PFA ANTIBACTERIAL WIPES- benzalkonium chloride cloth
Korea Living Goods Lab

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

Benzalkonium chloride 0.06%

INACTIVE INGREDIENTS

Sodium Levulinate, Levulic Acid, Laurylpyridium Chloride, Sodium Coco PG-dimonium Chloride Phosphate, Caprylhydroxamic Acid, Methylpropanediol, Water

PURPOSE

Antibacterial

WARNINGS

For external use only.

When using this product avoid contact with the eyes. In case of contact, flush eyes thoroughly with water

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Stop use and ask a doctor if irritation and redness develops and persists.

KEEP OUT OF REACH OF CHILDREN

If swallowed, get medical help or contact a Poison Control Center immediately.

Use(s)

■ For hand sanitizing to decrease bacteria on the skin.
■ Recommeded for repeated use.

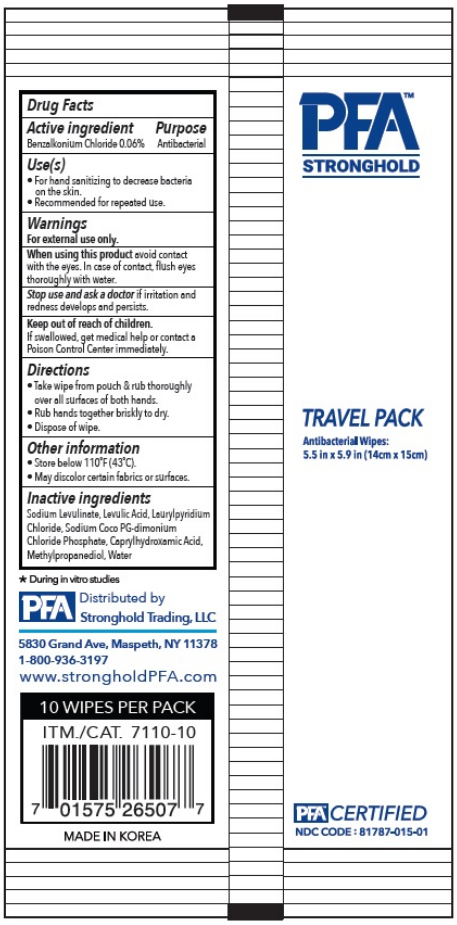
Directions

■ Take wipe from pouch & rub thoroughly over all surfaces of both hands.
■ Rub hands together briskly to dry.
■ Dispose of wipe

Other information

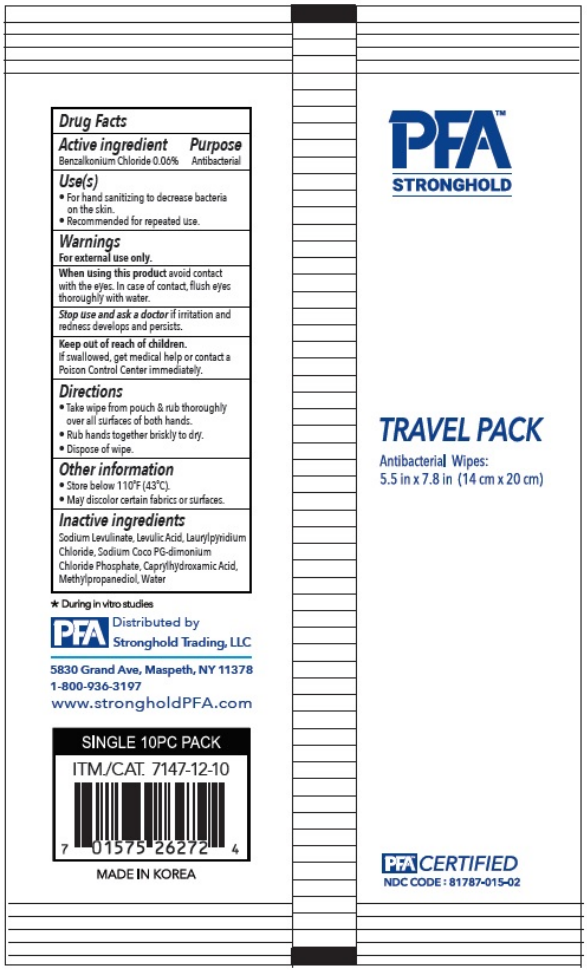
■ Store below 110℉(43℃)
■ May discolor certain fabrics or surfaces.

NDC: 81787-015-01
PFA ANTIBACTERIAL WIPES
5.5in x 5.9in (14cm x 15cm)



NDC: 81787-015-02
PFA ANTIBACTERIAL WIPES
5.5in x 7.8in (14cm x 20cm)

FRONT



BACK

PFA ANTIBACTERIAL WIPES

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81787-015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Benzalkonium chloride (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	Benzalkonium chloride	0.0006

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	
Sodium Levulinate (UNII: VK44E1MQU8)	
Methylpropanediol (UNII: N8F53B3R4R)	
Caprylhydroxamic Acid (UNII: UPY805K99W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81787-015-01	10 in 1 PACKET; Type 0: Not a Combination Product	10/01/2021	
2	NDC:81787-015-02	10 in 1 PACKET; Type 0: Not a Combination Product	10/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/01/2021	

Labeler - Korea Living Goods Lab (695538788)

Registrant - Korea Living Goods Lab (695538788)

Establishment

Name	Address	ID/FEI	Business Operations
Korea Living Goods Lab		695538788	manufacture(81787-015)

Revised: 11/2021 Korea Living Goods Lab