

**RAYON BAMBOO HEAVY DUTY ANTISEPTIC HAND SANITIZER WIPES-
benzalkonium chloride cloth
Orbizorb LLC**

Rayon Bamboo Heavy Duty Antiseptic Hand Sanitizer Wipes

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Use

For handwashing 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.
- if condition persists for more than 72 hours consult a doctor.

Keep out of reach of children.

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

Directions

- Peel back the foil seal and pull the center wipe through the slot in the lid.
- Wet hands thoroughly with product and allow to dry without wiping.
- Close lid after use to prevent wipes from drying out.

Inactive ingredients

Water, Estol, Poloxalene, PEG-75 Lanolin, Glycerin, Aloe Barbadensis Extract, Polyethylene glycol, Lauryl glycoide, Fragrance, Disodium EDTA, -Bromo-2-Nitropropane-1,3-Diol, 1,2-Benzisothiazolin-3-One

Package Labeling:

Manufactured for/ Elaborado por: Orbisorb LLC
 4715 N. 27th Street, Milwaukee, WI 53209
 TEL: 414-405-0223
 www.dirteezeus.com
 Made in China/Hecho en China
 General Purpose Cleaner – Max VOC 0.5%



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Limpiador de propósito general – Max VOC 0.5%
 NDC 90119-002-25

DIRTEEZE 



DIRTEEZE

25 Wipes/Toallitas 10.25" x 7.9" (26 cm x 20 cm)

DIRTEEZE

TESTED TO OEKO-TEX
 STANDARD 100 FOR
 HARMFUL SUBSTANCES

VOC FREE LOTION
 (NO VOLATILE
 ORGANIC COMPOUNDS)



BIODEGRADABLE WIFE AND LOTION

BIODEGRADABLE BAMBOO SOURCE

100% BIODEGRADABLE

ANTIBACTERIAL

DIRTEEZE

100% BIODEGRADABLE

Drug Facts	Purpose
Active ingredient Benzalkonium chloride 0.13%	Antiseptic
Use For handwashing 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. ■ If condition persists for more than 72 hours consult a doctor. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions ■ Peel back the foil seal and pull the center wipe through the slot in the lid. ■ Wet hands thoroughly with product and allow to dry without wiping. ■ Close lid after use to prevent wipes from drying out.	
Inactive ingredients Water, Ethanol, Polysorbate, PEG-75 Lanolin, Glycerin, Aloe Barbadensis Extract, Polyethylene glycol, Lauryl glycosides, Fragrance, Disodium EDTA, 2-Bromo-2-Nitropropane-1,3-Diol, 1,2-Benzothiazolin-3-One	

Información Sobre El Fármaco	Propósito
Ingrediente Activo Cloruro de Benzalconio al 0.13%	Antiséptico
Empleo Para lavarse las manos con el fin de disminuir las bacterias en la piel.	
Advertencias Solo para uso externo. No lo aplique en los ojos. Deje de usarlo y consulte a un médico si: ■ se presenta irritación y enrojecimiento. ■ la condición persiste por más de 72 horas consulte a su médico. Manténgase fuera del alcance de los niños. En caso de ingestión, busque ayuda médica o póngase en contacto con el Centro de Control de Envenenamientos inmediatamente.	
Modo de Empleo Despegue el sello de aluminio y jale la toallita central a través de la ranura de la tapa. Humedezca bien las manos con el producto y déjelo secar sin enjuagarlo. Cierre la tapa después de usarla para evitar que las toallitas se sequen.	
Ingredientes Inactivos Agua, Etanol, Polisorbato, Lanolina PEG-75, Glicerina, Extracto de Aloe barbadensis, Polietilenglicol, Lauril glucosido, Fragancia, EDTA disódico, 2-bromo-2-nitropropano-1,3-diol, 1,2-benzotiazolin-3-ona	

RAYON BAMBOO HEAVY DUTY ANTISEPTIC HAND SANITIZER WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:90119-002
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
BENZISOTHIAZOLINONE (UNII: HRA0F1A4R3)	
WATER (UNII: 059QF0KO0R)	
POLOXALENE (UNII: V8B3K56SW0)	
PEG-75 LANOLIN (UNII: 09179OX7TB)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
BRONOPOL (UNII: 6PU1E16C9W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:90119-002-25	25 in 1 PACKAGE	01/05/2021	
1		7.5 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 Drug	505G(a)(3)	01/05/2021	

Labeler - Orbizorb LLC (117447551)