PERIGIENE- antimicrobial drug product soap
Dermarite Industries LLC

DRUG LISTING: PERIGIENE

Active Ingredient

Chloroxylenol 0.5%

Purpose:

Antiseptic

Uses:

- Antiseptic cleanser for use in the perineal area.

Warnings:

- **For external use only.**
- **Avoid contact with eyes.** In case of contact, flush thoroughly with water.
- **Ask a doctor before use if** you have , deep of puncture wounds, animal bites, serious burns
- **When using this product,** do not apply to areas of raw or blistered skin in large quantities, do not use in or near the eyes
- **Stop use and ask a doctor if**, condition worsens, symptoms last for more than 7 days or clear up and recur within a few days

Warnings:

Keep out of reach of children. In case of accidental ingestion contact a physician or Poison Control Center right away

Directions:

- Apply to perineal area.
- Wipe with soft cloth.
- Repeat as needed to clean the area.

Other information:

Store at room temperature (59°-86°F). You may report a serious adverse event to DermaRite Industries, PO Box 7209, North Bergen, NJ 07047

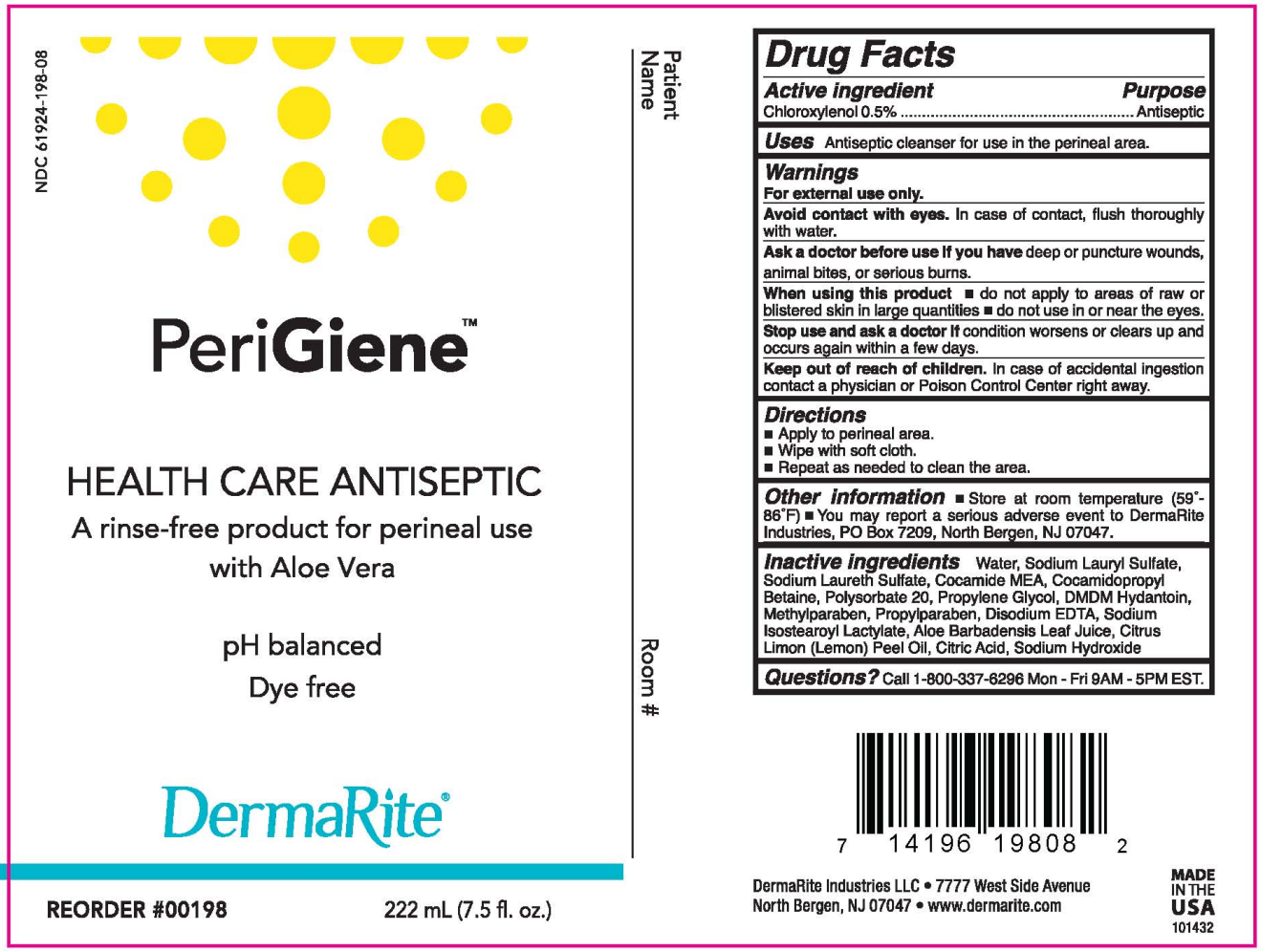
Inactive Ingredients:

Water, Sodium Lauryl sulfate, Sodium Laureth Sulfate, cocamide MEA, Cocamidopropyl Betaine, Polysorbate 20, Propylene glycol, DMDM Hydantoin, Methylparaben, Propylparaben, Disodium EDTA, Sodium Isostearoyl Lactylate, Aloe Barbadensis Leaf Juice, Citrus Limon (Lemon) Peel Oil, Citric Acid, Sodium Hydroxide.

Question?

Call 1-800-337-6296 Mon-Fri 9AM-5PM EST.

Perigiene Package Label Principal Display Panel



PERIGIENE

antimicrobial drug product soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:61924-198
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.005 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LEMON OIL (UNII: I9GRO824LL)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SODIUM ISOSTEAROYL LACTYLATE (UNII: 8730J0D3EV)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61924-198-08	222 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	12/12/2011	
2	NDC:61924-198-01	3800 mL in 1 JUG; Type 0: Not a Combination Product	12/12/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	12/12/2011	

Labeler - Dermarite Industries LLC (883925562)

Registrant - Dermarite Industries LLC (883925562)

Establishment

Name	Address	ID/FEI	Business Operations
Dermarite Industries LLC		883925562	manufacture(61924-198)

Revised: 12/2023 Dermarite Industries LLC