## OXYRUB PAIN RELIEF- menthol cream Pure Source, LLC

-----

OxyRub Pain Relief Cream

## OxyRub Pain Relief Cream

#### **Active Ingredient**

Menthol (1.25%)

#### **Purpose**

**Topical Analgesic** 

#### Uses

temporarily relieves minor aches and pains of muscles and joints associated

- simple backaches
- arthritis
- strains
- sprains
- bruises

#### **Warnings**

For external use only.

#### Do not use

redness over the affected area t **Ask a doctor before use if you haveWhen using this produc** 

# Stop use and consult a doctor if

- Use only as directed.
- with a heating pad, may blister skin
- on open wounds or damaged skin
- Avoid contact with eyes
- Do not bandage tightly
- skin redness or excessive skin irritation develops
- condition worsens or symptoms persist for more than 7 days
- symptoms clear up and occur again in a few days

## Keep out of reach of children

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

#### **Directions**

. Apply to affected area no more than 3-4 times daily. consult a doctor. Adults and children 12 years of age and olderChildren under 12 years of age,

#### Other information

Store at room temperature. For Lot Number and Expiration Date, see crimp at end of tube.

## **Inactive Ingredients**

Bitter Orange Oil, C13-14 Isoparaffin, Caprylhydroxomic Acid, Caprylyl Glycol, Eucalyptus Oil, Glycerin, Laureth-7, Oxidized Corn Oil, Polyacrylamide (1000 MW), Polysorbate 20, Water

OxyRub Pain Relief Cream 4oz/114g (65121-113-04)



## **OXYRUB PAIN RELIEF**

menthol cream

Product Type HUMAN OTC DRUG Item Code (Source) NDC:65121-113

Route of Administration TOPICAL

## **Active Ingredient/Active Moiety**

Ingredient NameBasis of StrengthStrengthMENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)MENTHOL1.25 g in 100 g

Inactive Ingredients					
Ingredient Name	Strength				
BITTER ORANGE OIL (UNII: 9TLV70SV6I)					
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)					
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)					
CAPRYLYL GLYCOL (UNII: 00YIU5438U)					
EUCALYPTUS OIL (UNII: 2R04ONI662)					
GLYCERIN (UNII: PDC6A3C0OX)					
LAURETH-7 (UNII: Z95S6G8201)					
CORN OIL (UNII: 8470G57WFM)					
POLYSORBATE 20 (UNII: 7T1F30V5YH)					
WATER (UNII: 059QF0KO0R)					

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:65121-113- 04	114 g in 1 TUBE; Type 0: Not a Combination Product	05/01/2013	

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M017	05/01/2013				

# Labeler - Pure Source, LLC (080354456)

# Registrant - Pure Source, LLC (080354456)

Establishment						
Name	Address	ID/FEI	<b>Business Operations</b>			
Pure Source, LLC		080354456	manufacture(65121-113)			

Revised: 11/2023 Pure Source, LLC