



**Australian Government**

**Department of Health and Aged Care**  
Therapeutic Goods Administration

## **Certificate of GMP Compliance of a Manufacturer**

**Certificate Number:**

MI-2023-CE-11486-1

**Issued to:**

USPL Nutritionals LLC

**Manufacturing Site Address:**

22B Van Dyke Avenue  
New Brunswick New Jersey 08901  
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

From the knowledge gained during inspection of this manufacturer, the latest of which was conducted on 07 November 2023, 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. 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.

**Issue Date: 02 July 2024**

**Expiry Date: 07 November 2026**

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	Capsule, hard	Listed Therapeutic Good	Storage
Medicine manufacture	Non Sterile	Tablet, uncoated	Listed Therapeutic Good	Storage
Medicine manufacture	Non Sterile	Capsule, hard	Listed Therapeutic Good	Dispensing

The following limitations are applicable to these manufacturing operations:

No further limitations are applicable.

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.