**RHINOMEL MANUKA- sodium chloride spray**
**Melcare Biomedical Pty Ltd**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.*

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**Rhinomel Manuka+ Nasal Spray**

**Drug Facts**

**Active ingredients**

Sodium chloride

**Purpose**

Nasal Wash

**Uses**

For persons prone to congestion

**Warnings**

• Do not use if hyper-sensitive to bee products, pollens or the preservative benzoic acid;
• Not suitable for persons with irritated nasal linings or suspected sinus infection. The natural acidity of honey may result in transient discomfort;
• For any reaction other than transient discomfort cease use and contact your health care professional;
• Consult your doctor if symptoms persist for more than 1 week, tend to reoccur

or are accompanied by pain or a persistent headache. These could be signs of a serious condition;
• Not recommended for pregnant women or children under 12 years without medical advice;
• Replace cap after use;
• Do not use if packaging is damaged or carton tamper seal is broken before first use.
• Keep out of reach of children

**Directions**

• Adults and Children ages over 12.
• Remove cap.
• Use 1-2 sprays per nostril as needed.
• Direct nozzle into your nostril. Fully depress the pump once or twice. Repeat in other nostril.
• After use, breathe normally – avoid sniffing.
• Wipe nozzle with clean dry tissue after use.

**Other Information**

- Single patient use only.
- Discard 2 months after opening.
- Store below 25°C (77°F)

**Inactive Ingredients**

Water, honey (Leptospermum spp), sodium gluconate, glycerine. Preservative sodium benzoate.

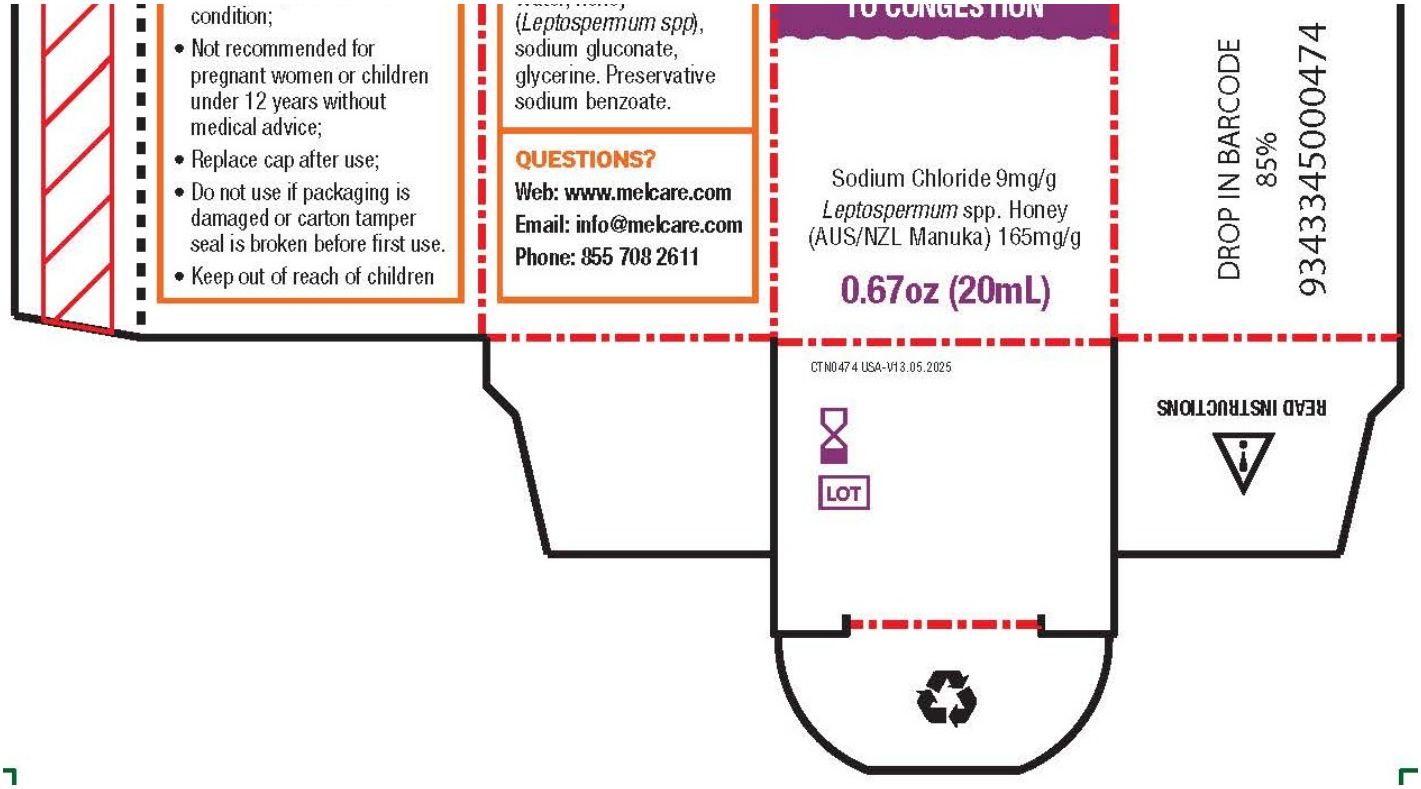
**Questions?**

**Web: www.melcare.com**

**Email: info@melcare.com**

**Phone: 855 708 2611**

**PRINCIPAL DISPLAY PANEL**




# RHINOMEL MANUKA

sodium chloride spray

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:73024-474 |
| **Route of Administration** | NASAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698) | SODIUM CHLORIDE | 9 mg in 1 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **SODIUM BENZOATE** (UNII: OJ245FE5EU) | |
| **HONEY** (UNII: Y9H1V576FH) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SODIUM GLUCONATE** (UNII: R6Q3791S76) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |

| Packaging | | | | |
|---|---|---|---|---|
| **#** | **Item Code** | **Package Description** | **Marketing Start Date** | **Marketing End Date** |
| **1** | NDC:73024-474-10 | 1 in 1 CARTON | 05/21/2025 | |

| | | | | |
|---|---|---|---|---|
| **1** | | 20 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| unapproved drug other | | 05/21/2025 | |

## Labeler - Melcare Biomedical Pty Ltd (743166886)

Revised: 5/2025 Melcare Biomedical Pty Ltd