#### OVERTIME- methyl salicylate, menthol and capsaicin lotion Physicians Science & Nature Inc.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

-----

### DB08-Overtime <sup>®</sup>Pain Relief Lotion

#### **Active Ingredients**

Methyl Salicylate 30%

Capsaicin 0.0375%

Menthol USP 10%

### 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 OUT OF REACH OF 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 or 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 persist for more than 7 days. If pain clears up and then recurs in a few days. If itching or rash occurs.

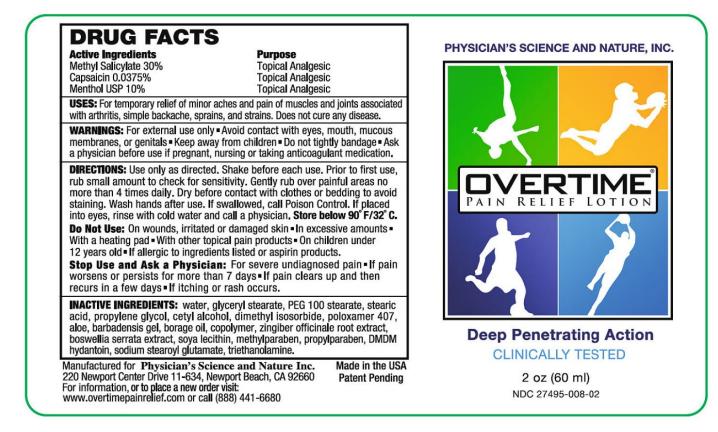
# **INACTIVE INGREDIENTS:**

water, glyceryl Stearate, PEG 100 Stearate, Stearic Acid, Propylene Glycol, Cetyl Alcohol, Dimethyl Isosorbide, Poloxamer 407, Aloe Barbadensis Gel, Borage Oil, Copolymer, Zingiber Officinale Root Extract, Boswellia Serrata Extract, Soy Lecithin, Methylparaben, Propylparaben, DMDM Hydantoin, Sodium Stearoyl Glutamate, Triethanolamine.

Manufactured for **Physician's Science and Nature Inc.** 220 Newport Center Drive 11-634, Newport Beach, CA 92660 Manufactured in the USA

## PHYSICIAN'S SCIENCE AND NATURE, INC.

OVERTIME<sup>®</sup> PAIN RELIEF LOTION Deep Penetrating Action With Natural Pain Relieving Ingredients Dermatologically Tested Hypoallergenic NDC 27495-008-02 2 Oz. Label



#### NDC 27495-008-04

4 Oz. Label



# OVERTIME

methyl salicylate, menthol and capsaicin lotion

Product Information				
Product Type	HUMAN OTC DRUG Item Code (Source)		NDC:27495-008	
Route of Administration	TOPICAL			
	<b>N A a b <b>b a b a b <b>b a b b <b>a b b a b <b>b a b b a b b a b b a b b a b b a b b a b b a b b a b b a b b a b</b></b></b></b></b>			
Active Ingredient/Active	emolety			
Ingre	edient Name		Basis of Strength	Strength
METHYL SALICYLATE (UNII: LAV UNII:0414PZ4LPZ)	5U5022Y) (SALICYLIC ACID -		METHYL SALICYLAT	E 18 g in 60 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)			MENTHOL	6 g in 60 mL
CAPSAICIN (UNII: S07044R1ZM) (CAPSAICIN - UNII:S07044R1ZM) CAPSAICIN				0.0225 g in 60 mL
Inactive Ingredients				
	Ingredient Name			Strength
WATER (UNII: 059QF0KO0R)				
GLYCERYL MONOSTEARATE (U				

PEG-100 STEARATE (UNII: YD01N1999R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
BORAGE SEED OIL (UNII: F8XAG1755S)	
GINGER (UNII: C5529G5JPQ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC10H)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
TROLAMINE (UNII: 903K93S3TK)	
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)	

# Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date						
1	NDC:27495-008- 02	50 in 1 CARTON	01/01/2007							
1		60 mL in 1 BOTTLE; Type 0: Not a Combination Product								
2	NDC:27495-008- 04	50 in 1 CARTON	01/01/2007							
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						
	approved drug ner		01/01/2007							

Labeler - Physicians Science & Nature Inc. (012485755)

Registrant - Westwood Laboratories, LLC (832280635)

Revised: 11/2023

Physicians Science & Nature Inc.