

ANTIBACTERIAL WET WIPES- benzethonium chloride cloth
YOYO LIP GLOSS, 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) (80%, 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. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. 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)

Alcohol 80% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer 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.

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away.

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

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

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel

LET'S BE FRIENDS!
#CollectLOL

WARNING!
TO AVOID DANGER OF SUFFOCATION, KEEP THIS PLASTIC BAG AWAY FROM BABIES AND CHILDREN. DO NOT USE THIS BAG IN CRIBS, BEDS, CARRIAGES OR PLAYPENS. THIS BAG IS NOT A TOY.

AVERTISSEMENT!
AFIN D'ÉVITER LE DANGER D'ASPHYXIE, VEUILLEZ CONSERVER CE SAC EN PLASTIQUE À DISTANCE DES BÉBÉS ET DES ENFANTS. N'UTILISEZ PAS CE SAC DANS LES BERCEAUX, LES LITS, LES LANDAUS OU PARCS POUR BÉBÉS. CE SAC N'EST PAS UN JOUET.

DIRECTIONS / CONSEILS:
PEEL BACK THE LABEL AND USE AS REQUIRED. RESEAL LABEL AFTER USE TO KEEP WIPES MOIST. ÉPILUCHER L'ÉTIQUETTE ET UTILISER SELON LES BESOINS. REFERMER L'ÉTIQUETTE APRÈS UTILISATION POUR GARDER LES LINGETTES HUMIDES.

FABRIQUÉ POUR
MANUFACTURED FOR
YOYO LIP GLOSS, INC
ASTORIA, NY 11103
1-800-YOYO-LIP
(1-800-969-6547)

YOYO WORLD
YOYO LipGloss.com

TRACKING: 44202.4422.0420 SKU: 44202

MADE IN CHINA / FABRIQUÉ EN CHINE
8 50018 07021 8

LOL SURPRISE!

ANTIBACTERIAL WET WIPES
LINGETTES HUMIDES ANTIBACTERIENNES
FRAGRANCE FREE | SANS PARFUM

Poids Net 16 Lingettes Net Wt. 16 Wipes

Drug Facts

Active Ingredients	Purpose
Benzethonium Chloride 0.1%	Antibacterial

Use decreases bacteria on skin

Warnings
For external use only

Do not use if you are allergic to any of the ingredients

When using this product, do not get into eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if irritation or rash develops 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
Adults and children 2 years & older:
• Apply to hands
• Allow to dry without wiping
Children under 2 years:
• Ask a doctor before use

Inactive ingredients:
Aqua, Propylene Glycol, Glycerin, Phenoxyethanol, Polyaminopropyl Biguanide Stearate, Hydroxypropyltrimonium Honey, Sodium Citrate

Faits sur les drogues

Objectif des ingrédients actifs	Purpose
Chlorure de benzalkonium 0,1%	Antibactérien

L'utilisation diminue les bactéries sur la peau

Avvertissements
Pour usage externe uniquement

Ne pas utiliser si vous êtes allergique à l'un des ingrédients

Lors de l'utilisation de ce produit, éviter tout contact avec les yeux. En cas de contact, rincer abondamment à l'eau.

Cessez l'utilisation et demandez à un médecin si une irritation ou une éruption cutanée se développe et persiste pendant plus de 72 heures.

Tenir hors de contact avec les enfants. En cas d'ingestion, consulter un médecin ou un centre antipoison immédiatement.

Instructions
Adultes et enfants de 2 ans et plus :
• Appliquer aux mains
• Laisser sécher sans essuyer
Les enfants de moins de 2 ans:
• Consulter un médecin avant utilisation

Ingrédients inactifs:
Aqua, Propylène Glycol, Glycérine, Phénoxyéthanol, Polyaminopropyl Biguanide Stéarate, Hydroxypropyltrimonium Honey, Sodium Citrate

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000 mL NDC: 00000-000-00

ANTIBACTERIAL WET WIPES

benzethonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70611-005
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
HONEY (UNII: Y9H1V576FH)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYAMINO PROPYL BIGUANIDE (UNII: DT9D8Z79ET)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70611-005-05	10 in 1 POUCH	03/30/2020	
1		1 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
2	NDC:70611-005-06	80 in 1 POUCH	03/30/2020	
2		1 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
3	NDC:70611-005-07	100 in 1 POUCH	03/30/2020	
3		1 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
4	NDC:70611-005-08	160 in 1 POUCH	03/30/2020	
4		1 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
5	NDC:70611-005-09	16 in 1 POUCH	03/30/2020	
5		1 mL in 1 APPLICATOR; Type 0: Not a Combination Product		
6	NDC:70611-005-10	72 in 1 POUCH	03/30/2020	
6		1 mL in 1 APPLICATOR; 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	part333A	03/30/2020	

Labeler - YOYO LIP GLOSS, INC (828881792)

Establishment			
Name	Address	ID/FEI	Business Operations
ZHEJIANG YOUQUAN CARE PRODUCTS TECHNOLOGY CO., LTD		421318425	manufacture(70611-005)

Revised: 5/2020

YOYO LIP GLOSS, INC