CORDIAL CLEAN AND PURE- benzalkonium chloride cloth
Lambi, S.A. de C.V.

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.

Hand Sanitizer Wipes

BENZALKONIUM CHLORIDE WIPES

BENZALKONIUM CHLORIDE 0.13% Purpose: Antimicrobial

Purpose

Antimicrobial

Use

To decrease bacteria on the skin

Warnings

For external use only.

When using this product

- Do not use in or near the eyes
- In case of contact rinse eyes thoroughly with water
- Avoid contact with open skin wounds
- Do not use on children under 2 months of age

Stop use and ask a doctor

If irritation or rash occurs.

Keep out of reach of children

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

Directions

- Adults and children 6 years and older - Rub wipe thoroughly over all surfaces of hands and allow to dry.
- Children under 6 years - Use only under adult supervision

Other information

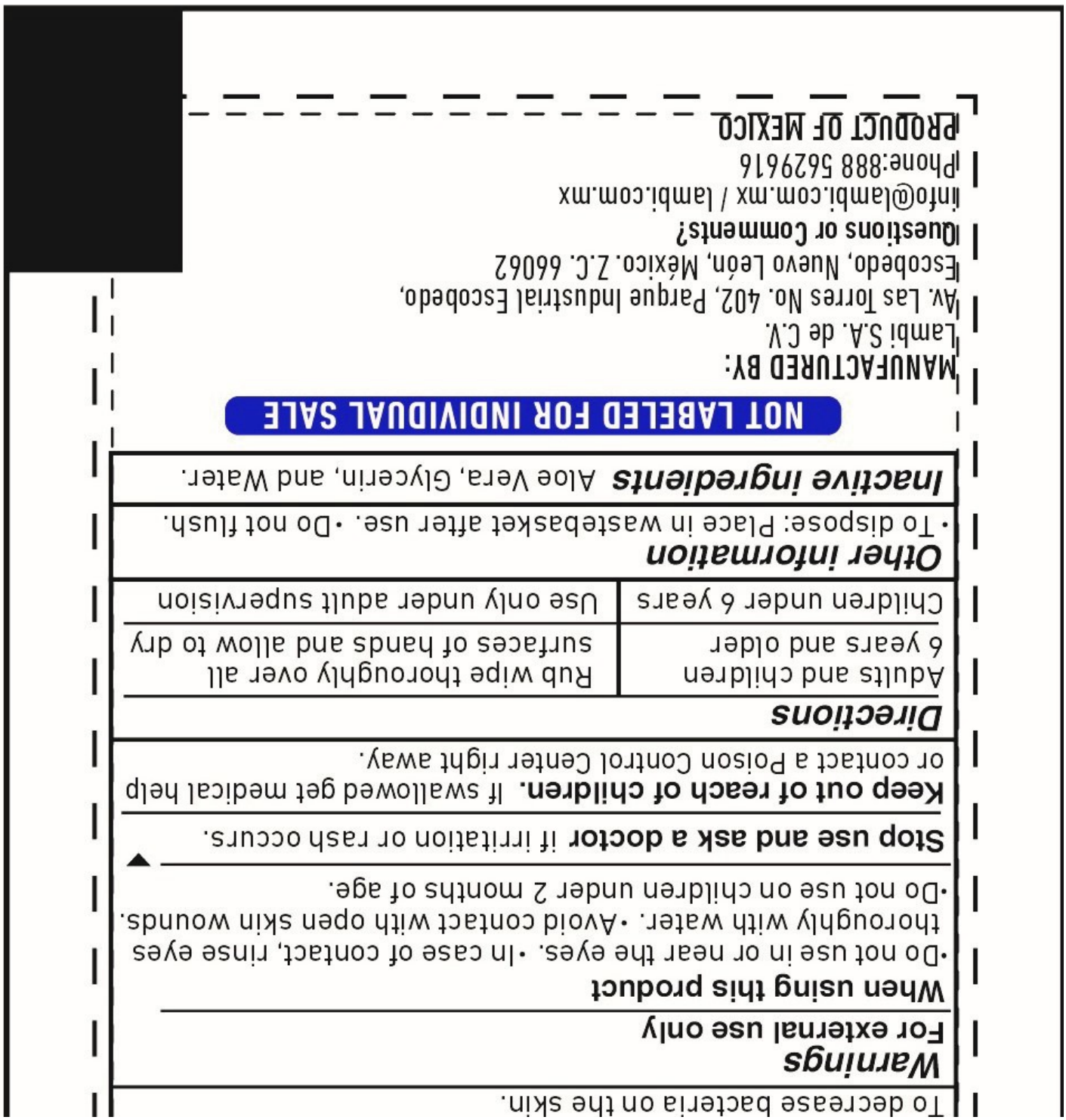
- To dispose - Place in waste basket after use
- Do not flush

Inactive ingredients

Aloe Vera, Glycerin and Water

Package Label - Sachet wipe

Sachet wipe NDC 81510-001-02



Drug Facts

Active Ingredient	Purpose
Benzalkonium chloride 0.13%	Antimicrobial

Use

CORDIAL

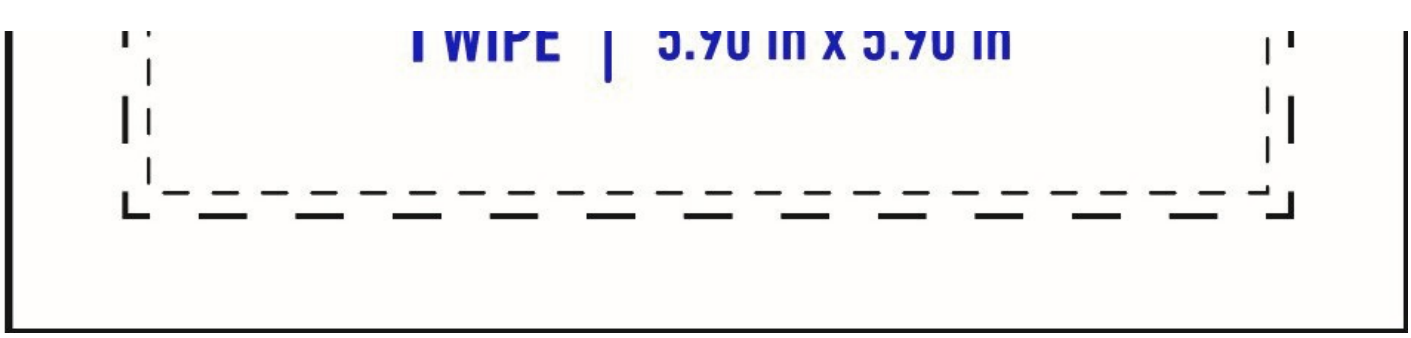
CLEAN & PURE™

HAND SANITIZING WIPE

MOISTURIZING FORMULA
WITH ALOE

NOT LABELED FOR INDIVIDUAL SALE

1 WIPE | 5.00 in x 5.00 in



LAMBI_BOX_100

CORDIAL CLEAN AND PURE

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81510-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81510-001-02	100 in 1 BOX	05/28/2021	
1		1 g in 1 PACKET; 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	05/28/2021	

Labeler - Lambi, S.A. de C.V. (811898915)

Registrant - Lambi, S.A. de C.V. (811898915)

Establishment

Name	Address	ID/FEI	Business Operations
Lambi, S.A. de C.V.		811898915	manufacture(81510-001)

Revised: 5/2021 Lambi, S.A. de C.V.