

ULTRA STRENGTH PAIN RELIEF- camphor 4%, menthol 10%, methyl salicylate 30% spray
Product Quest Mfg

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.

Drug Facts

Active ingredients	Purpose
Camphor – 4.00%	Topical Analgesic
Menthol – 10.00%	Topical Analgesic
Methyl Salicylate – 30.00%	Topical Analgesic

Uses

- temporarily relieves minor pain associated with:
- arthritis
- simple backache
- muscle strains
- bruises
- muscle sprains

Warnings

☐ **For external use only**

Flammable

- keep away from fire or flame. • avoid long term storage above 104°F. • do not puncture or incinerate. Contents under pressure. • do not store at temperatures above 120°F
- do not use while smoking or near heat or flame
- avoid long term storage above 104F
- do not puncture or incinerate. Contents under pressure
- do not store at temperature above 120F

When using this product

- avoid contact with eyes and mucous membranes
- do not apply to wounds or damages skin
- do not bandage tightly or use with a heating pad
- use only as directed

Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days

Keep out of reach of the 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

- shake well
- adults and children 2 years of age and older: spray on affected area, not more than 3 to 4 times daily
- children under 2 years of age: consult a doctor

Inactive ingredients

Alcohol Denat.





Since 1933, Scherer Labs™ has been providing Quality, Innovative and Value oriented family-care products.

Pain Relief

**Ultra Strength
Continuous Spray**

Camphor 4%
Menthol 10%
Methyl Salicylate 30%

Deep pain relieving spray
**Temporary relief from minor arthritis,
backache, muscle & joint pain**
Sprays at any angle



NET WT. 4 OZ. (113 g)

Drug Facts

Active ingredients	Purpose
Camphor 4%.....	Topical Analgesic
Menthol 10%.....	Topical Analgesic
Methyl Salicylate 30%.....	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
For external use only.

Flammable • keep away from fire or flame. • avoid long term storage above 104°F. • do not puncture or incinerate. Contents under pressure. • do not store at temperatures above 120°F.

When using this product • avoid contact with eyes and mucous membranes • do not apply to wounds or damaged skin • do not bandage tightly or use with a heating pad

Stop use and ask doctor if • condition worsens • 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 immediately.

If pregnant or breast-feeding, ask a health professional before use.

Directions • adults and children 2 years of age and older: spray product on affected area, not more than 3 to 4 times daily. • children under 2 years of age: consult a doctor.

Inactive ingredient Alcohol Denat.

Questions? 866-483-2846



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Manufactured by:
Product Quest Mfg, LLC.
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 Daytona Beach, FL 32117
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ULTRA STRENGTH PAIN RELIEF

camphor 4%, menthol 10%, methyl salicylate 30% spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:64048-5114
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Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	4 g in 100 g
Menthol (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	Menthol	10 g in 100 g
Methyl Salicylate (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	Methyl Salicylate	30 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64048-5114-4	113 g in 1 CAN; Type 0: Not a Combination Product	02/06/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/06/2014	

Labeler - Product Quest Mfg (927768135)

Registrant - Product Quest Mfg (927768135)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quest Mfg		927768135	manufacture(64048-5114) , label(64048-5114)

Revised: 6/2018

Product Quest Mfg