CITYWIPES 75% ALCOHOL WIPES- alcohol patch NIKIA MEDIA, INC.

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.

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (75%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Ethanol Alcohol, 75% v/v, antiseptic

Purpose

Daily cleaning of items and Antimicrobial

Use

This product is used to wipe hands, faces, skins, etc. It can be used on hands, skin, electronic products, office supplies, tableware, children's toys, and a variety of surfaces. Open the label, take out the wet towel, wipe the target area, and discard it in a trash receptacle. Close package immediately after use to maintain purity.

Warnings

Do not apply this product to red and swollen skin, wound, or any part sensitive to it. Keep this product out of children's reach. If swallowed, get medical help or contact a Poison Control Center immediately. For external use only. Avoid contact with eyes & wounded skin. Discontinue use if irritation and redness develop. If the condition persists for more than 72 hours, consult a physician.

Store it in a cool and dry place. Flammable; avoid direct sunlight and open fire. And avoid contact with eyes and damaged skin.

Do not use

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

This product is used to wipe hands, skins, electronic products, office supplies, tableware, children's toys, and a variety of surfaces. 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.

Directions

Use it as needed, after following the usage instructions. For external use only.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Water

Package Label - Principal Display Panel

010 mL NDC: 79935-001-10



Effective Sterilization without residue.

Drug Facts (continued)



DNIZITINAS QNAH VECOHOL WIPES



CityWipes.









- DISINFECTION & STERILIZATION -PREVENT CROSS VISRUS TRANSMISSION



ALCOHOL WIPES

10 **WIPES** WIPE SIZE 6 * 8 in (15.2 * 20.3cm)

75% ALCOHOL

ALCOHOL WIPES







(F) (C) (C)





CITYWIPES 75% ALCOHOL WIPES

alcohol patch

Product Information	roduct Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79935-001(NDC:75163-100)	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 U in 100 U	

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Product Characteristics			
Color		Score	
Shape	FREEFORM	Size	
Flavor		Imprint Code	
Contains			

l	Packaging				
l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:79935-001-10	10 U in 1 PATCH; Type 0: Not a Combination Product	05/01/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/01/2020	

Labeler - NIKIA MEDIA, INC. (003692231)

Inactive Ingredients

Revised: 7/2020 NIKIA MEDIA, INC.