

**DENDRACIN NEURODENDRAXCIN- methyl salicylate, menthol and capsaicin lotion**  
**Preferred Pharmaceuticals, Inc.**

-----  
**Dendracin Neurodendraxcin®, Topical Pain Relief Lotion**

**ACTIVE INGREDIENTS**

Methyl Salicylate 30%  
Menthol 10%  
Capsaicin 0.025%

**Purpose**

Topical Analgesic

**USES:**

For temporary relief of mild pain due to muscular strain, arthritis, and simple back pain. Does not cure any disease.

**Keep away from children and pets.**

If swallowed get medical help or contact a Poison Control Center right away.

**WARNINGS:**

For external use only. Do not use in eyes, mouth, on mucous membranes, or genitals. Do not allow treated skin to contact infants or pets. Do not tightly bandage. Do not use with heating pad. Do not use with other topical pain products. May stain furniture, clothing or bedding.

**DIRECTIONS:**

Use only as directed. Shake before each use. Prior to first use, rub small amount to check for sensitivity. Gently rub over painful areas. Dry before contact with clothes or bedding to avoid staining. Wash hands after use. Do not use more than 4 times daily or if pregnant or nursing. If placed into eyes, rinse with cold water and call a doctor.

**Do Not Use:**

On cuts or infected skin, on children less than 12 years old, in large amounts, especially over raw or blistered skin, if allergic to any ingredients, PABA, aspirin products, or sulfa.  
**Store below 90°F/32°C.**

## Stop Use and Ask a Physician:

For severe undiagnosed pain. If pain worsens or persists for more than 7 days. If pain clears up and then recurs in a few days. If itching or rash occurs.

## INACTIVE INGREDIENTS:

Aloe Barbadensis (Aloe Vera) Leaf Juice, Ammonium Acryloyldimethyltaurate/VP Copolymer, Benzocaine, Borago Officinalis Seed Oil (Borage Oil), Cetyl Alcohol, Dimethyl Sulfoxide, DMDM Hydantoin, Glyceryl Stearate, Methylparaben, PEG 100 Stearate, Poloxamer 407, Propylene Glycol, Propylparaben, Lecithin (Soy), Stearic Acid, Sodium Stearoyl Glucamate, Triethanolamine, Water, Officinale Root Extract.

Manufactured for **Physician's Science and Nature Inc.**

220 Newport Center Drive 11-634, Newport Beach, CA 92660

## Principal Display Panel

Physician's Science and Nature Inc.

**Relabeled by Preferred Pharmaceuticals, Inc.**

## DENDRACIN

Neurodendracin®

**Professional Formula**

*Dermatologically Tested*

*Hypoallergenic*

*Topical Pain Relief Lotion*

*Deep Penetrating Action*

NDC 68788-7696

|   |  |   |  |  |
|---|--|---|--|--|
| <p><b>Dendracin Topical Pain Relief Lotion</b><br/>Brand Name Medication<br/>Each mL Contains Methyl Salicylate 30% / Capsaicin 0.025% / Menthol 10%...topical analgesic<br/>Pkg Size: Exp Date:<br/>Lot#: Lot#: Batch#: Batch#: Ins: Ins:<br/>Mfg: Physicians Science and Nature Inc., Newport Beach, Prod#: Prod#:<br/>Warning<br/>For external use only. Do not use in eyes, mouth, on mucous membranes, or genitals. Keep away from children. Do not tightly bandage. Do not use with heating pad. Do not use with other topical pain products. Shake before each use. Wash hands after use. Do not use more than 4 times daily or if pregnant or nursing. If swallowed, call poison control. If placed in eyes, rinse with cold water and call a doctor. Do not use on cuts or infected skin, on children less than 12 years old, in large amounts, especially over raw or blistered skin. If allergic to any ingredients, PABA, aspirin products, or sulfas. Store below 50°F (10°C).</p> | <p><b>PREFERRED</b><br/>Pharmaceuticals, Inc. <small>ANALGESIC, CA 92660</small></p> | <p>CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed</p> | <p>Dendracin Topical Pain Relief Lotion<br/>Qty: Ins:<br/>Lot#: Bat#:<br/>Prod# (NDC):<br/>Dendracin Topical Pain Relief Lotion<br/>Qty: Ins:<br/>Lot#: Bat#:<br/>Prod# (NDC):<br/>Dendracin Topical Pain Relief Lotion<br/>Qty: Ins:<br/>Lot#: Bat#:<br/>Prod# (NDC):<br/>Dendracin Topical Pain Relief Lotion<br/>Qty: Ins:<br/>Lot#: Bat#:<br/>Prod# (NDC):</p> | <p>Log<br/>Chart<br/>Billing<br/>Patient</p> |
| <p>Directions English<br/>Use as directed by your doctor<br/>Apply externally _____ times a day.</p>  |  | <p>Instrucciones Espanol:<br/>Usó según lo dirigido por su doctor<br/>Aplique externamente _____ veces al día.</p>          |  |  |

120 ml (4 fl oz)

## DENDRACIN NEURODENDRAXCIN

methyl salicylate, menthol and capsaicin lotion

### Product Information

|                                |                |                           |                               |
|--------------------------------|----------------|---------------------------|-------------------------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:68788-7696(NDC:27495-014) |
| <b>Route of Administration</b> | TOPICAL        |                           |                               |

### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength            | Strength            |
|---|------------------------------|---------------------|
| <b>METHYL SALICYLATE</b> (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)                    | METHYL SALICYLATE            | 18 g<br>in 60 mL    |
| <b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A) | MENTHOL,<br>UNSPECIFIED FORM | 6 g in 60 mL        |
| <b>CAPSAICIN</b> (UNII: S07O44R1ZM) (CAPSAICIN - UNII:S07O44R1ZM)                                 | CAPSAICIN                    | 0.015 g<br>in 60 mL |

### Inactive Ingredients

| Ingredient Name   | Strength |
|---|----------|
| <b>WATER</b> (UNII: 059QF0KO0R)                                 |          |
| <b>BENZOCAINE</b> (UNII: U3RSY48JW5)                            |          |
| <b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)                 |          |
| <b>PEG-100 MONOSTEARATE</b> (UNII: YD01N1999R)                  |          |
| <b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)                          |          |
| <b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)                      |          |
| <b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)                         |          |
| <b>DIMETHYL SULFOXIDE</b> (UNII: YOW8V9698H)                    |          |
| <b>POLOXAMER 407</b> (UNII: TUF2IVW3M2)                         |          |
| <b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)                        |          |
| <b>BORAGE SEED OIL</b> (UNII: F8XAG1755S)                       |          |
| <b>AMMONIO METHACRYLATE COPOLYMER TYPE A</b> (UNII: 8GQS4E66YY) |          |
| <b>GINGER</b> (UNII: C5529G5JPQ)                                |          |
| <b>SOYBEAN LECITHIN</b> (UNII: 1DI56QDM62)                      |          |
| <b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)                         |          |
| <b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)                         |          |
| <b>DMDM HYDANTOIN</b> (UNII: BYR0546TOW)                        |          |
| <b>SODIUM STEAROYL GLUTAMATE</b> (UNII: 65A9F4P024)             |          |
| <b>TROLAMINE</b> (UNII: 9O3K93S3TK)                             |          |

### Packaging

| # | Item Code | Package Description   | Marketing Start Date | Marketing End Date |
|---|-----------|---|----------------------|--------------------|
|   | NDC 68788 | 60 mL in 1 BOTTLE, Topical Methyl Salicylate, Menthol, and Capsaicin Lotion |                      |                    |

|                              |   |   |                             |                           |
|------------------------------|---|---|-----------------------------|---------------------------|
| <b>1</b>                     | NDC:68788-7696-6                                | 60 mL in 1 BOTTLE; Type 0: Not a Combination Product  | 04/01/2020                  |                           |
| <b>2</b>                     | NDC:68788-7696-1                                | 120 mL in 1 BOTTLE; Type 0: Not a Combination Product | 04/01/2020                  |                           |
| <b>Marketing Information</b> |   |   |                             |                           |
| <b>Marketing Category</b>    | <b>Application Number or Monograph Citation</b> |   | <b>Marketing Start Date</b> | <b>Marketing End Date</b> |
| OTC Monograph Drug           | MO17  |   | 04/01/2020                  |                           |

**Labeler** - Preferred Pharmaceuticals, Inc. (791119022)

**Registrant** - Preferred Pharmaceuticals, Inc. (791119022)

|                                 |                |               |                            |
|---------------------------------|----------------|---------------|----------------------------|
| <b>Establishment</b>            |                |               |                            |
| <b>Name</b>                     | <b>Address</b> | <b>ID/FEI</b> | <b>Business Operations</b> |
| Preferred Pharmaceuticals, Inc. |                | 791119022     | RELABEL(68788-7696)        |

Revised: 1/2025

Preferred Pharmaceuticals, Inc.