

**SANIPRO DISINFECTANT WIPES- benzalkonium chloride cloth**  
**AJ Co., Ltd.**

*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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Benzalkonium Chloride

Propylene Glycol

Phenoxyethanol

Glycerin

Sodium Benzoate

Sodium Citrate

Disodium EDTA

Polysorbate 20

Citric Acid

Melaleuca Alternifolia (Tea Tree) Leaf Oil

Tocopheryl Acetate

Antibacterial Wipes to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

keep out of reach of the children

Thoroughly wipe hands with wipe.

Discard properly. Not flushable.

Be sure to close lid to keep wipes moist.

For external use only. Flammable. Keep away from heat or flame.

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 thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

for external use only



## SANIPRO DISINFECTANT WIPES

benzalkonium chloride cloth

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:75356-0005
<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	0.1 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75356-0005-1	400 g in 1 PACKAGE; Type 0: Not a Combination Product	08/08/2020	
2	NDC:75356-0005-2	80 g in 1 PACKAGE; Type 0: Not a Combination Product	08/08/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/08/2020	

**Labeler** - AJ Co., Ltd. (631079605)

**Registrant** - AJ Co., Ltd. (631079605)

**Establishment**

Name	Address	ID/FEI	Business Operations
AJ Co., Ltd.		631079605	manufacture(75356-0005) , label(75356-0005) , pack(75356-0005)

Revised: 10/2020

AJ Co., Ltd.