

DENDRACIN NEURODENDRAXCIN- methyl salicylate, menthol and capsaicin lotion
Proficient Rx LP

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.

Warnings:

For external use only. Do not use in eyes, mouth, on mucous membranes, or genitals. Do not tightly bandage. Do not use with heating pad. Do not use with other topical pain products.

Keep away from children.

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 swallowed, call poison control. 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:

Water, benzocaine, glyceryl stearate, PEG 100 stearate, stearic acid, cetyl alcohol, propylene glycol, dimethyl sulfoxide, triethanolamine, poloxamer 407, aloe barbadensis gel, borage oil, ammonium acryloyldimethyltaurate, zingiber officinale root extract, methylparaben, propylparaben, soya lecithin, DMDM hydantoin, sodium stearyl glutamate.

Manufactured for Physicians' Science and Nature, Inc.

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

Made in the USA

Patent Pending

Relabeled by:

Proficient Rx LP

Thousand Oaks, CA 91320

Principal Display Panel



Scan Here



NDC 63187-136-02

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Thousand Oaks, CA 91320

Dendracin 30% / 10% / 0.025%
60ml (2 fl oz) Lotion
Lot #:00000 SN# MASTER
NDC 63187-136-02 Exp:00/00/00

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Dendracin 30% / 10% / 0.025%

60ml (2 fl oz) Lotion

Each bottle contains: Methyl Salicylate 30%
Topical Analgesic; Menthol 10% Topical
Analgesic; Capsaicin 0.025% Topical Analgesic

See bottle

Product ID: RD013602

Mfr. For: Physicians' Science and Nature, Inc. 220 Newport Center Drive 11-634,
Newport Beach, CA 92660

Store below 90°F / 32°C

Keep medication out of the reach of children



GTIN: 00363187136024
SN# MASTER
Exp. 00/00/00
Lot #:00000

DENDRACIN NEURODENDRAXCIN

methyl salicylate, menthol and capsaicin lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63187-136(NDC:27495-014)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63187-136-02	50 in 1 CARTON	08/01/2014	
1		60 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:63187-136-04	50 in 1 CARTON	08/01/2014	
2		120 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/22/2011	

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	RELABEL(63187-136) , REPACK(63187-136)

Revised: 1/2024

Proficient Rx LP