

TRAVEL LITE HAND SANITIZING WIPES- non-drying alcohol-free cloth
Diamond Wipes International, Inc

Travel Lite Hand Sanitizing Wipes

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Uses

■ for hand sanitizing to decrease bacteria on the skin when water and soap are not available

Warnings

For external use only

When using this product do not use in or near eyes. If contact occurs, flush thoroughly with water

Discontinue use if irritation or redness develops if condition persists for more than 72 hours contact a doctor

Keep out of reach of children

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

Directions

■ tear open packet, remove wipe ■ wet hands thoroughly with product

Inactive ingredients

Aloe Barbadosensis (Aloe vera) Leaf Juice, Citric Acid, Fragrance, Glycerin, Hamamelis Virginiana (Witch Hazel) Bark/Leaf/Twig Extract, Phenoxyethanol, Polysorbate 20, Potassium Sorbate, Sodium Benzoate, Tocopheryl Acetate, Water

Label

Wipe hands thoroughly and let air dry

Great alternative when soap and water are not available.

Vegan • Cruelty-Free • No added Gluten • No added Paraben • No added Gmo • No added Sulfate • No added Phthalate • No added Triclosan

*Escherichia coli, Staphylococcus aureus, Salmonella typhimurium, Pseudomonas aeruginosa, Burkholderia cepacia, Enterobacter cloacae, Enterococcus faecalis, Streptococcus pyogenes, Candida albicans, Aspergillus brasiliensis

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travel lite™

HAND SANITIZING WIPE

MAKE IN U.S.A. with global components

99.9% Effective against most common germs*

Non-drying alcohol-free formula
with Aloe and Vitamin E

8" x 7" (20.3 cm x 17.8 cm)

Do Not Flush

Drug Facts

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TRAVEL LITE HAND SANITIZING WIPES

non-drying alcohol-free cloth

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:64709-107

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
HAMAMELIS VIRGINIANA TOP (UNII: UDA30A2JJY)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

Product Characteristics

Color	white (White cloth)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64709-107-10	1 g in 1 PACKET; Type 0: Not a Combination Product	11/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/10/2020	

Labeler - Diamond Wipes International, Inc (161104729)

Registrant - Diamond Wipes International, Inc (161104729)

Establishment

Name	Address	ID/FEI	Business Operations
Diamond Wipes International, Inc		161104729	manufacture(64709-107)

Revised: 12/2023

Diamond Wipes International, Inc