

MYTERRA HAND SANITIZER FOAM NA- benzalkonium chloride soap
PortionPac Chemical Corporation

MyTerra Hand Sanitizer Foam NA

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antimicrobial

Uses

- For hand sanitizing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

For external use only.

When using this product avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, or if 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

- Pump a small amount of foam into palm of hand.
- Rub thoroughly over all surfaces of both hands.
- Rub hands together briskly until dry.

Inactive Ingredients

Water, Glycerin, Propylene Glycol, Phenoxyethanol, Lauramine Oxide, Panthenol, Fragrance, Citric Acid

my.terra
Hand Sanitizer
Foam NA
Alcohol Free

Distributed by:
PortionPac Chemical Corp.
 85 Bradrock Drive
 Des Plaines, IL 60018
 (312) 226-0400

Drug Facts

Active Ingredient

Benzalkonium Chloride

Uses

- For hand sanitizing
- Recommended for use on

Warnings

For external use only

When using this product, avoid eye contact, flush eyes with water if contact occurs.

Stop use and ask a doctor if condition persists for more than 7 days.

Keep out of reach of children. If swallowed, contact a Poison Control Center or call a doctor immediately.

Directions

- Pump a small amount of foam onto your hands.
- Rub thoroughly over all surfaces of your hands.
- Rub hands together for at least 20 seconds.

Inactive Ingredients

Water, Glycerin, Propylamine Oxide, Propylamine Oxide, Propylamine Oxide, Propylamine Oxide

LOT #:
 EXP:

Contains: Eight Bags

Net Contents Each: 33.8 fl. oz. (1 qt. 1.8 fl. oz.) 1000 mL

Total Net Volume: 270.4 fl. oz. (2 gal. 14.4 fl. oz.) 8 L



Please Recycle This Container

MYTERRA HAND SANITIZER FOAM NA

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70780-151
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PANTHENOL (UNII: WW9CM0O67Z)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)

Product Characteristics

Color	white (water white - colorless, dispensed as a white foam)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70780-151-13	8 in 1 BOX	10/30/2017	
1		1000 mL in 1 BAG; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/30/2017	

Labeler - PortionPac Chemical Corporation (005448584)

Revised: 9/2025

PortionPac Chemical Corporation