GOOD NEIGHBOR PHARMACY COLD AND HOT THERAPY PAIN RELIEF BALM EXTRA STRENGTH- menthol, methyl salicylate ointment AmerisourceBergen Drug Corporation

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.

Good Neighbor Pharmacy Cold and Hot Therapy Pain Relief Balm

Drug Facts

Active ingredients

Menthol 7.6%

Methyl Salicylate 29%

Purpose

Topical Analgesic

Uses

Temporarily relieves minor pain associated with:

- arthritis simple backache muscle strains bruises
- muscle sprains cramps

Warnings

For external use only.

Allergy Alert:

If prone to allergic reaction from aspirin or salicylates, consult a doctor before use.

When using this product

• use only as directed • avoid contact with eyes or mucus membranes • do not bandage tightly or use with a heating pad • do not apply to wounds or damaged skin

Stop use and ask doctor if

• condition worsens • redness is present • irritation develops • symptoms persist for more than 7 days or clear up and occur again within a few days

If pregnant or breast-feeding

ask a healthcare professional before use.

Keep out of reach of children.

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

Directions

Adults and children over 12 years: • apply generously to affected area. • massage into painful area until thoroughly absorbed into skin. • repeat as necessary, but not more than 4 times daily. Children under 12 years: ask a doctor.

Inactive ingredients

Paraffin, White Petrolatum.

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menthol, methyl salicylate ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:24385-241
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
MENTHOL (UNII: L7T10 EIP3A) (MENTHOL - UNII:L7T10 EIP3A)	MENTHOL	7.6 g in 100 g		
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	29 g in 100 g		

Inactive Ingredients				
Ingredient Name	Strength			
PARAFFIN (UNII: 19 O 0 E 3 H 2 Z E)				
PETROLATUM (UNII: 4T6H12BN9U)				

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
1 NDC:24385-241-54	99.2 g in 1 JAR; Type 0: Not a Combination Product				
Marketing Inf	ormation				
Marketing Inf		Marketing Start Date	Marketing End Date		
	ry Application Number or Monograph Citation	Marketing Start Date 05/24/2000	Marketing End Date		

Labeler - AmerisourceBergen Drug Corporation (007914906)

Establishment				
Name	Address	ID/FEI	Business Operations	
Product Quest Mfg, LLC		927768135	manufacture(24385-241), label(24385-241)	

Revised: 7/2015 AmerisourceBergen Drug Corporation