STOPAIN CLINICAL- menthol, unspecified form gel
Troy Manufacturing, Inc.

Stopain Clinical Gel

Drug Facts

Active Ingredient

Menthol 10.0%

Purpose

Topical Analgesic

Uses

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

Warnings

Avoid contact with eyes. **For external use only.**

Flammable: Keep away from fire or flame.

When using this product
- use only as directed
- do not bandage tightly or use a heating pad
- do not apply to wounds or damaged skin

Stop use and ask a doctor if
- condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days
- redness is present
- excessive irritation of the skin develops

If pregnant or breast-feeding,

ask a health 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:

ask a doctor. **Children 12 years or younger:**

- apply directly onto affected area without the need to bandage
- repeat if necessary, but do not apply more than 4 times daily.

Other Information

- Store at room temperature

Inactive Ingredients

acrylic acid/vinyl ester copolymer, dimethylsulfone (MSM), eucalyptus oil, glucosamine sulfate, hydroxypropylcellulose, SD alcohol 39C, PEG-8 dimethicone, pentylene glycol, peppermint oil, triethanolamine, water (USP).

Package Labeling:118ml

Stopain®

CLINICAL

MAXIMUM PERFORMANCE
PAIN RELIEF

Immediate, Penetrating Relief from
Arthritis / Muscle Aches / Joint & Back Pain
With MSM & Glucosamine

GEL

4 FL OZ (118mL)

MAXIMUM PERFORMANCE PAIN RELIEF

FOR PROFESSIONAL RE-SALE ONLY

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For more information visit www.StopainClinical.com

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MADE IN USA

TROY HEALTHCARE

Manufactured for: Troy Healthcare, LLC
130 Lions Drive • Hazleton, PA 18202
800-201-PAIN (7246)

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Package Labeling:473ml

Package Labeling:946ml

Package Labeling:3785ml

STOPAIN CLINICAL

menthol, unspecified form gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63936-9750

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	100 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ACRYLIC ACID (UNII: J94PBK7X8S)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH)	
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63936-9750-3	118 mL in 1 TUBE; Type 0: Not a Combination Product	01/18/2017	
2	NDC:63936-9750-4	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/18/2017	
3	NDC:63936-9750-2	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/18/2017	
4	NDC:63936-9750-1	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/18/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/18/2017	

Labeler - Troy Manufacturing, Inc. (160075248)

Registrant - Troy Manufacturing, Inc. (160075248)

Revised: 10/2023 Troy Manufacturing, Inc.