package

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic

Use

■ Hand sanitizer to help reduce bacteria on the skin. For use when soap and water not available

Warnings

For external use only

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 Control Center immediately.

Directions

■ Thoroughly wipe hands with cloth

■ Rub hands together until dry

■ No rinsing required

Other information

■ Dispose of properly, do not flush

Inactive ingredients

water, polysorbate 20, glycerin, potassium sorbate, aloe (aloe barbadensis) leaf extract, dihydroacetic acid, phenoxyethanol, citric acid

Package Label

SANITIZING HAND WIPES MOISTURIZE WITH ALOE EXTRACT

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77642-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

POLYSORBATE 20 (UNII: 7T1F30V5YH)	
GLYCERIN (UNII: PDC6A3C0OX)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77642-001-20	20 in 1 PACKAGE; Type 0: Not a Combination Product	05/18/2020	
2	NDC:77642-001-50	50 in 1 PACKAGE; Type 0: Not a Combination Product	06/29/2020	
3	NDC:77642-001-40	400 in 1 PAIL; Type 0: Not a Combination Product	06/29/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/18/2020	

Labeler - COMMONWEALTH WHOLESALE CORP (079623736)

Revised: 6/2020 COMMONWEALTH WHOLESALE CORP