PURCOL ANTIBACTERIAL WIPES LEMON SCENT- benzalkonium chloride cloth Purocol 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.

Purcol Anti-Bacterial Wipes Lemon Scent

Drug Facts

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antimicrobial

Uses

- For hand washing to decrease bacteria on skin.
- Recommended for repeated use

Warnings

For external use only.

Do not use:

- in the eyes.
- if you are allergic to any of the ingredients.

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

irritation or rash appears and lasts for more than 72 hours.

Keep out of reach of children.

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

Directions

- **Storage**. Store at room temperature.
- **Dispensing.** Flip open the dispending cap. Peel back the security label. Remove wipes. To reseal: slightly press on dispensing cap.

- Use. Apply wipe thoroughly to hands or surfaces as desired. Allow to dry without wiping.
- **Disposal**. Dispose of used wipes in trash bin after use. Do not flush.

Inactive ingredients

Ethylhexylglycerin, Glycerin, Phenoxyethanol, Polyaminopropyl biguanide, Propylene Glycol, Purified water.

Package Labeling:



PURCOL ANTIBACTERIAL WIPES LEMON SCENT

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75266-001
Route of Administration	TOPICAL		

	Active Ingredient/Active Moiety		
ı	Ingredient Name	Basis of Strength	Strength
		BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYAMINOPROPYL BIGUANIDE (UNII: DT9D8Z79ET)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

F	Packaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75266- 001-01	80 in 1 POUCH	03/05/2021	
1		2.5 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/05/2021	

Labeler - Purocol LLC (117479056)

Establishment			
Name	Address	ID/FEI	Business Operations
Purocol LLC		117479056	manufacture(75266-001) , label(75266-001) , pack(75266-001)

Revised: 3/2021 Purocol LLC