NATURES SUNSHINE PRODUCTS EVERFLEX TOPICAL ANALGESIC PAIN RELIEF - menthol cream
Wasatch Product Development, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Everflex Topical Analgesic Pain Relief Cream

Active Ingredients	Purpose
Menthol 1.25%	Topical analgesic

Everflex pain relief cream provides temporary relief from aching joints and the surrounding tissues. Ever flex contains a proprietary blend of esterified fatty acids which is clinically proven to provide arthritis relief, plus MSM.

-Keep out of reach of children

-If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of product and consult a physician

-Do not apply to wounds or damaged skin

-Do not bandage tightly

Other Information: Store in a cool, dry place.

-For external use only

-Avoid contact with eyes

Directions for use: Massage Everflex into painful areas 3 to 4 times per day.

Inactive Ingredients: Water, Glyceryl Stearate, Glycerin, Methylsulfonylmethane (MSM), Cetyl Myristoleate, Olea Europaea (Olive) Fruit Oil, Cetyl Myristate, Lecithin, Menthol, Tocopheryl Acetate, Benzyl Alcohol, Phenoxyethanol, Carbomber, PEG-100 Stearate, Potassium Hydroxide, Cetyl Palmitoleate, Cetyl Oleate, Cetyl Palmitate, Cetyl Laurate, Methylparaben, Propylparaben, Ethylparaben, Butylparaben, Isobutylparaben.

Everflex Topical Analgesic Pain Relief Cream with MSM

2 Oz. (57 Grams)

NATURES SUNSHINE PRODUCTS EVERFLEX TOPICAL ANALGESIC PAIN RELIEF

menthol cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:44717-532
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1.25 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETYL MYRISTOLEATE (UNII: 87P8K33Q5X)	
OLIVE OIL (UNII: 6UYK2W1W1E)	
CETYL MYRISTATE (UNII: 7OPL833Q4D)	
MENTHOL (UNII: L7T10EIP3A)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PEG-100 STEARATE (UNII: YD01N1999R)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
CETYL PALMITOLEATE (UNII: 962I97113K)	
CETYL OLEATE (UNII: 78K2L26L8N)	

CETYL PALMITATE (UNII: 5ZA2S6B08X)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
ETHYLPARABEN (UNII: 14255EXE39)	
BUTYLPARABEN (UNII: 3QPI1U3FV8)	
ISOBUTYLPARABEN (UNII: 0QQJ25X58G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:44717-532-01	57 g in 1 JAR		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	08/13/2010	

Labeler - Wasatch Product Development, LLC (962452533)

Registrant - Nature's Sunshine Products, Inc. (081832388)

Establishment

Name	Address	ID/FEI	Business Operations
Wasatch Product Development, LLC		962452533	manufacture

Revised: 3/2011 Wasatch Product Development, LLC