ANTIBACTERIAL MOISTURIZING HAND WIPES- benzalkonium chloride liquid
Yinjing Medical Technology (Shanghai) Co., Ltd.

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.

Antibacterial Moisturizing Hand Wipes

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Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

Decrease bacteria on the 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 eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor

if irritation or rash occurs 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.

Directions

Lift package label to open and remove wipe.
Wipe hands thoroughly. Allow to dry.
Discard wipe in trash. Do not flush.
Re-seal package after each use to avoid evaporation.
For children under 2 year, Consult a physician before use.

Inactive ingredients

Water, Glycerin, Propylene Glycol, Polysorbate-20, Phenoxyethanol, Tocopheryl Acetate, Aloe Barbadensis Leaf Juice, Diazolidinyl Urea, Decyl Glucoside, Disodium EDTA, Sodium Citrate,

Fragrance.

IN IN

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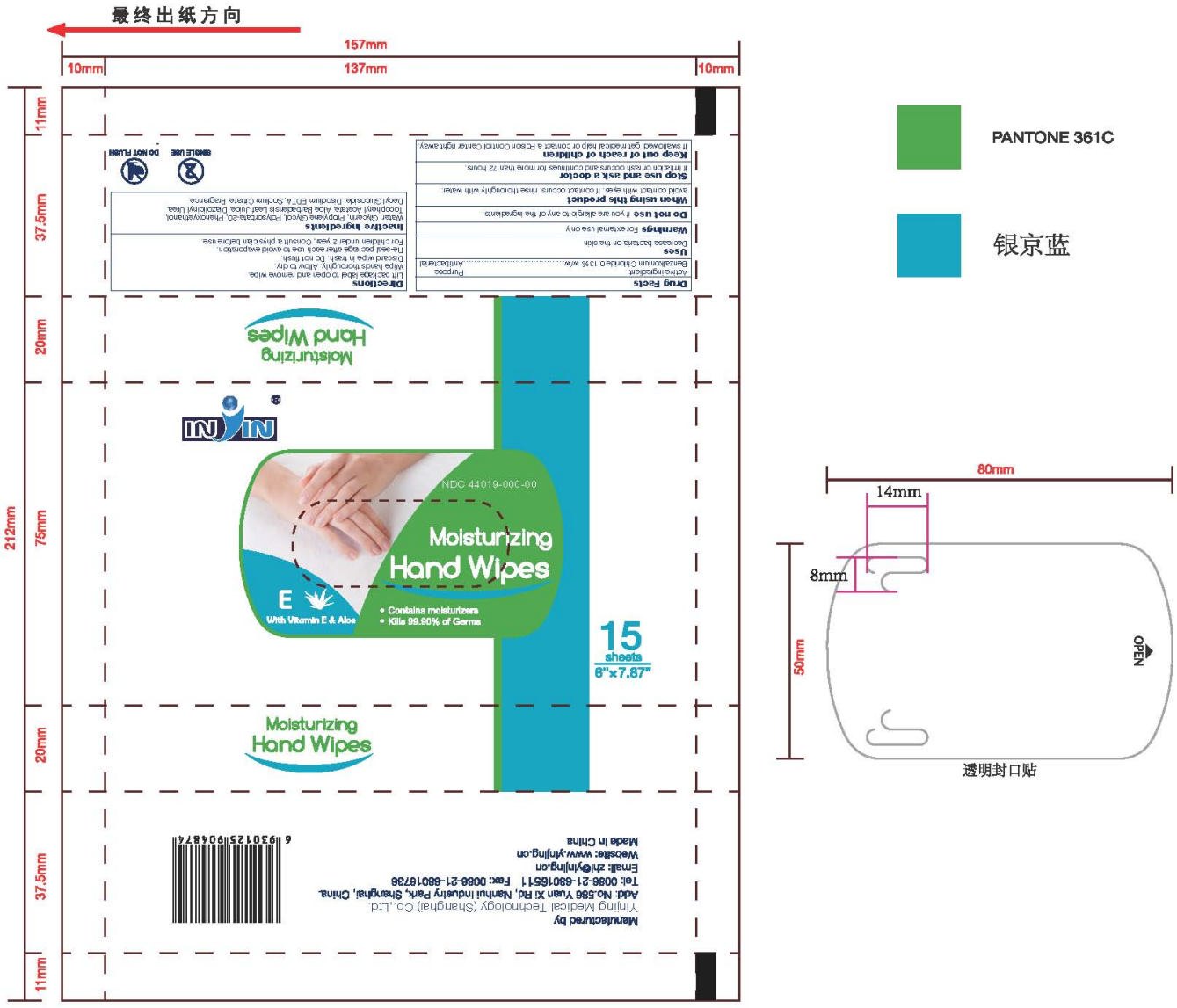
IN IN NDC: Antibacterial Moisturizing Hand Wipes 15 wipes 5.12 x 7.87 in (13cm x 20 cm) Kills 99.9% of Germs Contains Moisturizers, Vitamin E and Aloe

Antibacterial Moisturizing Hand Wipes 15 Wipes (44019-000-00)



ANTIBACTERIAL MOISTURIZING HAND WIPES
benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44019-000
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**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:44019-000-00	15 in 1 BAG	08/03/2016	
1		2.7 g in 1 PATCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	07/08/2013	

Labeler - Yinjing Medical Technology (Shanghai) Co., Ltd. (530501535)

Registrant - Yinjing Medical Technology (Shanghai) Co., Ltd. (530501535)

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