

FUBU ANTISEPTIC HAND SANITIZER WIPES- benzalkonium chloride cloth
TITUS KOZMETIK ARGE SANAYI VE TICARET LIMITED SIRKETI

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.

FUBU Antiseptic Hand Sanitizer Wipes

Drug Facts

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic

Use

For hand washing to decrease bacteria on the skin

Warnings

For external use only

Do not use

in the eyes

Stop use and ask a doctor if

- irritation and redness develop
- condition persists 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

- wet hands thoroughly with product and allow to dry without wiping

Other information

- Store at room temperature.
- Keep lid tightly closed when not in use.

Inactive ingredients

Benzoic Acid, C12-15 Pareth 12, Citric Acid, Dehydroacetic Acid, Fragrance, Glycerin, Phenoxyethanol, Water.

Package Labeling:15 WIPES





Distributed By :

Maha Global Partners Inc.
300 N. Midland Ave Saddle Brook NJ 07663

TO REPORT A SERIOUS ADVERSE EVENT, CONTACT
BEEKMAN INTERNATIONAL INC.
245 8TH AVE #101, NEW YORK, NY 10011
EMAIL: NY@BEEKMANUS.COM

MADE IN TURKEY

PRODUCTION DATE, EXPIRY DATE AND BATCH NUMBER ARE ON THE PACK.



ANTISEPTIC HAND SANITIZER WIPES
BENZALKONIUM CHLORIDE 0.13%



MULTIPURPOSE

15
wipes



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Package Labeling:72 WIPES



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benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81226-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
BENZOIC ACID (UNII: 8SKN0B0MIM)	
C12-15 PARETH-12 (UNII: 131316X18L)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
DEHYDROACETIC ACID (UNII: 2KAG279R6R)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81226-000-01	15 in 1 POUCH	01/01/2021	

1		1.8 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		
2	NDC:81226-000-02	72 in 1 POUCH	01/01/2021	
2		3.47 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

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

Labeler - TITUS KOZMETİK ARGE SANAYİ VE TİCARET LİMİTED SİRKETİ (595832524)

Revised: 12/2020

TITUS KOZMETİK ARGE SANAYİ VE TİCARET LİMİTED SİRKETİ