# BOOGIE HANDS ANTIBACTERIAL WET WIPES- benzalkonium chloride (0.115%) cloth ELEEO BRANDS 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.

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## **Boogie Hands Antibacterial Hand Wipes**

## **Active ingredients**

Benalkonium chloride (0.115%)

## **Purpose**

**Antibacterial** 

#### Use

For hand sanitizing to decrease bacteria on skin

## Warnings

For external use only

#### Do Not Use

if you are allergic to any of the ingredients

## When using this product

donot get into eyes.

if contact occurs, rinse thoroughly with water.

## Stop use and ask a doctor if

irritation or rash develops.

and continues 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:**

## \*adults and children 2 years and over

- apply to hands
- allow to dry without wiping

## \*children under 2 years

ask a doctor before use

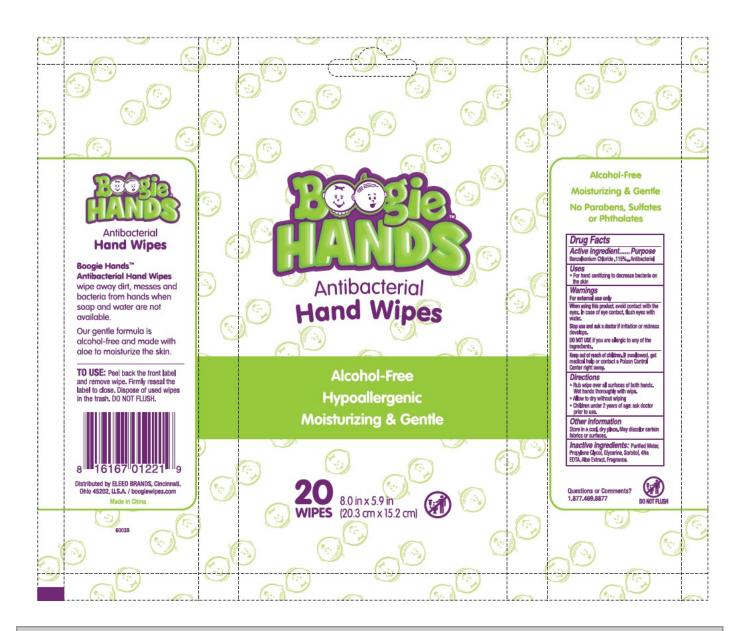
## **Inactive ingredients**

Water, Gycerin, Fragrance, Aloe extract, Disodium EDTA, Propylene Glycol, Sorbitol

## PRINCIPAL DISPLAY PANEL



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## **BOOGIE HANDS ANTIBACTERIAL WET WIPES**

benzalkonium chloride (0.115%) cloth

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79747-005	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZ ALKONIUM CHLORIDE	0.0012 g in 100 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
EDETATE SODIUM (UNII: MP1J8420LU)	

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79747-005- 20	3 in 1 CARTON	07/22/2020	
1		20 mL in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:79747-005- 02	20 mL in 1 PACKAGE; Type 0: Not a Combination Product	07/22/2020	
3	NDC:79747-005- 30	1 in 1 CARTON	07/22/2020	
3		30 mL in 1 POUCH; 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	07/22/2020	

## Labeler - ELEEO BRANDS LLC (078476782)

Revised: 12/2022 ELEEO BRANDS LLC