



Australian Government

Department of Health, Disability and Ageing
Therapeutic Goods Administration

Mr Geoff Acton
Managing Director
Antaria Pty Ltd
1821 Ipswich Road
ROCKLEA QLD 4106

TGA Reference: E19-522620

Subject: Reissue of licence to manufacture therapeutic goods MI-2019-LI-01711-1

Dear Mr Acton,

Please find enclosed the reissued Licence Part 1 (Authorisations) and Part 2 (Schedule of Conditions) 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

Maurice Makdessi
Delegate of the Secretary
Manufacturing Quality Branch

17 February 2026

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



Australian Government

Department of Health, Disability and Ageing
Therapeutic Goods Administration

Licence to Manufacture Therapeutic Goods – Part 1

Licence Number:

MI-2019-LI-01711-1

Granted to:

Antaria Pty Ltd
ABN: 36 092 404 727

Manufacturing Site Address:

1821 Ipswich Road
ROCKLEA QLD 4106

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
Active Pharmaceutical Ingredient manufacture	Non Sterile	API - Not Defined	Not Applicable	Testing chemical and physical
Sunscreen manufacture	Non Sterile	Topical Sunscreen Forms	Listed Therapeutic Good	Testing chemical and physical
Active Pharmaceutical Ingredient manufacture	Non Sterile	API - Not Defined	Not Applicable	Release for supply
Sunscreen manufacture	Non Sterile	Topical Sunscreen Forms	Listed Therapeutic Good	Release for supply

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: 7 February 2020

Date Revised: 17 February 2026

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, Disability and Ageing
Therapeutic Goods Administration

Licence to Manufacture Therapeutic Goods – Part 2: Schedule of Conditions

Licence Number:

MI-2019-LI-01711-1

Granted to:

Antaria Pty Ltd
ABN: 36 092 404 727

Manufacturing Site Address:

1821 Ipswich Road
ROCKLEA QLD 4106

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*:

Chemical and physical testing of non-sterile APIs is limited to zinc oxide and Proprietary Ingredient containing zinc oxide.

The release for supply of APIs is limited to non-sterile API and Proprietary Ingredient containing zinc oxide only.

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

Production: Charles Marchant

Quality Control: Dr. Vandana Chaudhary

Originally Granted: 7 February 2020

Date Revised: 16 February 2026

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