PROGRESS ANTIBACTERIAL WIPES LEMON SCENT- benzalkonium chloride cloth
Truechoicepack, 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.

Progress Anti-Bacterial Wipes Lemon Scent

Drug Facts

Active Ingredient

Benzalkonium chloride 0.13%

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.

When using this product

do not use in or contact the eyes.

Stop use and ask a doctor is

too much skin irritation or sensitivity develops or increases.

Keep this 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.

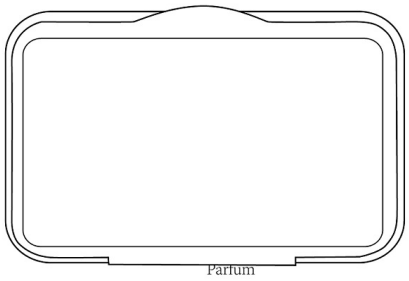
Other information

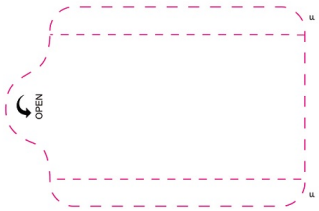
- Lot No. Manufacture date and Expiration date can be found on package.

Inactive Ingredients

Water, Phenoxyethanol, DMDM Hydantoin, Polysorbate 20, Ethylparaben, Methylparaben, Disodium Citrate, Aloe Barbadensis Leaf Juice, Parfum

Package Labeling:





PROGRESS ANTIBACTERIAL WIPES LEMON SCENT

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80882-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
ETHYLPARABEN (UNII: 14255EXE39)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
DISODIUM HYDROGEN CITRATE (UNII: 6FO62KCQ7A)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80882-002-50	50 in 1 BAG	11/20/2020	
1		4.04 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	11/20/2020	

Labeler - Truechoicepack, Corp (078786060)

Revised: 11/2020 Truechoicepack, Corp