TRICLARA- benzalkonium chloride liquid
Nightingale Pharmaceuticals, Inc.

TriClara Antiseptic Cold Sore Treatment

Drug Facts

Active Ingredient

Benzalkonium Chloride (0.13%)

Purpose

Antiseptic - First Aid

Uses

- treatment of cold sore / fever blister
- effective relief of symptoms of cold sore / fever blister
- protects against infections in sores, burns, cuts and scrapes

Warnings

- **For external use only:** Do not use in or around eyes or ears. Consult a physician before using on puncture wounds, serious burns or animal bites.
- **Allergy alert:** Do not use if you are allergic to any ingredients in this product.

Stop use and consult a doctor if the condition persists or gets worse.

Keep out of reach of children. If swallowed, get medical help or contact poison control at 1-800-222-1222.

Directions

- clean the affected area with soap and water and thoroughly dry the area.
- for best results apply a small quantity with a rubbing motion 8 times (once every 2 hours while awake) daily at onset of symptoms until cleared.
- wash hands after applying.
- do not share this product with others.
- retain these directions for help with use.
- for use on children under 12 years: ask a doctor.

Other information

- Store at 15 to 25°C (59 to 77°F), in a dry place, away from heat, humidity and direct sunlight
- **Discard 30 days after removeal from vauumed pouch.**

Inactive ingredients

glycerin, water, N-acetyl cysteine, sodium hydroxide (for pH), ascorbic acid

TriClara ® ON & GONE

New

Antiseptic Cold Sore Treatment

0.13% Benzalkonium Chloride

FAST

CLEAR

EFFECTIVE

BETTER APPEARANCE + SOOTHING RELIEF

Cold Sore / Fever Blister Antiseptic

NET WT 3.5G / .12 OZ

Distributed by:

Nightingale Pharmaceuticals, Inc.

Santee, CA 92071

www.triclara.com

RETAIN CARTON FOR COMPLETE PRODUCT INFORMAITON

NEW

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ON & GONE

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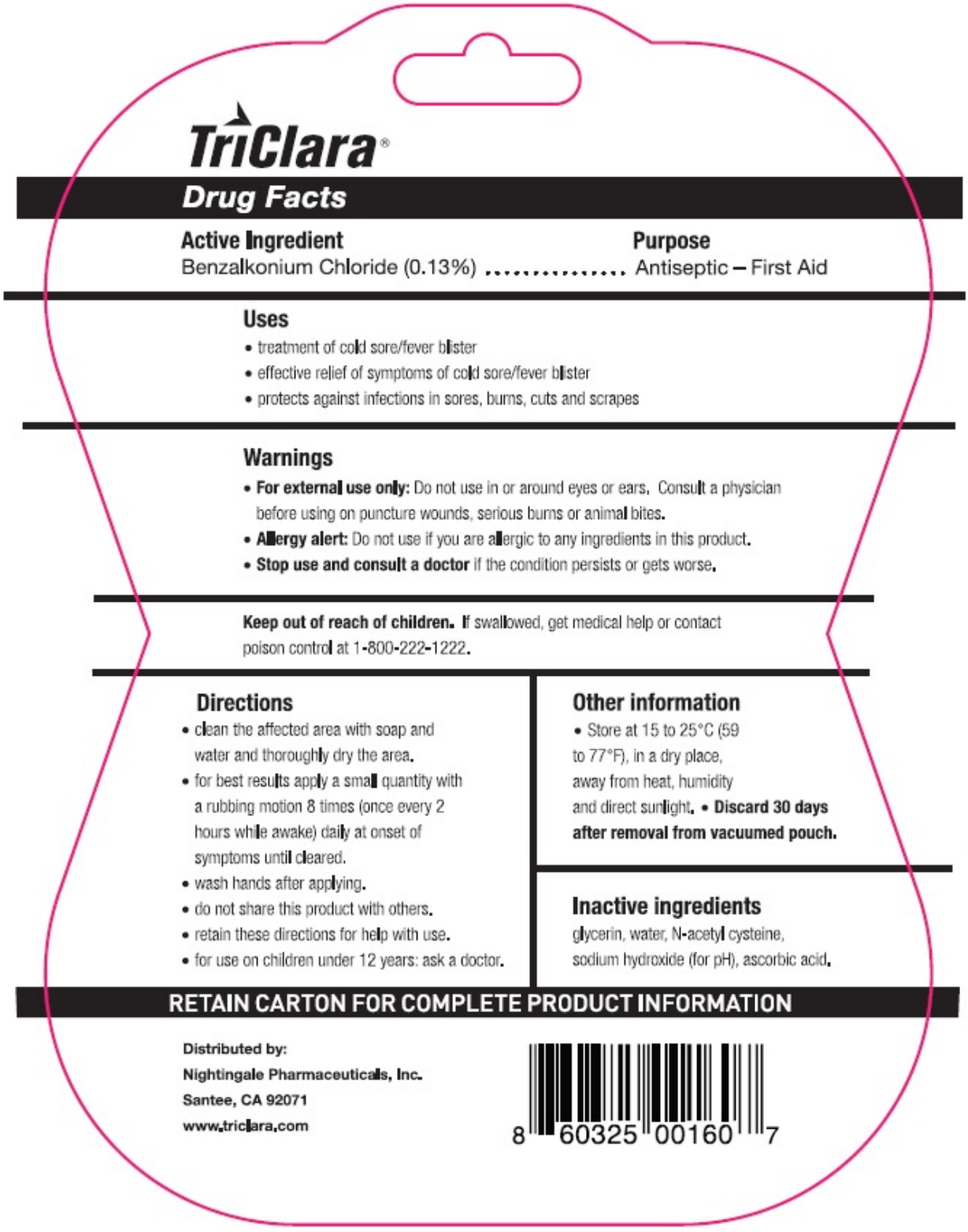

TRICLARA

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72518-001

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
WATER (UNII: 059QF0KO0R)	
ACETYLCYSTEINE (UNII: WYQ7N0BPYC)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ASCORBIC ACID (UNII: PQ6CK8PD0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72518-001-00	3.5 g in 1 BOTTLE; Type 0: Not a Combination Product	05/14/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/14/2019	

Labeler - Nightingale Pharmaceuticals, Inc. (081338827)

Revised: 12/2023 Nightingale Pharmaceuticals, Inc.