

**BENZALKONIUM CHLORIDE- sanitizing wipes cloth**  
**Purocor 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.*

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**Sanitizing Wipes**

***Active ingredients***

Benzalkonium Chloride 0.13%

***Purpose***

Antiseptic

***Use(s)***

Hand Sanitizer to help reduce bacteria that potentially can cause disease.

***Warnings***

**For external use only.**

**Do not use**

- In children less than 2 months of age
- On open skin wounds

**When using this product**

keep out of eyes, ears and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

**Stop use and ask doctor**

If irritation or rash appears.

**Keep out of reach of children.**

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

***Directions***

- Place enough product on hands to cover all surfaces. Rub hands together until dry.

- Supervise children under 6 years of age when using this product to avoid swallowing.

***Other information***

- Store between 15-30°C (59-86°F)

***Inactive ingredients***

Aloe Barbadensis Leaf Extract, Citric Acid, Decyl Glucoside, Disodium EDTA, Fragrance, Phenoxyethanol, Potassium Sorbate, Purified Water, Sodium Benzoate

**Principal Display Panel**

PuroCor

CLINICAL STRENGTH FORMULA

SANITIZING

WIPES

1200 WET WIPES

Lemon Scent

ALCOHOL

FREE

8" X 5.65" (20.3cm x 14.4cm)



CLINICAL STRENGTH FORMULA

# SANITIZING WIPES



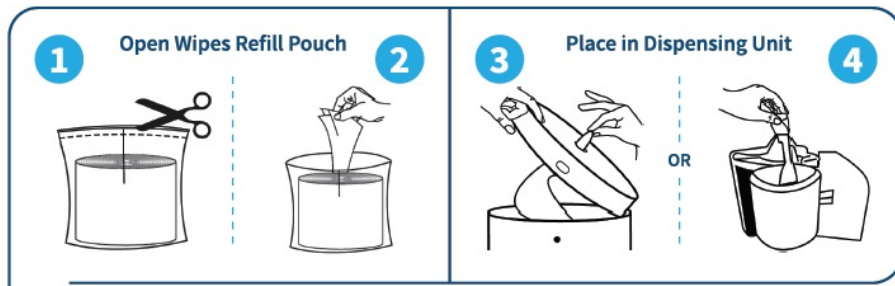
**1200 WET WIPES**  
*Lemon Scent*



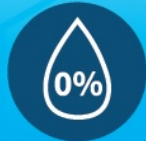
**11 lb**

8" x 5.65" (20.3cm x 14.4cm)





1. Cut open top of bag.
2. Pinch center wipe from middle of roll and pull up then place roll into dispenser.
3. Pass first wipe through opening of dispenser.
4. Securely close cover of dispenser.



ALCOHOL FREE



SOFT ON HANDS



NO ANIMAL TESTING



LEMON SCENT

**Drug Facts**

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**Distributed by:**  
PuroCor LLC  
8 The Green  
Suite#12589  
Dover, DE 19901  
[www.purocor.com](http://www.purocor.com)

**BENZALKONIUM CHLORIDE**

sanitizing wipes cloth

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:82663-001
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.013 mg in 100 mg

**Inactive Ingredients**

Ingredient Name	Strength
<b>DECYL GLUCOSIDE</b> (UNII: Z17H97EA6Y)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>POTASSIUM SORBATE</b> (UNII: 1VPU26JZZ4)	

**WATER** (UNII: 059QF0KO0R)

**EDETATE DISODIUM ANHYDROUS** (UNII: 8NLQ36F6MM)

**CITRIC ACID MONOHYDRATE** (UNII: 2968PHW8QP)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82663-001-01	1200 in 1 CANISTER	04/12/2022	
1		1 mg in 1 CANISTER; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/12/2022	

**Labeler** - Purocor LLC (118531080)

Revised: 11/2022

Purocor LLC