MEDSAFE
New Zealand Medicines and Medical Devices Safety Authority

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Good Manufacturing Practice Certificate

To whom it may concern

TT60-631-16-3

This is to certify that the **Cawthron Institute** operating at **98 Halifax St. East, Nelson, New Zealand,** has been audited to the *New Zealand Code of Good Manufacturing Practice for Manufacture and Distribution of Therapeutic Goods, Part 1: Manufacture of Pharmaceutical Products*, and has been found to comply with the requirements with respect to the following activities:

1. The microbiological, viscosity and chemical laboratory assessment of raw materials, including purified water, in-process and finished products for:

- Nutritional or 'nutraceutical' (vitamin, mineral & herbal) products where minor therapeutic claims are made, including products intended for the Australian market where they are termed 'listable'.
- Related products intended for the New Zealand market, as defined in the Medicines Act 1981.

2. The analytical instrumental techniques as applied to 1:

- Spectrometry (UV/Visible & FTIR)
- Gas chromatography
- High performance liquid chromatography & UPLC
- Thin layer chromatography & HPTLC
- ICP-MS

3. The storage of stability samples under ICH conditions

All testing performed according to pharmacopoeial requirements, client-supplied methods, in-house methods or methods specified in published standards, as required by the appropriate Regulatory Authority.

This certification is based on an audit carried out by an officer of the Ministry of Health at the Company's site on **16 and 17 May 2023**.

This certificate is valid until **30 April 2026**.

Derek Fitzgerald
Manager, Compliance Management Branch
30 October 2024

MINISTRY OF HEALTH
133 MOLESWORTH STREET
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