



Australian Government
Department of Health
Therapeutic Goods Administration

Certificate of GMP Compliance of a Manufacturer

Certificate Number:

MI-2013-CE-04797-1

Issued to:

Nature's Sunshine Products Inc.

Manufacturing Site Address:

1655 North Main Street
Spanish Fork Utah
UNITED STATES OF AMERICA

The Therapeutic Goods Administration, the Competent Authority of Australia, confirms that this manufacturer has been inspected following section/s 25(1)(g), 26(1)(g) and/or 26A(3) of the *Therapeutic Goods Act 1989* in connection with marketing authorisation/s listing manufacturers located outside Australia.

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 23 to 25 June 2014, 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 – 15 January 2009.

This certificate reflects the status of the manufacturing site at the time of the inspection noted above. This certificate remains valid until the expiry date provided that re-inspections are conducted as determined by the Therapeutic Goods Administration as the issuing Authority. This certificate should not be relied upon to reflect the compliance status after the expiry date.

EXPIRY DATE: 25 December 2017

ISSUE DATE: 09 September 2014

Name and signature of an authorised person of the Competent Authority of Australia:

Signed:

.....
Dr Dragana Milic, Delegate of the Secretary
Office of Manufacturing Quality

This certificate is valid only if the security provisions (blue and grey curved dotted lines on the bottom half of each page) are visible. 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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MANUFACTURING OPERATIONS

This certificate covers 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	Solid Unit Dosage Forms - Tablets	Registered Therapeutic Good	Finished Product Manufacture
Medicine manufacture	Non Sterile	Solid Unit Dosage Forms - Hard Capsules	Listed Therapeutic Good	Finished Product Manufacture
Medicine manufacture	Non Sterile	Liquids	Registered Therapeutic Good	Finished Product Manufacture
Medicine manufacture	Non Sterile	Powder, oral	Listed Therapeutic Good	Finished Product Manufacture
Medicine manufacture	Non Sterile	Herb, dried	Listed Therapeutic Good	Finished Product Manufacture
Medicine manufacture	Non Sterile	Capsule, soft	Listed Therapeutic Good	Packaging, Labelling and Release for Supply
Medicine manufacture	Non Sterile	Capsule, soft	Listed Therapeutic Good	Testing chemical and physical
Medicine manufacture	Non Sterile	Capsule, soft	Listed Therapeutic Good	Testing microbial

The following limitations are applicable to these manufacturing operations:

No further conditions are applicable.

Name and signature of an authorised person of the Competent Authority of Australia:

Signed:

Dragana Milic

Dr Dragana Milic, Delegate of the Secretary
 Office of Manufacturing Quality

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