# K19 RELIEF- menthol liquid K19 LLC

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**K19 SPRAY RELIEF** 

#### **DRUG FACTS:**

### **Active Ingredient:**

Menthol 1.25%

**Topical Analgesic** 

#### Indications:

For the temporary relief of minor aches and pains of the muscles and joints associated with arthritis, simple backache, sprains, bruises and stains.

# Warnings:

- For external use only.
- Avoid contact with eyes and mucous membranes.
- If symptoms persist for more than 7 days, discontinue use and consult physician.

# Keep Out of reach of children.

- If swallowed, consult physician.
- Do not apply to wounds or damaged skin.
- Do not bandage tightly.

# If pregnant or breast feeding,

contact physician prior to use.

#### Directions:

- Adults and children 2 years of age or older: Apply to affected area not more than 3 to 4 times daily.
- Chidlren under 2 years of age: consult a physician.

#### Additional Information:

Store at room temperature.

# Other Ingredients:

Aqua (Deionized Water), Camphor, Glycerin, Glyceryl Laurate, SD-Alcohol 40B.

# Package Labeling:





Dermatologist Tested and Approved

10 FL 0Z (295 mL)

**Spray Relief™** is formulated for longer lasting, powerful pain relief. Keep moving, keep active, keep pain at bay. **Spray Relief** is the only topical analgesic that uses solubilized crystalline menthol, providing long lasting relief without the side effects of ingestible analgesics. Directional 360° spray targets directly at the pain source.

**Solubilized Crystalline Menthol + Glycerin =** activation and reactivation of the pain relieving menthol, at levels that do not burn the skin or create malodor. Movement and sweating reactivates the crystals which remain on the skin ready for reactivation. Combining with compression gear contains the moisture for up to 8 hours of relief.

Don't be sidelined from aches and pains.

Spray Relief.

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Distributed K19, LLC 4280 Winchell Rd • Mantua, Ohio 44255 (866) 300-7300 • www.kosar19.com Made in the USA menthol liquid

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:83574-413

Route of Administration TOPICAL

# **Active Ingredient/Active Moiety**

Ingredient Name		<b>Basis of Strength</b>	Strength
	MENTHOL (LINII: L7T10FIP3A) (MENTHOL - LINII: L7T10FIP3A)	MENTHOL	12.5 mg in 1 ml

# **Inactive Ingredients**

Ingredient Name	Strength
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WATER (UNII: 059QF0KO0R)

**CAMPHOR (SYNTHETIC)** (UNII: 5TJD82A1ET)

**GLYCERIN** (UNII: PDC6A3C0OX)

GLYCERYL LAURATE (UNII: Y98611C087)

#### **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		295 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/10/2023	

# **Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/10/2023	
OTC Monograph Drug	M017	07/10/2023	

# **Labeler -** K19 LLC (090893478)

Revised: 11/2023 K19 LLC