



Australian Government
Department of Health
 Therapeutic Goods Administration

Licence to Manufacture Therapeutic Goods – Part 1

Licence Number:

MI-2012-LI-00154-3

Granted to:

The Green Dispensary Pty Ltd t/a GD Pharma

ABN: 95 083 276 724

Manufacturing Site Address:

46a Beulah Road

NORWOOD SA 5067

The manufacturer above is hereby 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	Sterile	Eye Drops	Therapeutic Goods for Clinical Trials	Sterile Finished Product Manufacture - excluding Testing
Medicine manufacture	Sterile	Injections	Therapeutic Goods for Clinical Trials	Sterile Finished Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	Oral Liquid, solution	Therapeutic Goods for Clinical Trials	Sterile Finished Product Manufacture - excluding Testing
Medicine manufacture	Sterile	Eye Drops	Registered Therapeutic Good	Sterile Finished Product Manufacture - excluding Testing
Medicine manufacture	Sterile	Injections	Registered Therapeutic Good	Sterile Finished Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	Nasal Drops	Registered Therapeutic Good	Full Product Manufacture - excluding Testing

Signed:



Jenny Hantzinikolas, Delegate of the Secretary

This Licence is valid only if the security provisions (blue and grey curved dotted lines across the bottom half of each page) are visible.
 This Licence is the property of the Therapeutic Goods Administration and must be returned or destroyed upon demand.
 This Licence remains valid until otherwise suspended or revoked by the Therapeutic Goods Administration.
 The status of an Australian Licence may be viewed at <https://www.ebs.tga.gov.au/>



Australian Government
Department of Health
 Therapeutic Goods Administration

Licence to Manufacture Therapeutic Goods – Part 1

Licence Number:
 MI-2012-LI-00154-3

Medicine manufacture	Non Sterile	Nasal Drops	Therapeutic Goods for Clinical Trials	Full Product Manufacture - excluding Testing
Medicine manufacture	Non Sterile	Liquids	Registered Therapeutic Good	Full Product Manufacture - excluding Testing

This licence is subject to the requirements of the *Therapeutic Goods Act 1989*, and its Regulations.

Section 40(4) of the *Therapeutic Goods Act 1989* and Regulation 19, 20, 21 and 22 of the Therapeutic Goods Regulations 1990 impose various statutory conditions on all licences to manufacture therapeutic goods.

In addition to that, the specific conditions mentioned in Part 2 of this licence have been imposed under Section 40(1) or 40(2) of the *Therapeutic Goods Act 1989*.

Originally Granted: **19 March-2013**

Date Revised: **12 February 2015**

Signed:



Jenny Hantzinikolas, Delegate of the Secretary

This Licence is valid only if the security provisions (blue and grey curved dotted lines across the bottom half of each page) are visible.
 This Licence is the property of the Therapeutic Goods Administration and must be returned or destroyed upon demand.
 This Licence remains valid until otherwise suspended or revoked by the Therapeutic Goods Administration.
 The status of an Australian Licence may be viewed at <https://www.ebs.tga.gov.au/>



Australian Government

Department of Health
Therapeutic Goods Administration

Licence to Manufacture Therapeutic Goods – Part 2:

Schedule of Conditions

Licence Number:

MI-2012-LI-00154-3

Issued to:

The Green Dispensary Pty Ltd t/a GD Pharma

ABN: 95 083 276 724

Manufacturing Site Address:

46a Beulah Road

NORWOOD SA 5067

In addition to the statutory conditions that have been imposed on all licences to manufacture therapeutic goods under Section 40(4) of the *Therapeutic Goods Act 1989* and Regulations 19, 20 and 21 of the Therapeutic Goods Regulations 1990, the conditions specified below have been imposed on this licence under Section/s 40(1) and/or 40(2) of the *Therapeutic Goods Act 1989*:

Sterile products are limited to terminally sterilised products only.

Quality Control testing is excluded from this licence.

The licence is limited to the manufacture of liquid dosage forms including nasal sprays. The licence excludes the manufacture of Penicillins and Cytotoxic drugs.

Persons currently nominated under Section 37(1)(e) of the Act as having control:

Production: Andrew Gia-Phong Vuong

Quality Control: Antony Condina

Originally imposed: **19 March 2013**

Date Revised: **12 February 2015**

Signed:

Jenny Hantzinikolas, Delegate of the Secretary

This Licence is valid only if the security provisions (blue and grey curved dotted lines across the bottom half of each page) are visible.
This Licence is the property of the Therapeutic Goods Administration and must be returned or destroyed upon demand.
This Licence remains valid until otherwise suspended or revoked by the Therapeutic Goods Administration.
The status of an Australian Licence may be viewed at <https://www.ebs.tga.gov.au/>