

**PERIGIENE- antimicrobial drug product soap**  
**DermaRite Industries LLC**

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**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:**

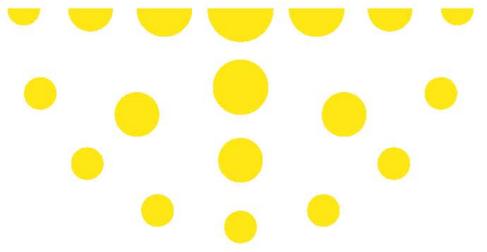
Water, Sodium Lauryl sulfate, Sodium Laureth Sulfate, cocamide MEA, Cocamidopropyl Betaine, Polysorbate 20, Propylene glycol, DMDM Hydantoin, Methylparaben, Propylparaben, Disodium EDTA, Sodium Isostearoyl Lactylate, Aloe Barbadensis Leaf Juice, Citrus Limon (Lemon) Peel Oil, Citric Acid, Sodium Hydroxide.

**Question?**

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

**Perigiene Package Label Principal Display Panel**

NDC 61924-198-08



# PeriGiene™

**HEALTH CARE ANTISEPTIC**  
A rinse-free product for perineal use  
with Aloe Vera

pH balanced  
Dye free



REORDER #00198      222 mL (7.5 fl. oz.)

Patient Name

**Drug Facts**

Active ingredient	Purpose
Chloroxylenol 0.5%	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 or puncture wounds, animal bites, or 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 or clears up and occurs again within a few days.  
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

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

Room #



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DermaRite Industries LLC • 7777 West Side Avenue  
North Bergen, NJ 07047 • www.dermarite.com

**MADE  
IN THE  
USA**  
101432

**PERIGIENE**

antimicrobial drug product soap

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:61924-198
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

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

**Inactive Ingredients**

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

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**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	

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**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: 12/2024

Dermarite Industries LLC