

INSTANT HAND SANITIZING WIPE- alcohol cloth
SHANDONG HENGFA HYGIENIC PRODUCTS CO., LTD.

Instant Hand Sanitizing Wipe

Instant Hand Sanitizing Wipe

Active Ingredient(s)

Alcohol 1mL/pcs. Purpose: Antiseptic

Purpose

Antiseptic

Use

To help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

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.

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

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Directions

- Wipe the surface of the skin

Other information

- Store between 15-30C (59-86F)

- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

WATER

HYDROGEN PEROXIDE

Package Label - Principal Display Panel

Gerigentle

Antimicrobial Alcohol Gel HAND WIPES

NDC 69771-001-01
Cleans and Moisturizes
An antiseptic handwash to help prevent cross-contamination

- Kills 99.99% of germs
- Contains moisturizing aloe, glycerin and vitamin E
- Removes soil and bacteria more effectively than gels alone

135 Wipes 6x7.5 in (15x19cm)
12/CS
Made in China

Reorder #23130

Manufactured for:
Geri-Gentle Corporation
Brooklyn, NY 11231

Antimicrobial Alcohol Gel Hand Wipes

Drug Facts		Directions
Active ingredient	Purpose	<ul style="list-style-type: none"> ■ To start: feed of the wipes. Remove cover and discard seal from container. From center of toweled roll, pull up wipe corner, tear off fist wipe for use. Twist next wipe into a point and thread through the hole in the container cover. Pull through about one inch. Replace cover. Pull out wipes as needed and snap off at 90° angle. Keep cap closed to prevent moisture loss. ■ Wipe hands, fingers, inter-digital areas and wrists thoroughly with towelette. Be sure to utilize the entire wipe surface. Allow to dry. ■ If hands are visibly soiled or contaminated, use first wipe clean hands, then discard wipe, sanitize with a single use. ■ Discard after single use.
Alcohol 65.9% by volume Antiseptic		
Uses		<p>Other Information Lot No. and Expiration Date can be found on canister</p> <p>Inactive ingredients: Aloe Barbadosis Leaf Juice, Carbomer, Glycerin, Propylene Glycol, Tetrasodium EDTA, Tocopheryl Acetate, Triethanolamine, Water</p>
<ul style="list-style-type: none"> ■ Antiseptic ■ For hand washing to decrease bacteria on the skin, after assisting ill persons, and before contact with a person under medical care or treatment. ■ Apply topically to the skin to help prevent cross-contamination ■ Recommended for repeated use ■ Dries in seconds 		
Warnings		
<ul style="list-style-type: none"> ■ Flammable, keep away fire or flame ■ For external use only 		
Do Not Use in or make contact with the eyes		
Stop Use if irritation and redness develop. If condition persists for more than 72 hours consult a physician.		
Keep Out Of Reach Of Children unless under adult supervision. If swallowed, get medical help or contact a Poison Center immediately		

INSTANT HAND SANITIZING WIPE

alcohol cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79913-201
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	1 mL

Inactive Ingredients

Ingredient Name	Strength
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79913-201-01	1 in 1 BAG; Type 0: Not a Combination Product	03/09/2021	
2	NDC:79913-201-02	12 in 1 BAG; Type 0: Not a Combination Product	03/09/2021	
3	NDC:79913-201-03	40 in 1 BAG; Type 0: Not a Combination Product	03/09/2021	
4	NDC:79913-201-04	80 in 1 BAG; Type 0: Not a Combination Product	03/09/2021	
5	NDC:79913-201-05	100 in 1 BAG; Type 0: Not a Combination Product	03/09/2021	
6	NDC:79913-201-06	135 in 1 PAIL; Type 0: Not a Combination Product	03/09/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/09/2021	

Labeler - SHANDONG HENGFA HYGIENIC PRODUCTS CO., LTD. (546195384)

Establishment

Name	Address	ID/FEI	Business Operations
SHANDONG HENGFA HYGIENIC PRODUCTS CO., LTD.		546195384	manufacture(79913-201)

Revised: 10/2023

SHANDONG HENGFA HYGIENIC PRODUCTS CO., LTD.