



Australian Government

Department of Health, Disability and Ageing
Therapeutic Goods Administration

Mr Fergus O'Connell
Head of Quality
Eurofins BioPharma Product Testing Australia Pty Ltd
179 Magowar Road
GIRRAWEEEN NSW 2145

TGA Reference: E21-367678

Subject: Issue of GMP certificate MI-2026-LI-00518-1

Dear Mr O'Connell,

Please find enclosed the GMP certificate for your manufacturing premises as requested.
Please do not hesitate to contact the Manufacturing Quality Branch if you require any further information.

Yours sincerely,

Signed and authorised by

Scott Pearce
Assistant Director
Licensing, Certification and Engagement Section
Manufacturing Quality Branch

16 February 2026

Contact: GMP@health.gov.au, Phone: 1800 020 653



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Department of Health, Disability and Ageing
Therapeutic Goods Administration

Certificate of GMP Compliance of a Manufacturer

Certificate Number:

MI-2026-LI-00518-1

Issued to:

Eurofins BioPharma Product Testing Australia Pty Ltd
ABN: 63 114 804 572

Manufacturing Site Address:

179 Magowar Road
GIRRAWEEEN NSW 2145
AUSTRALIA

The Therapeutic Goods Administration, the Competent Authority of Australia, confirms that this manufacturer holds a licence with number **MI-2021-LI-08995-1** to manufacture therapeutic goods under Section 38 of the *Therapeutic Goods Act 1989* and is included in the national inspection program following Section 40(4)(b) of the *Therapeutic Goods Act 1989*.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 14 to 16 May 2024, it is considered that the manufacturer complies with the Good Manufacturing Practice requirements of the PIC/S Guide to Good Manufacturing Practice for Medicinal Products – 01 May 2021.

This certificate reflects the status of the manufacturing site at the time of the inspection noted above and should not be relied upon to reflect the compliance status after the expiry date. This certificate should also not be relied upon where the status of the licence to manufacture therapeutic goods is not current. Where required, the Therapeutic Goods Administration as the issuing authority should be consulted.

Issue Date: 16 February 2026

Expiry Date: 16 May 2027

This certificate remains valid only if re-inspections are conducted when scheduled by the Therapeutic Goods Administration.
The authenticity of this certificate may be verified with the Therapeutic Goods Administration as the issuing authority.



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Certificate of GMP Compliance of a Manufacturer

Certificate Number:

MI-2026-LI-00518-1

MANUFACTURING OPERATIONS

The manufacturer above is authorised under Section 38 of the *Therapeutic Goods Act 1989* to carry out the following steps in the manufacture of therapeutic goods at the manufacturing site address specified above.

Manufacturing Type	Sterility	Dosage Form	Product Category	Manufacturing Step
Testing Laboratory	Sterile & Non Sterile	All Dosage Forms	Not Applicable	Testing microbial
Testing Laboratory	Sterile & Non Sterile	All Dosage Forms	Not Applicable	Testing chemical and physical
Testing Laboratory	Sterile	All Dosage Forms	Not Applicable	Testing sterility
Testing Laboratory	Sterile & Non Sterile	All Dosage Forms	Not Applicable	Endotoxin Testing

In addition to the statutory conditions that apply to all licences granted under Section 38 of the *Therapeutic Goods Act 1989*, the following specific conditions have been imposed on the licence under Sections 40(1) and/or 40(2) of the *Therapeutic Goods Act 1989*:

The authorisation for chemical testing is restricted to the testing of Ethylene oxide residues by gas-liquid chromatography.

The authorisation for physical testing is restricted to the testing of sub-visible particles.

The licence also authorises the potency testing of antibiotics using bioassay, and disinfectant testing.

This certificate remains valid only if re-inspections are conducted when scheduled by the Therapeutic Goods Administration. The authenticity of this certificate may be verified with the Therapeutic Goods Administration as the issuing authority.