

TOMMIE COPPER PAIN RELIEF- menthol 16% spray
Tommie Copper, Inc.

Pain Relief Liquid Spray

Menthol 16%

Topical Analgesic

For the temporary relief of minor aches and pain associated with simple backaches, arthritis, strains, bruises, and sprains.

For external use only. Flammable--Keep away from fire or flame. **When using this product** avoid contact with eyes. In case of contact with eyes, flush thoroughly with water, do not apply to wounds or damaged skin, and do not bandage tightly, do not use with a heating pad. Contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120F. . **Stop use and ask a doctor if** condition worsens, if symptoms persist for more than 7 days or clear up and occur again within a few days.

If pregnant or breast-feeding ask a health professional.

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

Shake well before use. Adults and children 12 years of age and older, apply to the affected area not more than 3 to 4 times daily. Children under 12 years of age: ask a doctor.

Alcohol Denat., Aloe Barbadensis (Aloe Vera) Leaf Extract, Arnica Montana (Arnica) Flower Extract, Calendula Officinalis (Marigold) Flower Extract, Camellia Sinensis (Green Tea) Leaf Extract, Echinacea Angustifolia (Coneflower) Extract, Glycerin, Ilex Paraguariensis (Yerba Mate) Leaf Extract, Juniperus Communis (Juniper) Fruit Extract, Propylene Glycol, Water

FOLD



PAIN RELIEF LIQUID SPRAY

TOMMIE COPPER

COOLING MENTHOL 16%

Temporary relief for minor pain from arthritis, backache, joint & muscle pain



ALOE



CHAMOMILE

NET WT 4 OZ (113g)

FOLD

Drug Facts

Active ingredient	Purpose
Menthol 16%	Topical Analgesic

Uses For the temporary relief of minor aches and pain associated with • simple backache • arthritis • strains • bruises • sprains.

Warnings

For external use only.

Flammable - Keep away from fire or flame.

When using this product avoid contact with eyes. In case of contact with eyes, flush thoroughly with water • do not apply to wounds or damaged skin • do not bandage tightly • do not use with heating pad • contents under pressure. Do not puncture or incinerate. Do not store at temperature above 120° F.

Stop use and ask a doctor if • condition worsens • if symptoms persist for more than 7 days or clear up and occur again within a few days.

If pregnant or breast-feeding ask a health professional.

Keep out of the reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions • shake well before use • adults and children 12 years or older: apply to affected area not more than 3 to 4 times daily • children under 12 years of age: ask a doctor.

Inactive ingredients Alcohol Denat., Aloe Barbadensis (Aloe Vera) Leaf Extract, Arnica Montana (Arnica) Flower Extract, Calendula Officinalis (Marigold) Flower Extract, Camellia Sinesis (Green Tea) Leaf Extract, Chamomilla Recutita (Matricaria) Flower Extract, Echinacea Angustifolia (Coneflower) Extract, Glycerin, Ilex Paraguariensis (Yerba Mate) Leaf Extract, Juniperus Communis (Juniper) Fruit Extract, Propylene Glycol, Water.

THIS PRODUCT DOES NOT CONTAIN COPPER.

DISTRIBUTED BY:
TOMMIE COPPER, INC.
74 SOUTH MOGER AVE. MOUNT KISCO,
NEW YORK 10549 1.855.692.8291
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TOMMIE COPPER PAIN RELIEF

menthol 16% spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72562-910
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	16 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CALENDULA ARVENSIS LEAF (UNII: 3U3U118F2L)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

GLYCERIN (UNII: PDC6A3C00X)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALCOHOL (UNII: 3K9958V90M)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B404F)	
MATRICARIA CHAMOMILLA FLOWERING TOP OIL (UNII: SA8AR2W4ER)	
ECHINACEA ANGUSTIFOLIA (UNII: VB06AV5US8)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
JUNIPER BERRY (UNII: O84B5194RL)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72562-910-04	113 g in 1 CAN; Type 0: Not a Combination Product	11/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/30/2020	

Labeler - Tommie Copper, Inc. (081176569)

Registrant - Derma Care Research Labs, LLC (116817470)

Establishment

Name	Address	ID/FEI	Business Operations
Derma Care Research Labs, LLC		116817470	manufacture(72562-910)

Revised: 12/2024

Tommie Copper, Inc.