

**ADVANCE ANTIBACTERIAL WIPES- benzalkonium chloride swab
NP Korea**

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.

Active Ingredient(s)

Benzalkonium Chloride 0.1%Antimicrobial

Purpose

Antimicrobial, Hand Wipes

Use

- To help reduce bacteria on the skin.
- For use when soap and water are not available.

Warnings

For external use only.

Do not use

- Do not use if you are allergic to any of the ingredients.
- Do not get into eyes. In case of contact, rinse eyes thoroughly with water.
- Avoid contact with broken skin.
- Do not inhale or ingest.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

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Directions

Adults and children 2 years and over;

- ☐ Apply wipe thoroughly to hands.
- ☐ Allow to dry without wiping.
- ☐ Children under 2 years, ask a doctor before use.

Other information

- Store at room temperature.
- Keep away from direct sunlight.

- Dispose of used wipes in trash receptacle after use.
- Do not flush.

Inactive ingredients

Water, Propylene Glycol, Phenoxyethanol, Glycerin, Sodium Benzoate, Sodium Citrate, Disodium EDTA, Polysorbate 20, Citric

Acid, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Tocopheryl Acetate.

Package Label - Principal Display Panel

AdvanceWipes.com



Drug Facts	<p>Warnings</p> <ul style="list-style-type: none"> For use when soap and water are readily available. Antibacterial Wipes to help reduce bacteria that potentially can cause disease. <p>Directions</p> <ul style="list-style-type: none"> Adults and children 2 years and over: Apply thoroughly to hands. Children under 2 years: Ask a doctor before use. Adhere to any labels on the off. <p>Other Information</p> <ul style="list-style-type: none"> Keep away from direct sunlight. Some are room temperature. Dispose of used wipes in trash receptacle after use. <p>Contains</p> <ul style="list-style-type: none"> Disinfectant Antibacterial Antiseptic Antifungal Antiviral Antiparasitic Anticorrosive Antistatic Antibacterial Antifungal Antiviral Antiparasitic Anticorrosive Antistatic <p>Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center right away. Some may cause drowsiness or dizziness. If you experience any of these symptoms, stop use immediately. Do not use if you are pregnant, nursing, or taking any medication. Do not use if you are allergic to any of the ingredients. Avoid contact with broken skin. Do not use on face or eyes.</p> <p>When using this product, do not get this wipe in case of contact, rinse eyes thoroughly with water.</p> <p>Do not use if you are allergic to any of the ingredients.</p> <p>For external use only.</p>
Active Ingredients <p>Benzalkonium Chloride 0.1%</p> <p>Antibacterial</p>	<p>Purpose</p> <p>Antibacterial</p>

Gentle on Skin



50 WIPES

Gentle on Skin

ALCOHOL FREE



Fresh Citrus Scent

ALCOHOL FREE



50 WIPES

Gentle on Skin

Kills 99% of most germs & illness causing bacteria*
(*Lab tested bacteria: Esherichiahia coli, Staphylococcus aureus, Salmonella typhimurium, Pseudomonas aeruginosa, Klensiella Pneumoniae)

FDA OTC Drug Registered

Distributed by: PackcoLLC Bellevue, WA

Manufactured by: NP Korea Corp. Made in Korea

AdvanceWipes.com

benzalkonium chloride swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79 178 -300
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.1 in 100

Inactive Ingredients

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

Product Characteristics

Color		Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79 178 -300 -50	50 in 1 PACKAGE; Type 0: Not a Combination Product	01/19/2021	

AdvanceWipes.com



8 80739 980155 5

Drug Facts

Active Ingredients: Benzalkonium Chloride 0.1%

Uses:
 For use when soap and water are readily available.
 Disinfectant Wipes to help reduce bacteria that potentially can cause disease.

Warnings:
 Do not use if you are allergic to any of the ingredients.
 The disinfectant may sting.
 Avoid contact with broken skin. Do not wash or repeat.
 Stop use and ask a doctor if irritation or rash appears 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.
 Contains Benzalkonium Chloride (BKC) and Citrus Fragrance.

Directions:
 Adults and children 2 years and over: Apply wipe thoroughly to hands.
 Rub for 20 seconds.
 Children under 2 years: Ask a doctor before use.

Other Information:
 Store at room temperature.
 Keep away from direct sunlight.
 Dispose of used wipes in trash receptacle after use.
 Do not flush.

Contains:
 No parabens.
 No dyes.
 No phthalates.
 No formaldehyde.
 No triclosan.
 No benzophenone.
 No sodium lauryl sulfate.
 No sodium hypochlorite.
 No sodium hydroxide.
 No sodium peroxide.
 No sodium silicate.
 No sodium stearate.
 No sodium tallowate.
 No sodium lauryl ether sulfate.
 No sodium lauryl ether sulfonate.
 No sodium lauryl ether phosphate.
 No sodium lauryl ether carboxylate.
 No sodium lauryl ether sulfate sodium salt.
 No sodium lauryl ether sulfonate sodium salt.
 No sodium lauryl ether phosphate sodium salt.
 No sodium lauryl ether carboxylate sodium salt.
 No sodium lauryl ether sulfate sodium salt sodium salt.
 No sodium lauryl ether sulfonate sodium salt sodium salt.
 No sodium lauryl ether phosphate sodium salt sodium salt.
 No sodium lauryl ether carboxylate sodium salt sodium salt.

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FDA OTC Drug Registered

Distributed by:
Packco LLC
Bellevue, WA

Manufactured by:
NP Korea Corp.
Made in Korea

AdvanceWipes.com

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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OTC monograph not final	part333A	01/19/2021	
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Labeler - NP Korea (689007752)

Registrant - NP Korea (689007752)

Establishment

Name	Address	ID/FEI	Business Operations
NP Korea		689007752	label(79178-300)

Establishment

Name	Address	ID/FEI	Business Operations
Daol G&C Co., Ltd.		693891755	manufacture(79178-300)

Revised: 1/2021

NP Korea