MOTION MEDICINE- camphor, menthol cream
Motion Medicine Inc.

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.

Active Ingredients (Percent by weight)	**Purpose**
Menthol 4.0% .....Analgesic	(Pain reliever)
Camphor 4.0% .....Analgesic	(Pain reliever)

For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains, bruises and sprains, etc.

Warnings

- Avoid getting into eyes or on mucous membranes.
- If condition worses, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product and consult a doctor.
- Do not apply to wounds or damaged skin. Do not bandage tightly.

Keep out of the reach of children

- Use only as directed.
- If swallowed, get medical help or contact a Poison Control center right away.

Directions

Adults and children 12 years of age or older: Rub this soothing cream on the affected area not more than 3 to 4 times daily.

Children under the age of 12: Do not use, consult a doctor.

Water purified, methylsulfonylmethane (MSM), cetyl alcohol, glyceryl stearate, steareth-100, glycerin, glucosamine sulfate,

stearyl alcohol, polysorbate 20, urea, phenoxyethanol, caprylyl glycol, sorbic acid, polyacrylamide, C-13-14 isoparaffin, laureth-7,

eucalyptus oil, grape seed oil, methyl salicylate, tocopheryl acetate, alcohol denatured, thymol, lavender oil, chondroitin sulfate.

Pain Reliever

Distributed by:

Motion Medicine Inc.

84 Sunmeadows Crescent SE
Calgary, AB CANADA T2X 3H3

1-866-972-8466

120g

M otion Medicine

Eases pain, improves movement, and is highly effective

Topical Remedy

MOTION MEDICINE

camphor, menthol cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50963-809
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	4 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	4 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
STEARETH-100 (UNII: 4OH5W9UM87)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
UREA (UNII: 8W8T17847W)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
SORBIC ACID (UNII: X045WJ989B)	
POLYACRYLAMIDE (1500 MW) (UNII: 5D6TC4BRWV)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
LAURETH-7 (UNII: Z95S6G8201)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GRAPE SEED OIL (UNII: 930MLC8XGG)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALCOHOL (UNII: 3K9958V90M)	
THYMOL (UNII: 3J50XA376E)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
CHONDROITIN SULFATE SODIUM (BOVINE) (UNII: 8QTV3DTT8W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50963-809-94	1 in 1 BOX	09/01/2009	
1		120 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:50963-809-05	500 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2009	
3	NDC:50963-809-16	7 g in 1 JAR; Type 0: Not a Combination Product	09/01/2009	
4	NDC:50963-809-88	2 g in 1 POUCH; Type 0: Not a Combination Product	09/01/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	09/01/2009	

Labeler - Motion Medicine Inc. (248579919)

Registrant - Nutri-Dyn Products Ltd. dba Professional Health Products (209901511)

Establishment

Name	Address	ID/FEI	Business Operations
Enterprises ImportFab Inc.		248586117	manufacture(50963-809)

Revised: 10/2022 Motion Medicine Inc.