DOCTOR HOYS PAIN RELIEF GEL ROLL-ON- menthol, camphor (synthetic) gel
Dynamic Blending Specialists, Inc.

Doctor Hoy's Pain Relief Gel Roll-On

Drug Facts:

Active ingredients:

Menthol (5%)

Camphor (5%)

Purpose:

Topical Analgesic

Uses:

Temporary relief of minor aches and pains of muscles and joints due to: simple backache • arthritis • sprains • strains

Warnings:

• For external use only

• Use as directed

• Avoid contact with eyes or mucus membranes

• Do not apply to open wounds, irritated or damaged skin

Keep Out of reach of children

If swallowed, get medical help or contact a Poison Control Center immediately

If pregnant or breast feeding,

• ask a health professional before use

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

• Use caution with wrap or heating pad.

Directions:

For adults and children 2 years and older: For best results, shake well and apply generously. Use 1 layer for minor pain, 2 layers for medium and 3 layers for severe symptoms • Rub in well and allow to dry between layers (usually 2-3 minutes) • Repeat

process up to 4 times daily • for children under 2 years of age: Consult a physician before use.

Inactive ingedients:

Alcohol, Ammonium Acryloyldimethyltaurate/VP Copolymer, Arnica Montana (Arnica Flower) Extract, Biosaccharide Gum-1, Butyrospermum Parkii (Shea) Butter Extract, Ethylhexylglycerin, Euphorbia Cerifera (Candelilla) Wax, Glycerin, Hamamelis Virginiana (Witch Hazel) Water, Hydrated Silica, Jojoba Esters, Phenoxyethanol, Polyglyceryl-10 Caprylate/Caprate, SD Alcohol 40-B, Sodium Starch Octenylsuccinate, Tocopherol Acetate, Water, Zanthoxylum Bungeanum Fruit Extract.

75638-011-00

DOCTOR HOYS PAIN RELIEF GEL ROLL-ON

menthol, camphor (synthetic) gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75638-011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	50 mg in 1 mL
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ARNICA MONTANA WHOLE (UNII: O80TY208ZW)	
BIOSACCHARIDE GUM-1 (UNII: BB4PU4V09H)	
ALCOHOL (UNII: 3K9958V90M)	
SHEA BUTTER (UNII: K49155WL9Y)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CANDELILLA WAX (UNII: WL0328HX19)	
GLYCERIN (UNII: PDC6A3C0OX)	
HAMAMELIS VIRGINIANA TOP WATER (UNII: NT00Y05A2V)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
ZANTHOXYLUM BUNGEANUM FRUIT (UNII: 3CIP16A418)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75638-011-00	89 mL in 1 TUBE, WITH APPLICATOR; Type 0: Not a Combination Product	03/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/01/2023	

Labeler - Dynamic Blending Specialists, Inc. (085704438)

Revised: 1/2025 Dynamic Blending Specialists, Inc.