HEMORRHOIDAL PAIN RELIEF- hemorrhoidal pain relief ointment ointment
Neilmed Pharmaceuticals Inc.

Hemorrhoidal Pain Relief Ointment

Drug Facts

Active Ingredients:

Mineral Oil 14%

Petrolatum 74.9%

Phenylephrine HCl 0.25%

Drug Facts:

Purpose

Protectant

Protectant

Vasoconstrictor

Uses:

- Helps relieve the local itching and discomfort associated with hemorrhoids
- Temporarily shrinks hemorrhoidal tissue and relieves burning
- Temporarily provides a coating for relief of anorectal discomforts
- Temporarily protects the inflamed, irritated anorectal surface to help make bowel movements less painful

Warnings:

For external and/or intrarectal use only

Ask a doctor before use if you have:

- Heart disease
- High blood pressure
- Thyroid disease
- Diabetes
- Difficulty in urination due to enlargement of prostate gland

Ask a doctor or pharmacist before use if you are:

Presently taking a prescription drug for high blood pressure or depression

When using this product:

- Do not exceed the recommended daily dosage unless directed by a doctor

Stop use and ask doctor if:

- Bleeding occurs
- Condition worsens or does not improve within 7 days
- Introduction of applicator into the rectum causes additional pain

If pregnant or breast-feeding:

Ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions:

- Adults: When practical, cleanse the affected area by patting or blotting with an appropriate cleansing wipe. Gently dry by patting or blotting with a tissue or a soft cloth before applying ointment.
- When first opening the tube, puncture foil seal with top end of cap.
- Apply to the affected area up to 4 times daily, especially at night, in the morning or after each bowel movement.
- Intrarectal use: Remove cover from the applicator, attach applicator to tube, lubricate applicator well and gently insert applicator into the rectum, thoroughly cleanse applicator after each use and replace cover.
- Also apply ointment to external area.
- Regular use provides continual therapy for relief of symptoms.

Other Information:

Store at 15°C – 30°C (59°F – 86°F)

- Children under 12 years of age: ask a doctor

Inactive ingredients:

Benzoic acid, butylated hydroxytoluene, corn oil, glycerin, lanolin, lanolin alcohol, methylparaben, propylparaben, paraffin wax, purified water, thymus vulgaris oil, tocopherol, acetate, white wax

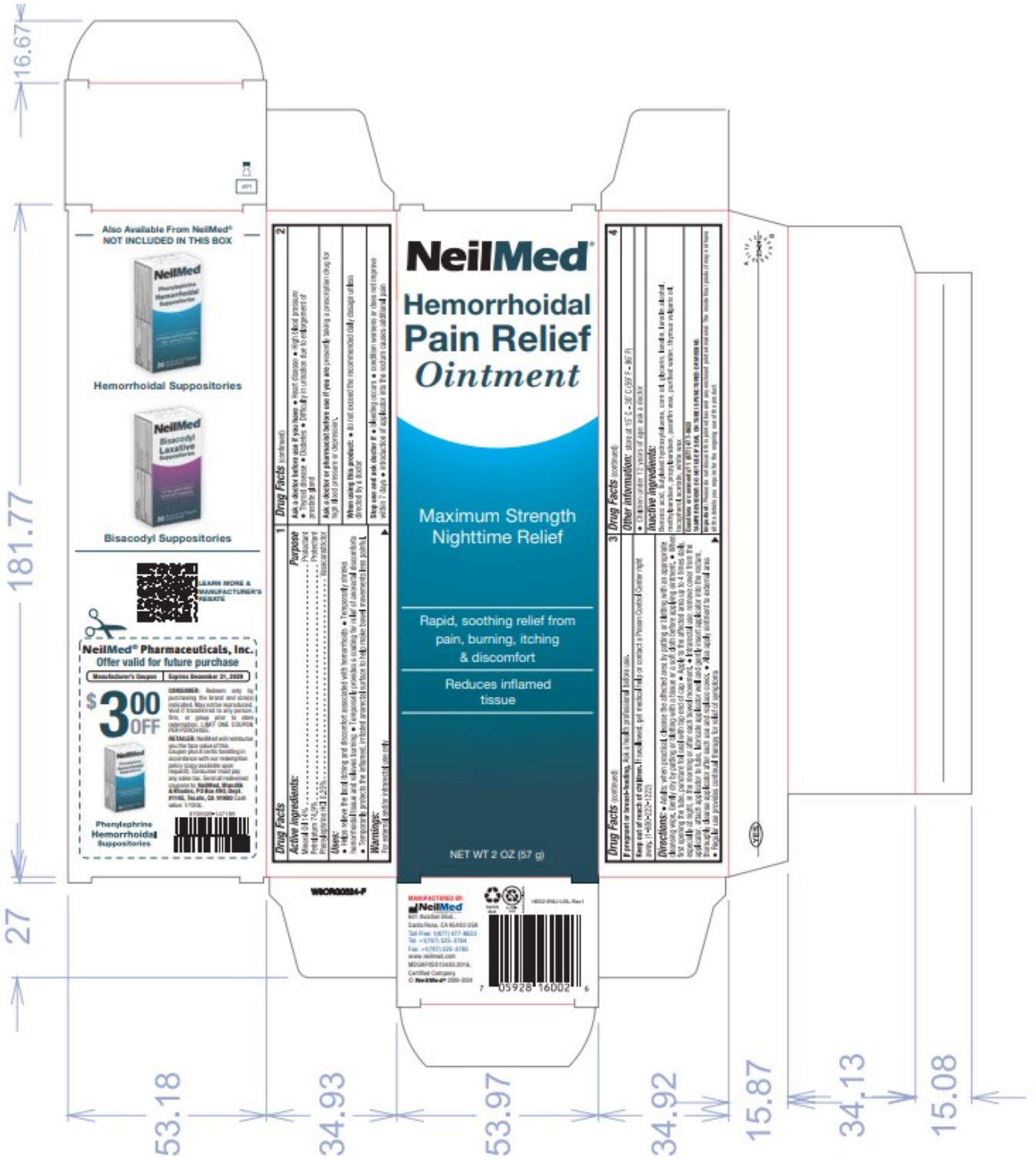
Questions or comments?

1 (877) 477-8633

TAMPER EVIDENT: DO NOT USE IF SEAL ON TUBE IS PUNCTURED OR MISSING

Important:

Please do not discard this printed box and any enclosed printed material. The inside final product may not have all the details you require for the ongoing use of the product.



HEMORRHOIDAL PAIN RELIEF

hemorrhoidal pain relief ointment ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:13709-334
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	749 mg in 1 g
MINERAL OIL (UNII: T5L8T28FGP) (MINERAL OIL - UNII:T5L8T28FGP)	MINERAL OIL	140 mg in 1 g
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	2.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
METHYLPARABEN (UNII: A2I8C7HI9T)	
PARAFFIN (UNII: I9O0E3H2ZE)	
WATER (UNII: 059QF0KO0R)	
THYMUS VULGARIS (THYME) OIL (UNII: 2UK410MY6B)	
LANOLIN (UNII: 7EV65EAW6H)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
GLYCERIN (UNII: PDC6A3C0OX)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
LANOLIN ALCOHOL (UNII: 884C3FA9HE)	
WHITE WAX (UNII: 7G1J5DA97F)	
CORN OIL (UNII: 8470G57WFM)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13709-334-01	1 in 1 CARTON	06/11/2025	
1		57 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	06/11/2025	

Labeler - Neilmed Pharmaceuticals Inc. (799295915)

Establishment

Name	Address	ID/FEI	Business Operations
Neilmed Pharmaceuticals Inc.		799295915	manufacture(13709-334)

Revised: 6/2025

Neilmed Pharmaceuticals Inc.