RENEW ANTIBACTERIAL HAND WIPES- benzalkonium chloride cloth
Kairon 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.

Renew Antibacterial Hand Wipes

Active Ingredient

Benzalkonium Chloride 0.13% v/v. Purpose: Antiseptic

Purpose

Antiseptic

Use

helps decrease bacteria on the skin

Warnings

Flammable. Keep away from fire or flame. For external use only

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 if irritation or rash appears and lasts.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Do not use

on children under 2 months of age or on open skin wounds

Directions

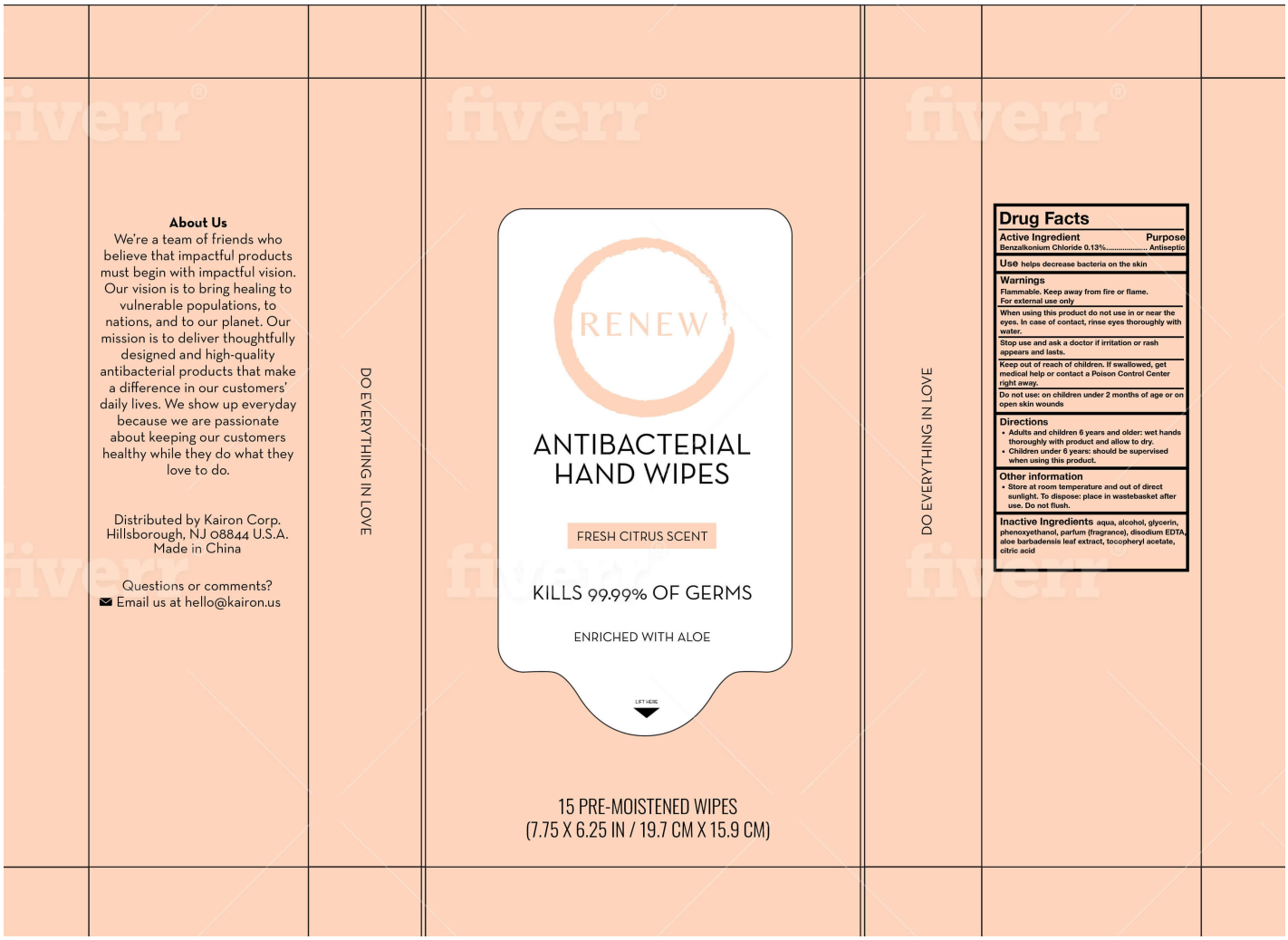
- Adults and children 6 years and older: wet hands thoroughly with product and allow to dry.
- Children under 6 years: should be supervised when using this product.

Other information

- Store at room temperature and out of direct sunlight. To dispose: place in wastebasket after use. Do not flush.

Inactive ingredients

aqua, alcohol, glycerin, phenoxyethanol, parfum (fragrance), disodium EDTA, aloe barbadensis leaf juice, tocopheryl acetate, citric acid



15 PRE-MOISTENED WIPES

NDC: 78899 -001-01

RENEW ANTIBACTERIAL HAND WIPES

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78899-001
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 in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
PHENOXYETHANOL (UNII: HIE492ZZ3T)	0.3 g in 100 g
GLYCERIN (UNII: PDC6A3C0OX)	0.5 g in 100 g

WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	10 g in 100 g
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	0.05 g in 100 g
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	0.01 g in 100 g
ALOE VERA LEAF (UNII: ZY81Z83H0X)	0.05 g in 100 g
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	0.01 g in 100 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78899-001-01	100 g in 1 PACKAGE; Type 0: Not a Combination Product	08/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/01/2020	

Labeler - Kairon Corp. (117413044)

Registrant - Kairon Corp. (117413044)

Establishment

Name	Address	ID/FEI	Business Operations
Kairon Corp.		117413044	manufacture(78899-001)

Revised: 6/2020 Kairon Corp.