PAIN RELIEF- analgesic menthol spray
Unifirst First Aid Corporation

Pain Relief Spray Green Guard

Drug Facts

Active ingredients

USP Menthol 7%

Purpose

Topical analgesic

Uses

Temporary relief of minor aches and pains of muscles and joints.

Warnings

For external use only. Flammable. Keep away from flame.

Do not use
- on open wounds or damaged skin

Ask a doctor before use if
- you are prone to allergic reactions to salicylates, including aspirin

When using this product
- avoid contact with eyes

Stop use and ask a doctor if
- condition worsens
- symptoms persist for more than 7 days or clear up and recur again within a few days
- excessive skin irritation occurs

If pregnant or breast feeding,

ask a doctor 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 (2 years and older):

Shake well and aply to affected araea not more than 3-4 times daily.

Do not bandage tightly.

Children under 2 years:

Consult a doctor

Other information

- store at room temperature 56°-86°F (15°-30°C)
- will not stain clothing

Inactive ingredients

coconut oil, eucalyptus lkeaf oil, glycerol, isopropyl alcohol, peppermint oil, water, wintergreen leaf oil

Questions or comments?

1-800-869-6970

Green Guard Pain Relief Spray Label

Pain Relief spray

Fast Acting pain Relief

For Temporary Relief of Minor Aches and Pains

Green Guard®

2 Fl. Oz. (59.1 mL)

PAIN RELIEF

analgesic menthol spray

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47682-321
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	70 mg in 1 L

Inactive Ingredients

Ingredient Name	Strength
COCONUT OIL (UNII: Q9L0O73W7L)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-321-02	0.059 L in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	04/01/2019	

Labeler - Unifirst First Aid Corporation (832947092)

Revised: 1/2024

Unifirst First Aid Corporation