DOCTORS TOUCH ERASE THE PAIN- menthol liquid
DOCTOR'S CHOICE MEDICAL, LLC

DOCTOR'S TOUCH Erase The Pain

Drug Facts

Active Ingredients

Menthol 5.00%

Purpose

Topical Analgesic

Uses:

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

Warnings:

For external use only

Do not use

- on damaged or broken skin

When using this product

- Avoid contact with the eyes.
- Do not bandage tightly.

Stop use and ask a doctor if

- rash or irritation develops and lasts.
- condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children.

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

If pregnant or breast-feeding,

ask a health professional before use

Directions:

- Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily.

- Children under 2 years of age: consult a doctor.

Other information:

- Protect the product in this container from excessive heat and direct sun.

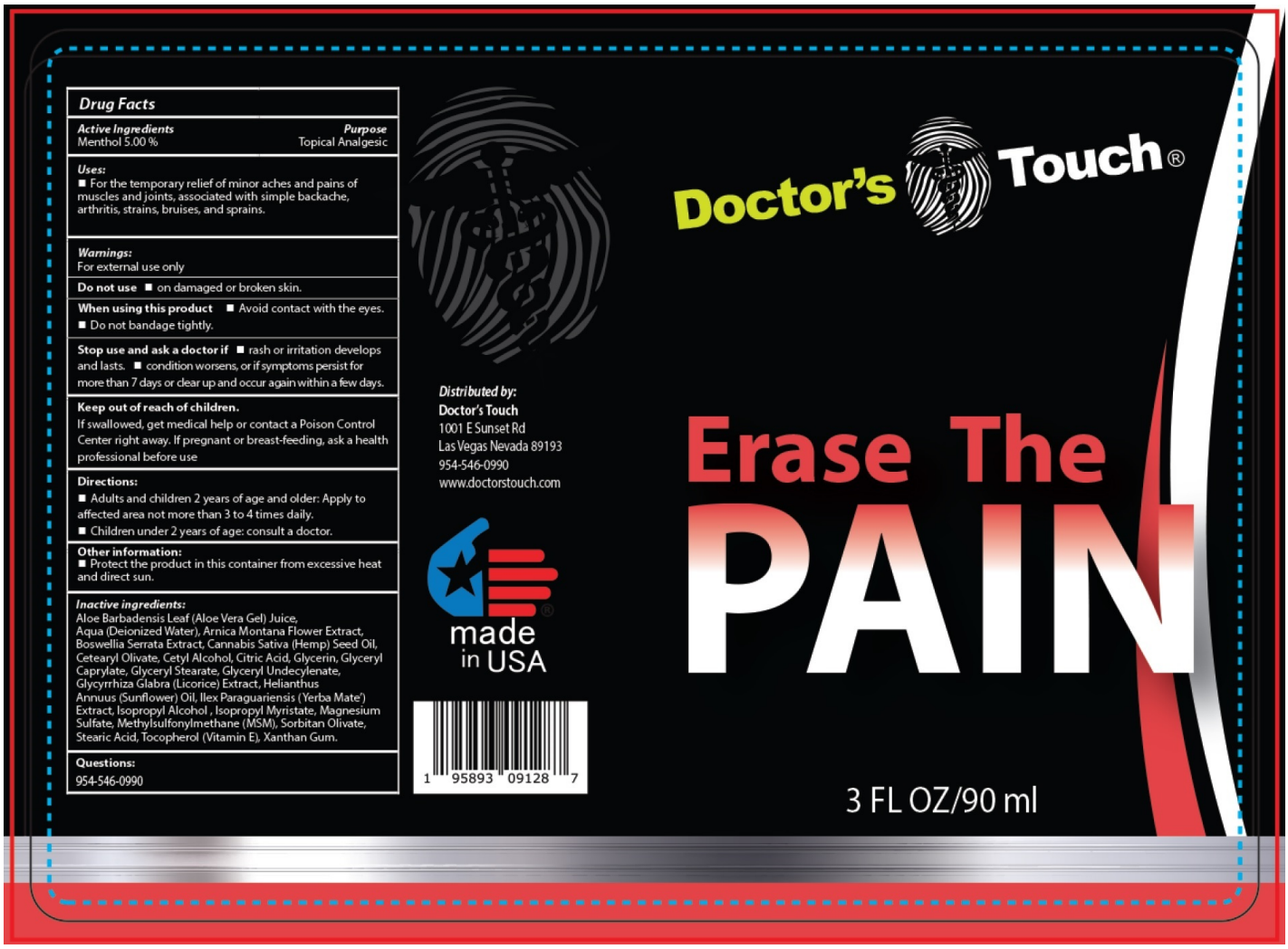
Inactive ingredients:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, Cannabis Sativa (Hemp) Seed Oil, Cetearyl Olivate, Cetyl Alcohol, Citric Acid, Glycerin, Glyceryl Caprylate, Glyceryl Stearate, Glyceryl Undecylenate, Glycyrrhiza Glabra (Licorice) Extract, Helianthus Annuus (Sunflower) Oil, Ilex Paraguariensis (Yerba Mate') Extract, Isopropyl Alcohol, Isopropyl Myristate, Magnesium Sulfate, Methylsulfonylmethane (MSM), Sorbitan Olivate, Stearic Acid, Tocopherol (Vitamin E), Xanthan Gum.

Questions:

954-546-0990

Package Labeling:



DOCTORS TOUCH ERASE THE PAIN

menthol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80045-396
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

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOCAPRYLATE (UNII: TM2TZD4G4A)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TOCOPHEROL (UNII: R0ZB2556P8)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:80045-396-00	90 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/27/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/27/2023	

Labeler - DOCTOR'S CHOICE MEDICAL, LLC (081045011)

Revised: 11/2023

DOCTOR'S CHOICE MEDICAL, LLC