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

ADVANCE ANTIBACTERIAL WIPES

benzalkonium chloride swab

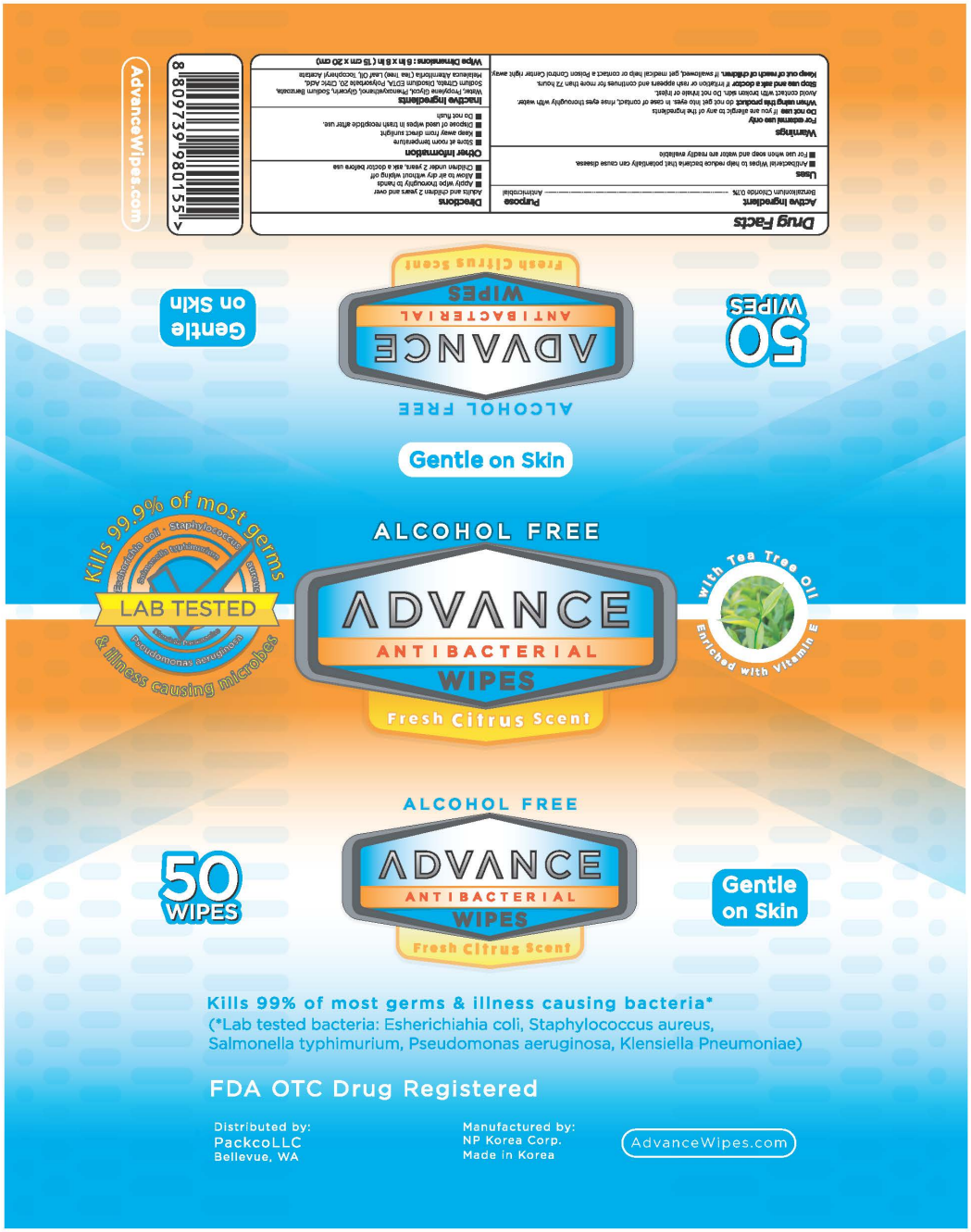
Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:79178-300
Route of Administration	TOPICAL		

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:79178-300-50	50 in 1 PACKAGE; Type 0: Not a Combination Product	01/19/2021	



Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

OTC monograph not final	part333A	01/19/2021	

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