

OPTIMEL MANUKA DRY EYE DROPS- glycerin solution/ drops
Melcare Biomedical Pty Ltd

Optimel Manuka+ Dry Eye Drops

Drug Facts

Active ingredients

Glycerin 0.2%

Purpose

Lubricant

Uses

- for the temporary relief of burning and irritation due to dryness of the eye.
- for use as a protectant against further irritation or to relieve dryness of the eye.

Warnings

For external use only

Do not use

- if you have a history of hypersensitivity or sensitive to honey or benzoate preservatives without seeking medical advice.
- if solution changes color or becomes cloudy.
- if packaging is damaged.
- serious conditions should be managed under the supervision of a doctor.

When using this product

- Before first use: check tamper ring is attached to cap base – remove cap and discard ring.
- to avoid contamination, do not touch tip of container to any surface.
- replace cap after using.
- remove contact lenses before using.
- wait at least 10 minutes before contact lens insertion.
- Initial stinging and redness on drop application is to be expected.

Stop use and ask a doctor if

- You experience eye pain change in vision, continued redness or irritation of the eye.
- Condition worsens

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wash hands.
- Instill 1 drop to the inner corner of the affected of the affected eye up to 3 times daily.
- Children under 16 years of age ask a doctor.

Other information

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

Inactive ingredients

Honey (*Leptospermum* spp.), Sodium benzoate, Sodium chloride, Water.

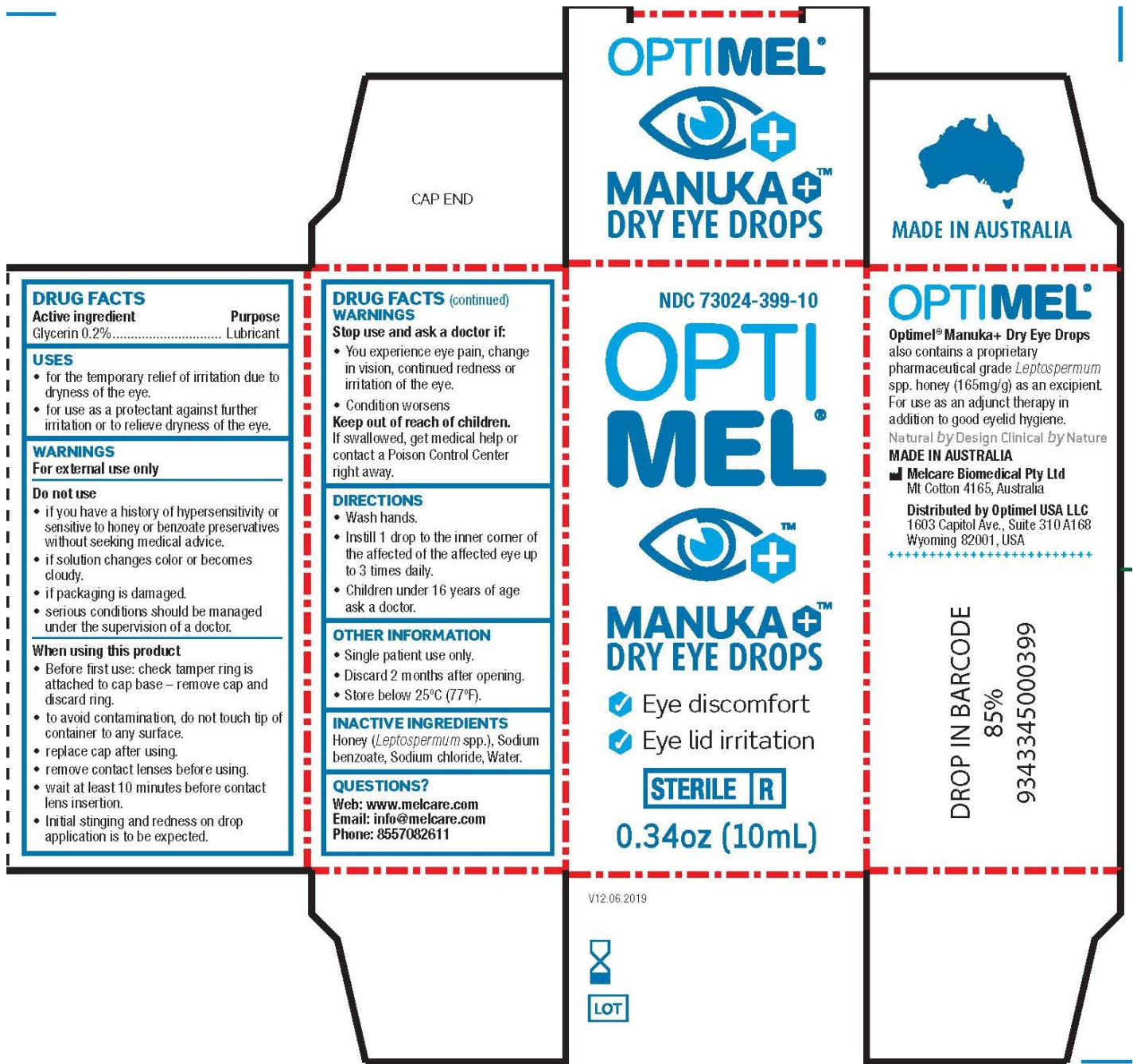
Questions?

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PRINCIPAL DISPLAY PANEL



OPTIMEL MANUKA DRY EYE DROPS

glycerin solution/ drops

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73024-399
Route of Administration	OPHTHALMIC		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GLYCERIN (UNII: PDC6A3C0OX) (GLYCERIN - UNII:PDC6A3C0OX)	GLYCERIN	2 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
HONEY (UNII: Y9H1V576FH)	
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73024-399-10	1 in 1 CARTON	07/19/2019	
1		10 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M014	07/19/2019	

Labeler - Melcare Biomedical Pty Ltd (743166886)

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