Australian Government
Department of Health, Disability and Ageing
Therapeutic Goods Administration

Dr Fiona Zhang
Assistant QA Manager
Homart Pharmaceuticals Pty Ltd
59 Kirby Street
RYDALMERE NSW 2116

TGA Reference: 2014/017137

Subject: Issue of GMP certificate MI-2026-LI-01579-1

Dear Dr Zhang,

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

05 March 2026

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

Australian Government

Department of Health, Disability and Ageing

Therapeutic Goods Administration

Certificate of GMP Compliance of a Manufacturer

Certificate Number:

MI-2026-LI-01579-1

Issued to:

Homart Pharmaceuticals Pty Ltd
ABN: 74 057 411 640

Manufacturing Site Address:

59 Kirby Street
RYDALMERE NSW 2116
AUSTRALIA

The Therapeutic Goods Administration, the Competent Authority of Australia, confirms that this manufacturer holds a licence with number **MI-2012-LI-04100-3** 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.*

This certificate is issued based on a Surveillance Inspection of GMP compliance. From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 11 to 12 August 2025, 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 February 2022.

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: **05 March 2026** **Expiry Date:** **12 August 2028**

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.

PO Box 100 Woden ACT 2606 ABN 40 939 406 804
Phone: 1800 020 653 Fax: 02 6203 1605 Email: info@tga.gov.au www.tga.gov.au

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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
Medicine manufacture	Non Sterile	Tablet; uncoated	Listed Therapeutic Good	Full Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	Powders and Granules	Listed Therapeutic Good	Full Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	Solid Unit Dosage Forms - Hard Capsules	Listed Therapeutic Good	Full Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	All Dosage Forms	Registered Therapeutic Good	Packaging, Labelling and Release for Supply
Medicine manufacture	Non Sterile	All Dosage Forms	Listed Therapeutic Good	Packaging, Labelling and Release for Supply
Medicine manufacture	Non Sterile	Tablet, chewable	Listed Therapeutic Good	Full Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	Capsule, soft	Listed Therapeutic Good	Full Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	Liquids Group	Listed Therapeutic Good	Full Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	Topical Sunscreen Forms	Listed Therapeutic Good	Full Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	Tablet, film coated	Listed Therapeutic Good	Full Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	All Dosage Forms	Listed Therapeutic Good	Testing Physical
Medicine manufacture	Non Sterile	Semi Solids	Listed Therapeutic Good	Full Product Manufacture - excluding Testing

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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 licence does not authorise the manufacture of medicines listed for export that include substances only permitted in medicines contained within schedules 2, 3, 4 & 8 of the Poisons Standard.

Testing is restricted to physical testing only.

The manufacture of registered therapeutic goods excludes therapeutic products to which any part of the Standard for the Uniform Scheduling of Medicines and Poisons (SUSMP) applies, except for those goods containing solely the active ingredient macrogol.

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