

CARESOUR ANTIBACTERIAL WIPES LEMON SCENTED- benzalkonium chloride solution
Onecare Products LLC

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.

Caresour

Active Ingredient

Benzalkonium Chloride 0.3%

Purpose

Antiseptic

USE

For hand sanitizing to decrease bacteria on the skin.

Apply topically to the skin to help prevent cross contamination.

Recommended for repeated use.

Dries in seconds.

Warning

● For external use only.

Do not use in or contact the eyes.

● Stop use and ask a doctor if too much skin irritation or sensitivity develops or increases

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

Directions

● Gently pull back reusable label, remove and use wipe as required.

● Reseal back after use

Other Information

Lot No, Manufacture date, and Expiration date can be found on package

Inactive ingredients

Water, Phenoxyethanol, DMDM Hydantoin, Ethylparaben, Methylparaben, Polysorbate 20, Fragrance, Disodium EDTA, Sodium Citrate, Cetylpyridinium Chloride

Distributed by: Onecare Products LLC.
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www.onecareus.com



Item Code: #W-076 Made in China



50 Wipes
6 in x 8 in (15.2 cm x 20.3 cm)

Lemon Scent

Antibacterial
Wipes

Caresour



Alcohol Free

50 Wipes
6 in x 8 in (15.2 cm x 20.3 cm)

Lemon Scent

Antibacterial
Wipes

Caresour



Caresour

Antibacterial
Wipes

Lemon Scent

50 Wipes

6 in x 8 in (15.2 cm x 20.3 cm)



Drug Facts

Active Ingredient Benzalkonium chloride 0.3%.....**Purpose** Antiseptic

Uses
• For hand sanitizing to decrease bacteria on the skin.
• Apply topically to the skin to help prevent cross contamination.
• Recommended for repeated use.
• Dries in seconds.

Warnings
• For external use only.
• Do not use in or contact the eyes.
• Stop use and ask a doctor if too much skin irritation or sensitivity develops or increases.

Drug Facts (continued)

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

Directions
• Gently pull back resealable label, remove and use wipe as required.
• Reseal back after use.

Other information
• Lot No. Manufacture date and Expiration date can be found on package.

Inactive Ingredients
Water, Phenoxyethanol, DMDM Hydantoin, Ethylparaben, Methylparaben, Polysorbate 20, Fragrance, Disodium EDTA, Sodium Citrate, Cetylpyridinium Chloride.

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75513-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	9.6 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
ETHYLPARABEN (UNII: 14255EXE39)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
CETYLPIRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
DISODIUM HEDTA (UNII: KME849MC7A)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75513-002-01	1 in 1 POUCH; Type 0: Not a Combination Product	06/15/2020	
2	NDC:75513-002-02	10 in 1 POUCH; Type 0: Not a Combination Product	06/15/2020	
3	NDC:75513-002-03	25 in 1 POUCH; Type 0: Not a Combination Product	06/15/2020	
4	NDC:75513-002-04	30 in 1 POUCH; Type 0: Not a Combination Product	06/15/2020	
5	NDC:75513-002-05	50 in 1 POUCH; Type 0: Not a Combination Product	06/15/2020	
6	NDC:75513-002-06	80 in 1 POUCH; Type 0: Not a Combination Product	06/15/2020	
7	NDC:75513-002-07	100 in 1 POUCH; Type 0: Not a Combination Product	06/15/2020	
8	NDC:75513-002-08	70 in 1 CANISTER; Type 0: Not a Combination Product	06/15/2020	
9	NDC:75513-002-09	100 in 1 CANISTER; Type 0: Not a Combination Product	06/15/2020	
10	NDC:75513-002-10	300 in 1 CANISTER; Type 0: Not a Combination Product	06/15/2020	
11	NDC:75513-002-11	800 in 1 PACKAGE; Type 0: Not a Combination Product	06/15/2020	
12	NDC:75513-002-12	1200 in 1 PACKAGE; Type 0: Not a Combination Product	06/15/2020	
13	NDC:75513-002-13	1500 in 1 PACKAGE; Type 0: Not a Combination Product	06/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/14/2020	

Labeler - Onecare Products LLC (129515451)

Establishment

Name	Address	ID/FEI	Business Operations
Kangna (Zhejiang) Medical Supplies Co., Ltd.		554530173	manufacture(75513-002)

Revised: 6/2020

Onecare Products LLC