BENZALKONIUM CHLORIDE- hand sanitizing wipes cloth
Progressive

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.

Hand Sanitizing Wipes

Active ingredients

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.

May irritate eyes.

Keep out of reach of children unless under adult supervision

Directions

- Remove lid
- Pull wipe from center of roll and thread through opening in lid

Do not push finger through opening

- Replace lid, pull wipe up, and then out 45º angle

The next wipe dispenses automatically

- Close lid to retain moisture

Inactive ingredient

Aloe Barbadensis Leaf Juice, Disodium EDTA, DMDM Hydantoin, Ethylparaben, Fragrance, Methylparaben, Phenoxyethanol, Polysorbate 20, Sodium Citrate, Water

Principal Display Panel

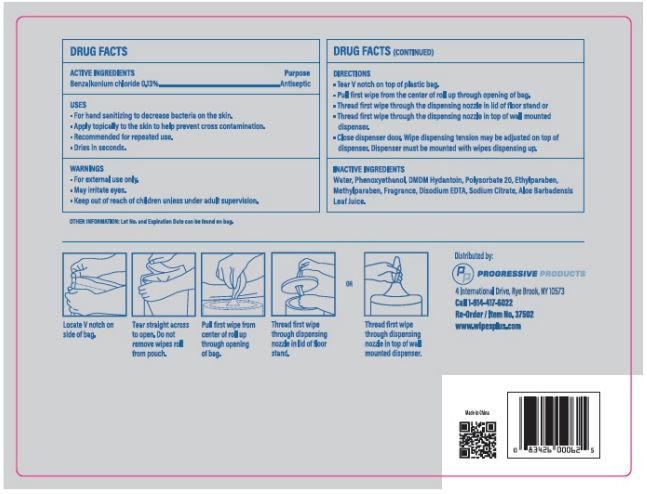
wipesplus

HAND SANITIZING

alcohol wipes

lemon scent

Kills 99.9% of germs



BENZALKONIUM CHLORIDE

hand sanitizing wipes cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67151-313
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.0013 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
ETHYLPARABEN (UNII: 14255EXE39)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67151-313-04	240 in 1 CANISTER	03/29/2022	
1		1 mg in 1 CANISTER; Type 0: Not a Combination Product		
2	NDC:67151-313-09	1500 in 1 CANISTER	03/29/2022	
2		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	03/29/2022	

Labeler - Progressive (127111792)

Revised: 10/2022 Progressive