BOOGIE HANDS ANTIBACTERIAL WET WIPES- benzalkonium chloride (0.13%) cloth
ELEEO BRANDS LLC

Boogie Hands Antibacterial Hand Wipes

Active ingredients

Benalkonium chloride (0.13%)

Purpose

Antimicrobial

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

avoid contact with the eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if

if irritation or redness develops.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away

Directions:

- Rub wipe over all surfaces of both hands. Wet hands thoroughly with wipe.
- Allow to dry without wiping.
- Children under 2 years of age:ask doctor prior to use.

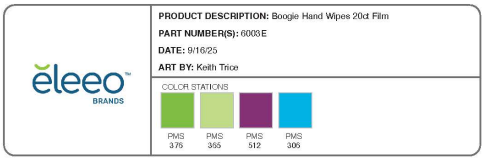
- Children under 6 years of age:use adult supervision.

Inactive ingredients

Purified water, Decyl glucoside, Aloe barbadensis(aloe) leaf extract, Tocopheryl (Vitamin E) acetate, Disodium EDTA, Potassium sorbate, Sodium benzoate, Phenoxyethanol, Citric acid, Fragrance

PRINCIPAL DISPLAY PANEL

20ct NDC:79747-014-02



BOOGIE HANDS ANTIBACTERIAL WET WIPES

benzalkonium chloride (0.13%) cloth

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79747-014
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.0013 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CITRIC ACID (UNII: 2968PHW8QP)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79747-014-20	3 in 1 CARTON	09/30/2025	
1		20 mL in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:79747-014-02	20 mL in 1 PACKET; Type 0: Not a Combination Product	09/30/2025	
3	NDC:79747-014-30	1 in 1 CARTON	09/30/2025	
3		30 mL in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:79747-014-40	2 in 1 CARTON	09/30/2025	
4		20 mL in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:79747-014-80	4 in 1 CARTON	09/30/2025	
5		20 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 Drug	505G(a)(3)	09/30/2025	

Labeler - ELEEO BRANDS LLC (078476782)

Revised: 9/2025 ELEEO BRANDS LLC