DOCTOR HOYS NATURAL PAIN RELIEF- camphor, menthol gel
DOCTOR HOY'S, 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.

Pain Relief Gel

Active Ingredients	Purpose
Camphor (5%)	External Analgesic
Menthol (5%)	External Analgesic

Uses:
for the temporary relief of minor aches and pains of muscles and joints due to:
* simple backache * arthritis * sprains * strains * bruises

Warnings:
* for external use only * do not apply to wounds or damaged skin or bandage tightly * avoid contact with eyes * Keep out of reach of children. if swallowed, get medical help or contact a Poison Control Center immediately * 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 this product and consult a physician * Pregnancy-breast feeding warning: if pregnant or breast feeding, ask a health professional before use * do not bandage tightly

Inactive Ingredients:

distilled water, hamamelis virginiana (witch hazel extract), isopropyl alcohol, arnica Montana, oleyl alcohol and zanthoxylum alatum (scezhuan pepper), Bio-saccharide Gum -1, glycerin, ammonium acryloyldimethyltaurate/VP copolymer, beta cyclodextrin, menthol, potassium hydroxide, mannitol, cellulose chromium, hydroxide green, tocopheryl acetate (vitamin E), hydroxypropyl methycellulose

Directions:
for adults and children 2 years and older; apply to affected area not more than 3 to 4 times daily. "Children under 2 years of age: consult a doctor." Apply generously to affected and surrounding areas. Rub in well. Use 1 application for minor pain, 2 for medium and 3 applications for severs symptoms. Allow to dry between applications (usually just 2-3 minutes).

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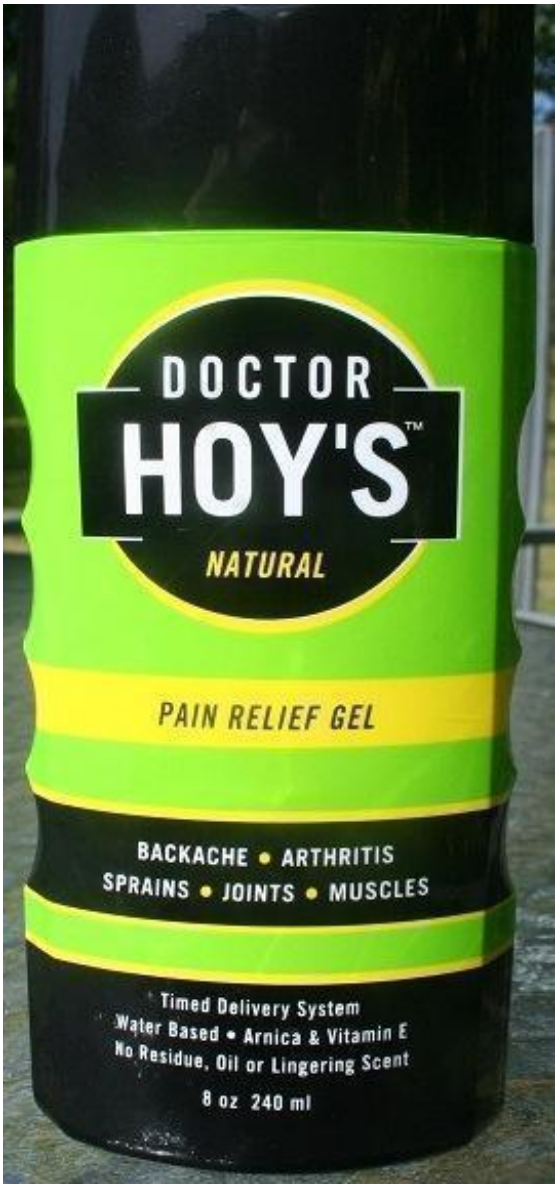
An image/jpeg of the container label is included in this section.

ID: MM1



DOCTOR HOYS NATURAL PAIN RELIEF

camphor, menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10842-102
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 (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	50 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
EUPHORBIA ANTISYPHILITICA WHOLE (UNII: 82A88H0RIQ)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
GLYCERIN (UNII: PDC6A3C0OX)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
ARNICA MONTANA (UNII: O80TY208ZW)	
WATER (UNII: 059QF0KO0R)	
WITCH HAZEL (UNII: 101I4J0U34)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYGLYCERYL-10 CAPRYLATE (UNII: YS396CQX5C)	
ZANTHOXYLUM BUNGEANUM FRUIT (UNII: 3CIP16A418)	
BIOSACCHARIDE GUM-1 (UNII: BB4PU4V09H)	
HYDRATED SILICA (UNII: Y6O7T4G8P9)	
OLEYL ALCOHOL (UNII: 172F2WN8DV)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
MANNITOL (UNII: 3OWL53L36A)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
MODIFIED CORN STARCH (1-OCTENYL SUCCINIC ANHYDRIDE) (UNII: 461P5CJN6T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10842-102-01	237 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	05/10/2005	
2	NDC:10842-102-02	1814 mL in 1 JUG; Type 0: Not a Combination Product	05/10/2005	
3	NDC:10842-102-04	5 mL in 1 PACKET; Type 0: Not a Combination Product	05/10/2005	
4	NDC:10842-102-05	946 mL in 1 JUG; Type 0: Not a Combination Product	08/05/2015	
5	NDC:10842-102-08	89 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	09/01/2015	
6	NDC:10842-102-09	3785 mL in 1 JUG; Type 0: Not a Combination Product	05/10/2005	
7	NDC:10842-102-03	118 mL in 1 TUBE; Type 0: Not a Combination Product	09/01/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	05/10/2005	

Labeler - DOCTOR HOY'S, LLC (791882322)

Registrant - DOCTOR HOY'S, LLC (791882322)

Establishment

Name	Address	ID/FEI	Business Operations
Specialty Formulations and Manufacturing LLC		003989912	manufacture(10842-102) , label(10842-102)

Establishment

Name	Address	ID/FEI	Business Operations
Dynamic Blending Specialists, Inc.		085704438	manufacture(10842-102)

Establishment

Name	Address	ID/FEI	Business Operations
United Laboratories Manufacturing, LLC		807878116	manufacture(10842-102) , label(10842-102)

Revised: 12/2022

DOCTOR HOY'S, LLC