

**BONBLISSITY HAND SANITIZING WIPES- benzalkonium chloride cloth  
BONBLISS BEAUTY, INC.**

*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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**73582-0002\_BONBLISSITY HAND SANITIZING WIPES**

**Active ingredient**

Benzalkonium Chloride 0.1%

**Purpose**

Antimicrobial

**Use**

- Hand sanitizer to help reduce bacteria

**Warnings**

**For external use only**

**When using this product** do not use in or near the eyes. If contact, rinse eyes thoroughly with water.

**Stop use and ask a doctor if** irritation develops and persists for more than 72 hours.

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

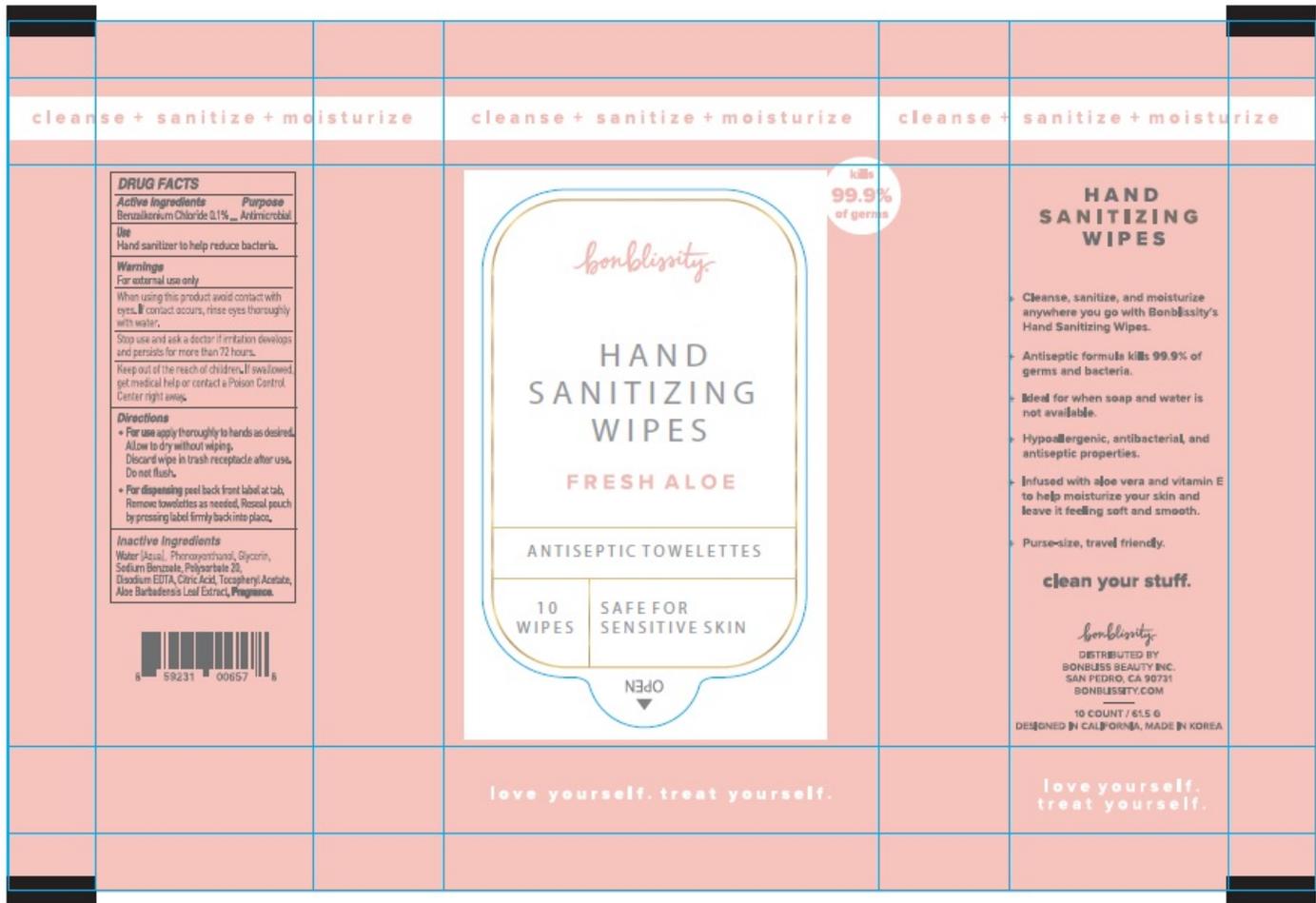
**Directions**

For use apply thoroughly to hands as desired. Allow to dry without wiping. Discard wipe in trash receptacle after use. Do not flush.

For dispensing peel back front label at tab. Remove towelettes as needed. Reseal pouch by pressing label firmly back into place.

**Inactive ingredients**

Water, Phenoxyethanol, Glycerin, Sodium Benzoate, Polysorbate 20, Disodium EDTA, Citric Acid, Tocopheryl Acetate, Aloe Barbadensis Leaf Extract, Fragrance



## BONBLISSITY HAND SANITIZING WIPES

benzalkonium chloride cloth

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:73582-0002
<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	0.01 g in 1 g

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)	
<b>EDETATE DISODIUM ANHYDROUS</b> (UNII: 8NLQ36F6MM)	

<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73582-0002-1	58 g in 1 POUCH; Type 0: Not a Combination Product	06/02/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/02/2020	

**Labeler** - BONBLISS BEAUTY, INC. (055735749)

**Registrant** - BONBLISS BEAUTY, INC. (055735749)

### Establishment

Name	Address	ID/FEI	Business Operations
FIRSTCHAM CO., LTD		694707327	manufacture(73582-0002)

Revised: 2/2023

BONBLISS BEAUTY, INC.