

**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**

do not 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, Glycerin, 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

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:79747-005
<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	0.0012 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

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

### 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