SANIPRO DISINFECTANT WIPES- benzalkonium chloride liquid
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.

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.

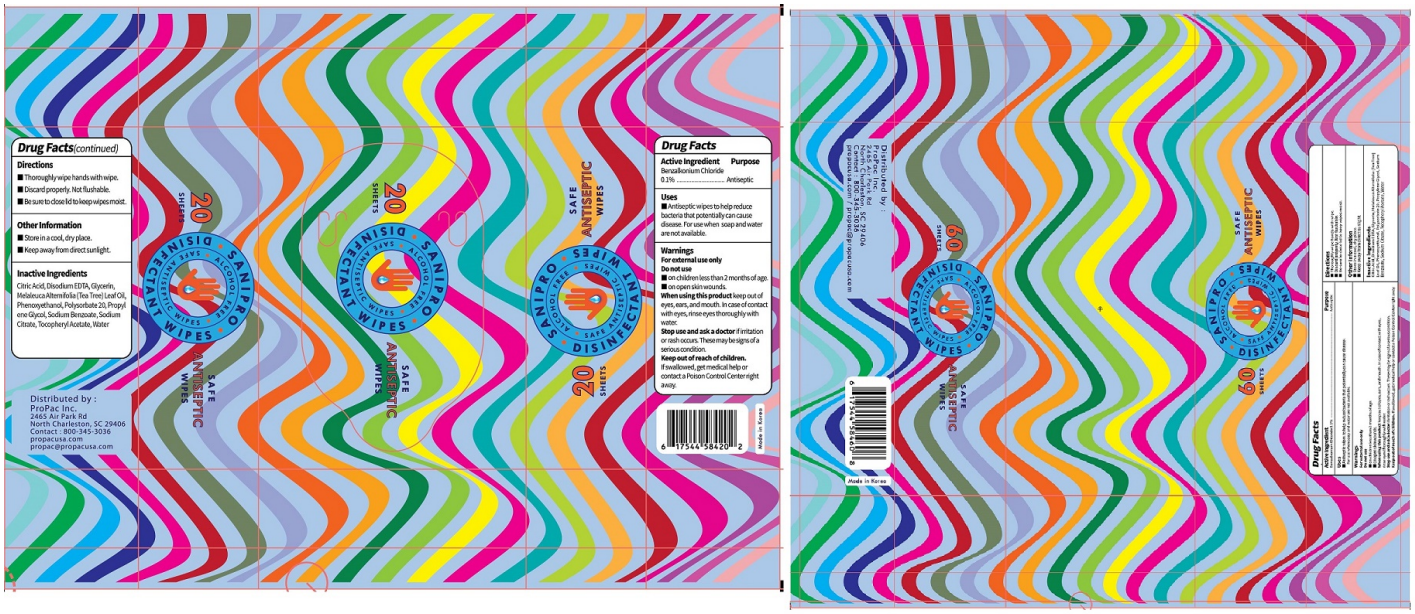
Do not use
- in children less than 2 months of age
- on open skin wounds

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 liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75356-0004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:75356-0004-1	400 g in 1 PACKAGE; Type 0: Not a Combination Product	08/08/2020	
2	NDC:75356-0004-2	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(75356-0004) , label(75356-0004) , pack(75356-0004)

Revised: 8/2020 AJ Co., Ltd.