WIPESPLUS SANITIZING ALCOHOL FREE WIPES- benzalkonium chloride cloth Progressive 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.

Drug Facts

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 the reach of children unless under adult supervision

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Directions

- Tear V notch on top of plastic bag.
- Pull first wipe from the center of roll up through opening of bag.
- Thread first wipe through the dispensing nozzle in lid of floor stand or
- Thread first wipe through dispensing nozzle in top of wll mounted dispenser.
- Close dispenser door. Wipe dispensing tension may be adjusted on top of dispenser. Dispenser must be mounted with wipes dispensing up.

Inactive Ingredient

Purified Water, Polysorbate 20, Glycerin, Phenoxyethanol, Sodium Benzoate, Citric Acid, Fragrance, Potassium Sorbate, Decyl Glucoside, Ethylhexylglycerin, Tocopheryl Acetate, Aloe Barbadensis Leaf Juice.

Principal Display Panel

Wipesplus

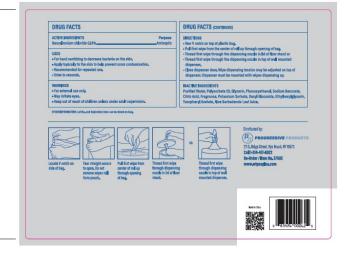
SANITIZING ALCOHOL FREE WIPES

Kills 99.9% of germs

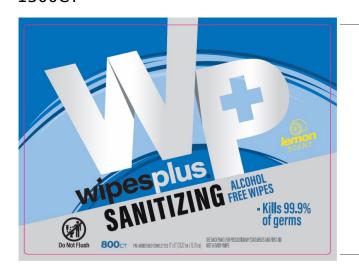


240CT



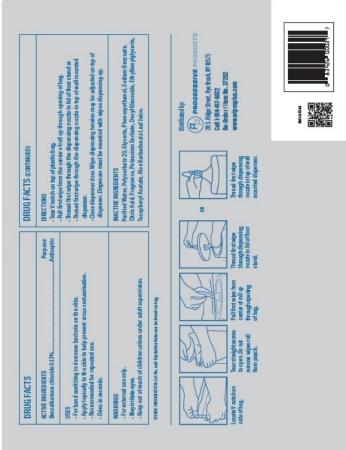


1500CT











400CT

WIPESPLUS SANITIZING ALCOHOL FREE WIPES benzalkonium chloride cloth Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:67151-343 Route of Administration TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZ ALKONIUM CHLORIDE	0.13 g in 100			

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)			
POLYSORBATE 20 (UNII: 7T1F30V5YH)			
GLYCERIN (UNII: PDC6A3C0OX)			
PHENOXYETHANOL (UNII: HIE492ZZ3T)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)			
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)			
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)			

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:67151-343- 01	240 in 1 CANISTER; Type 0: Not a Combination Product	07/13/2022		
2	NDC:67151-343- 02	1500 in 1 PACKAGE; Type 0: Not a Combination Product	07/13/2022		
3	NDC:67151-343- 03	800 in 1 PACKAGE; Type 0: Not a Combination Product	07/13/2022		
4	NDC:67151-343- 04	400 in 1 PACKAGE; Type 0: Not a Combination Product	07/13/2022		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	07/13/2022			

Labeler - Progressive Products, LLC (127111792)

Revised: 3/2023 Progressive Products, LLC