ANTISEPTIC WIPES- zylast hand wipes swab Westwood Laboratories

Zylast Antibacterial Hand Wipes

Ethyl Alcohol - 76%

Antiseptic

- To decrease bacteria on skin.
- Recommended for repeated use.
- For external use only
- Flammable. Keep away from flames or fires.
- Do not used in or near eyes.
- In case of contact, wash eyes thoroughly with water.
- Avoid contact with broken skin.
- Do not inhale or ingest.
- Irritiation or redness develops.
- Condition persists for more than 72 hours.

If swallowed, get medical help or call a Poison Control Center right away.

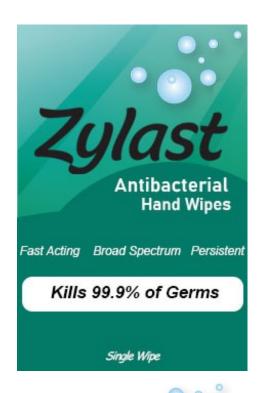
- Wet hands thoroughly with product and let dry without wiping.
- For children under 6, use only with adult supervision.
- Not recommended for use on infants.

Store 20-25*C (66-77*F)

Benzethonium Chloride, Dimethicone, Farnesol, Panthenol, PEG-12, Polyaminopropyl Biguanide, Water.

Single Wipe Labeling

NDC: 57362-469-18





Drug Facts

Active Ingredient

Purpose

Ethyl Alcohol - 76%.....Antiseptic

Uses

- To decrease bacteria on skin.
- Recommended for repeated use.

Warnings

For external use only

Flammable. Keep away from flames or fires.

When using this product Do not use in or near eyes

- In case of contact, wash eyes thoroughly with water.
- Avoid contact with broken skin. Do not inhale or ingest.

Stop use and ask a doctor if irritation or redness develops condition persists for more than 72 hours

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

Directions

 Wet hands thoroughly with product and let dry without wiping. =For children under 6, use only with adult supervision. =Not recommended for use on infants.

Other Information Store 20-25 C (66-77 F)

Inactive Ingredients Benzethonium Chloride, Dimethicone, Farnesol, Panthenol, PEG-12, Polyaminopropyl Biguanide, Water.

Manufactured for Innovative Biodefense, Inc. 9272 Jeronimo Rd. Suite #121, Irvine, CA 92618; Made in USA US Patent 10,524,997; Patents Pending

ANTISEPTIC WIPES

zylast hand wipes swab

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:57362-469
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	76 mL in 100 mL

Inactive Ingredients		
	Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57362-469- 19	60 mL in 1 PACKAGE; Type 0: Not a Combination Product	08/01/2020	
2	NDC:57362-469- 18	5 mL in 1 POUCH; Type 0: Not a Combination Product	08/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/01/2020	

Labeler - Westwood Laboratories (078363903)

Revised: 1/2023 Westwood Laboratories