

**QUALITY CHOICE ULTRA STRENGTH PAIN RELIEVING CREAM ULTRA STRENGTH- camphor, menthol, methyl salicylate cream
QUALITY CHOICE (Chain Drug Marketing Association)**

Quality Choice Ultra Strength Pain Relieving Cream

Active ingredient

Camphor 4%

Menthol 10%

Methyl salicylate 30%

Purpose

Camphor - Topical analgesic

Menthol - Topical analgesic

Methyl salicylate - Topical analgesic

Uses

temporarily relieves minor aches and pains of muscles and joints associated with:

- simple backache
- arthritis
- strains
- sprains
- bruises

Warnings

For external use only

Do not Use

- on wounds or damaged skin
- with a healing pad
- on a child under 12 years of age with arthritis-like conditions

Ask a doctor before use if you have

redness over the affected area

When using this product

- use only as directed
- avoid contact with eyes or mucous membranes
- do not bandage tightly

Stop use and ask a doctor if

- skin redness or excessive irritation of the skin occurs
- 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 right away (1-800-222-1222)

Directions

- adults and children 12 years and over; apply to affected area not more than 3 to 4 times daily
- children under 12 years of age; consult a physician

Other Information

- Store at 20-25 C (68-77F)
- do not use if tube seal under cap is broken

Inactive Ingredients

carbomer homopolymer type c, cetostearyl alcohol, cetyl alcohol, glyceryl monostearate, methylparaben, peg-100 stearate, polysorbate 60, propylparaben, stearic acid, trolamine, water

Principal Display Panel

GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)
CETYL ALCOHOL (UNII: 936JST6JCN)
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)
POLYOXYL 100 STEARATE (UNII: YD01N1999R)
POLYSORBATE 60 (UNII: CAL22UVI4M)
WATER (UNII: 059QF0KO0R)
METHYLPARABEN (UNII: A2I8C7HI9T)
STEARIC ACID (UNII: 4ELV7Z65AP)
TROLAMINE (UNII: 9O3K93S3TK)
PROPYLPARABEN (UNII: Z8IX2SC1OH)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-692-04	1 in 1 CARTON	03/01/2021	
1		113 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/01/2021	

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Establishment

Name	Address	ID/FEI	Business Operations
ANICARE PHARMACEUTICALS PRIVATE LIMITED		916837425	manufacture(63868-692)

Revised: 12/2024

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