THREE

March 10, 2023

Re: Three International: Manufacturing Philosophy and Certifications

Here at Three, we provide curated proactive wellness solutions using our proprietary Cellular Absorption Technology, proven to help you live a life of greater health and purpose.

The following pages contain the certifications of the contract manufacturers Three International partners with to produce its first-in-class products. These manufacturers are as follows: 1.) CSB Nutrition Corporation, 2.) Elevate Health Sciences, and 3.) United 1 Laboratories. These manufacturers have been producing nutritional supplements for decades and are held to the highest level of excellence. These powerful certifications attest that every Three product is manufactured to the highest quality standards to ensure they are pure, safe, and effective.

All ingredients in Three's products are source controlled to ensure the amounts of curated phytonutrients in the products are consistent every time. Every ingredient undergoes a battery of rigorous testing before it is deemed acceptable to use in the product.

Before a Three product is manufactured, it undergoes intense pilot testing to make sure the product formulated on the laboratory benchtop by our Ph.D. scientists is the same when made at metric ton scale. Thorough analytical analysis, content uniformity, and other techniques are used to verify they are identical in every detail.

During the manufacturing process, we never use fillers, binders, or excipients. At Three, we use the highest quality ingredients, backed by the best science, to make sure your body gets the nutrients it needs.

After the product is manufactured, a stringent Quality Analysis/Quality Control process is followed, along with third-party testing, before the product is released. Only then is it ready to be shipped to your home.

Thank you for joining us on this journey and for trusting us with your proactive wellness needs.

Be well,

Dr. Dan Gubler
Chief Scientific Officer
Three International

12180 S 300 E, Unit 1590
Draper, UT 84020

eurofins | Food Assurance

Eurofins Food Assurance
2120 Rittenhouse Street, Suite A
Des Moines, IA 50321, USA
Ph: (515) 299-6979
www.eurofinsus.com/assurance/food

DATES OF AUDIT:
01/20/2025 – 01/21/2025

NEXT RE-CERTIFICATION DATE:
12/22/2025

DATE OF DECISION:
04/17/2025

EXPIRATION DATE:
03/07/2026

CERTIFICATE NUMBER:
61774

CERTIFICATION TYPE:
Unannounced Recertification

Certificate of Registration

This acknowledges that

CSB Nutrition
2600 N. Main St.
Spanish Fork, UT 84601

is registered as meeting the requirements for the

SQF Food Safety Code for Dietary Supplements Manufacturing, Edition 9

Registration schedule

Scope of registration [food sector categories and products]:

Food sector category: FSC 31: Dietary Supplements Manufacturing

Products: Dietary Supplements

Signature of issuing officer
Brian Neal

SQF INSTITUTE
One world. One standard.
SQF Institute is a division of FMI.

ANAB
ANSI National Accreditation Board
ACCREDITED
ISO/IEC 17065
PRODUCT CERTIFICATION BODY
9166

CSB Nutrition Corporation

Certificate of Manufacturer

Product: Imune
Product: Purifl (30)

This document is to declare that *Purifl (30)* and *Imune* are exclusively manufactured for iii International at CSB Nutrition Corporation, an independent food and dietary supplement manufacturer, located in Spanish Fork, Utah, USA.

The methods used in the facilities, and the controls used for the design, manufacture, process, packaging, labeling, testing, and holding at CSB Nutrition Corporation, as a Food Manufacturer, adhere to the Current Good Manufacturing Practices and Quality System regulations as defined in 21 CFR parts 110 and 111, and meet these regulatory requirements.

Signed,

3-3-23

Amanda Huffman — Date
Document Control
CSB Nutrition Corporation

2600 North Main Street
Spanish Fork, UT 84660
Tel: 801-804-7552 Fax: 801-798-7944

State of Utah
SPENCER J. COX
Governor
DEIDRE M. HENDERSON
Lieutenant Governor

Department of Agriculture and Food
Craig W. Buttars
Commissioner
Kelly Pehrson
Deputy Commissioner
Travis Waller
Director, Regulatory Services

Certificate No.: REG-2023-14086

GOOD MANUFACTURING PRACTICE CERTIFICATE

We hereby certify that ELEVATE HEALTH SCIENCES, located at, 3421 SIERRA VISTA WAY, PROVO, UT 84606 is currently under inspection as a manufacturer of health food or dietary supplements. ELEVATE HEALTH SCIENCES has all the facilities to comply with the GOOD MANUFACTURING PRACTICE for food and dietary supplements (Code of Good Manufacturing Practice for food).

We also certify that ELEVATE HEALTH SCIENCES, is an inspected facility and the manufacturing plant in which their products are produced are subject to inspections at suitable intervals.

Inspection evaluates and assures compliance with the Utah Wholesome Food Act and Utah Food Protection Rule, which identifies the standard for proper facility construction, good manufacturing practices for food and dietary supplements (GMP), and fulfills requirements of those applicable laws and rules in the State of Utah.

UTAH DEPARTMENT OF AGRICULTURE AND FOOD

Travis Waller

Division of Regulatory Services

State of Utah, County of Salt Lake.

On this date FEB 01 2023 before me, the notary, personally appeared Travis Waller, proved on the basis of satisfactory evidence to be person, whose name is subscribed to this document, and acknowledge that he/she executed the same.

Notary Public

ALEXANDRA MATHEWS
Notary Public - State of Utah
Comm. No. 708180
My Commission Expires on
Oct 1, 2023

ELEVATE
HEALTH SCIENCES

3421 Sierra Vista Way
Provo, UT 84606
801-292-1217
www.elevatehs.com

CERTIFICATE OF MANUFACTURE

This certificate confirms that the product(s) listed below was manufactured, and tested by Elevate Health Sciences, USA, in accordance with the formula and specification provided and authorized by iii International.

Product: 3I Vitalite Capsule

Product: 3I OmeGo Softgel

Product: 3I Revive Softgel

All associated manufacturing, and testing documents are reviewed and released when found satisfactory. This product is manufactured in compliance with current good manufacturing practices and internal standard operating procedures.

Kristen Mitchell — 03/03/2023

Quality Systems Manager — Date

Australian Government
Department of Health and Aged Care
Therapeutic Goods Administration

Mr Nic Bryan
Vice President of Quality
Elevate Health Sciences
3421 Sierra Vista Way
Provo Utah 84606
United States of America

TGA Reference: E18-368931

Subject: Issue of GMP certificate MI-2019-CE-11110-1

Dear Mr Bryan,

Please find enclosed the GMP certificate for your manufacturing premises.

The certificate remains valid only if re-inspections are conducted when scheduled by the Therapeutic Goods Administration. The inspection frequency is not a reflection of the expiry date shown on the certificate but is consistent with the re-inspection frequency applicable to Australian manufacturers of the same class of products.

The Therapeutic Goods Administration will contact the relevant sponsor/s to arrange the re-inspection of your facility.

Yours sincerely,

Signed and authorised by

Matt Davis
Senior GMP Inspector
Manufacturing Quality Branch

17 November 2022

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

Australian Government

Department of Health and Aged Care
Therapeutic Goods Administration

Certificate of GMP Compliance of a Manufacturer

Certificate Number:

MI-2019-CE-11110-1

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, soft	Listed Therapeutic Good	Full Product Manufacture - excluding Chemistry
Medicine manufacture	Non Sterile	Capsule, hard	Listed Therapeutic Good	Full Product Manufacture - excluding Chemistry

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.

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

2023

CERTIFICATE OF REGISTRATION

This certifies that:

United Laboratories Manufacturing, LLC
1541 Champion Dr
Carrollton, TX 75006-6814
United States

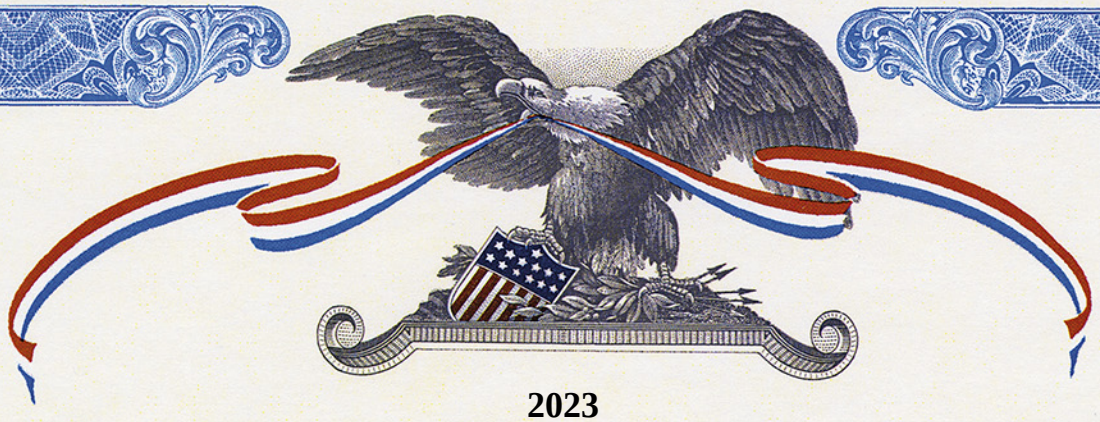
is registered with the U.S. Food and Drug Administration pursuant to the Federal Food Drug and Cosmetic Act, as amended by the Bioterrorism Act of 2002 and the FDA Food Safety Modernization Act, such registration having been verified as currently effective on the date hereof by Registrar Corp:

U.S. FDA Registration No.:	**18261284888**
U.S. FDA UFI (DUNS) No.:	**807878116**
U.S. Registration Agent:	**Registrar Corp** 144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

This certificate affirms that the above stated facility is registered with the U.S. Food and Drug Administration pursuant to the Federal Food Drug and Cosmetic Act, as amended by the Bioterrorism Act of 2002 and the FDA Food Safety Modernization Act, such registration having been verified as effective by Registrar Corp as of the date hereof, and Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate until December 31, 2023, unless such registration has been terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Registrar Corp assumes no liability to any person or entity in connection with the foregoing. The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

Registrar Corp
144 Research Drive, Hampton, Virginia, 23666, USA
Telephone: +1-757-224-0177 • Fax: +1-757-224-0179
info@registrarcorp.com • www.registrarcorp.com

David Lennarz
Executive Director
Registrar Corp
Dated: October 14, 2022



2023

CERTIFICATE OF REGISTRATION

This certifies that:

United Laboratories Manufacturing LLC
10685 King William Dr
Dallas, TX 75220-2412
United States

is registered with the U.S. Food and Drug Administration pursuant to the Federal Food Drug and Cosmetic Act, as amended by the Bioterrorism Act of 2002 and the FDA Food Safety Modernization Act, such registration having been verified as currently effective on the date hereof by Registrar Corp:

U.S. FDA Registration No.:	**15177704584**
U.S. FDA UFI (DUNS) No.:	**116910554**
U.S. Registration Agent:	**Registrar Corp** 144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

This certificate affirms that the above stated facility is registered with the U.S. Food and Drug Administration pursuant to the Federal Food Drug and Cosmetic Act, as amended by the Bioterrorism Act of 2002 and the FDA Food Safety Modernization Act, such registration having been verified as effective by Registrar Corp as of the date hereof, and Registrar Corp will confirm that such registration remains effective upon request and presentation of this certificate until December 31, 2023, unless such registration has been terminated after issuance of this certificate. Registrar Corp makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. Registrar Corp assumes no liability to any person or entity in connection with the foregoing. The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Registrar Corp is not affiliated with the U.S. Food and Drug Administration.

Registrar Corp
144 Research Drive, Hampton, Virginia, 23666, USA
Telephone: +1-757-224-0177 • Fax: +1-757-224-0179
info@registrarcorp.com • www.registrarcorp.com

David Lennarz
Executive Director
Registrar Corp
Dated: January 16, 2023

INW Dallas One Solutions

Certificate of Manufacture

This certifies that the products listed below will be manufactured by United Laboratories Manufacturing, LLC dba Dallas One Solutions, located at 1541 Champion Drive, Carrollton, Texas 75006, USA. These products will be produced exclusively for iii International, for their distribution and will be manufactured in accordance with the current United States Food and Drug Administration's (FDA) Good Manufacturing Practices, 21 CFR part 111, 211 and Dallas One Solutions' master formulations.

PRODUCT	FORMULA
iii International Collagene Gel 10 Pack	D-1188
iii International Eternel Gel 30 Pack	D-1189

Verified by: R. Pratibha _____ Date: 3/6/2023

Pratibha Ramanu, Quality Manager

Certificate of Conformity

Print Date

December 02, 2024

Certification Number

C0175333-HSCDS-4

Initial Certification

November 23, 2022

Expiration Date

December 01, 2025

NSF International has assessed and confirmed compliance of

CSB Nutrition Corporation

Facility:2600 North Main Street, Spanish Fork, UT, 84660, United States

Scope: NSF/ANSI 455-2 - 2021

which includes 21CFR Part 111, 21 CFR Part 117, 21 CFR Part 11, 21 CFR Part 1.5 Subpart L & 21 CFR Part 1.9 Subpart O

Product Technologies:

Dry Formulation, Encapsulation, Mixing, Packaging/Labeling Operation, Packaging/Labeling Operation - Bulk Packaging, Packaging/Labeling Operation - Primary Packaging, Packaging/Labeling Operation - Secondary Packaging, Quality Unit Operations, Warehousing

Product Categories:

Capsule, Oral, Dissolving Films, Powder, Soft Gel

Signed on behalf of
NSF International

David Trosin
Senior Director Global Certification,
Health Sciences

ANAB
ANSI National Accreditation Board
ACCREDITED
ISO/IEC 17065
PROCESS CERTIFICATION BODY

NSF International
789 N. Dixboro Road, Ann Arbor, MI 48105 USA

This certificate is the property of NSF International and must be returned upon request.
For the most current and complete information, please access NSF's website (nsf.org).

GMP CERTIFIED
NSF/ANSI 455-2
Dietary Supplements

Certificate of Conformity

Print Date

January 24, 2025

Certification Number

C0178332-HSCDS-10

Initial Certification

December 20, 2021

Expiration Date

January 17, 2026

NSF International has assessed and confirmed compliance of

UNITED LABORATORIES MANUFACTURING LLC DBA DALLAS ONE SOLUTIONS

Facility:1541 Champion Drive, Carrollton, TX, 75006, United States

Scope: NSF/ANSI 455-2 - 2021

which includes 21CFR Part 111, 21 CFR Part 117, 21 CFR Part 11, 21 CFR Part 1.5 Subpart L & 21 CFR Part 1.9 Subpart O

Product Technologies:

Liquid Formulation, Mixing, Packaging/Labeling Operation, Packaging/Labeling Operation - Dispensing, Packaging/Labeling Operation - Primary Packaging, Packaging/Labeling Operation - Secondary Packaging, Quality Unit Operations

Product Categories:

Ingestible Liquid

Signed on behalf of
NSF International

David Trosin
Senior Director Global Certification,
Health Sciences

ANAB
ANSI National Accreditation Board
ACCREDITED
ISO/IEC 17065
PROCESS CERTIFICATION BODY

NSF International
789 N. Dixboro Road, Ann Arbor, MI 48105 USA

GMP CERTIFIED
NSF/ANSI 455-2
Dietary Supplements

Certificate of Conformity

Print Date

January 20, 2025

Certification Number

C0178332-HSCDS-9

Initial Certification

December 20, 2021

Expiration Date

January 17, 2026

NSF International has assessed and confirmed compliance of

INW

Facility:1541 Champion Drive, Carrollton, TX, 75006, United States

Scope: NSF/ANSI 455-2 - 2021

which includes 21CFR Part 111, 21 CFR Part 117, 21 CFR Part 11, 21 CFR Part 1.5 Subpart L & 21 CFR Part 1.9 Subpart O

Product Technologies:

Liquid Formulation, Mixing, Packaging/Labeling Operation, Packaging/Labeling Operation - Dispensing, Packaging/Labeling Operation - Primary Packaging, Packaging/Labeling Operation - Secondary Packaging, Quality Unit Operations

Product Categories:

Ingestible Liquid

Signed on behalf of
NSF International

David Trosin
Senior Director Global Certification, Health Sciences

ANAB
ANSI National Accreditation Board
ACCREDITED
ISO/IEC 17065
PROCESS CERTIFICATION BODY

NSF International
789 N. Dixboro Road, Ann Arbor, MI 48105 USA

GMP CERTIFIED
NSF/ANSI 455-2
Dietary Supplements

Certificate of Conformity

Print Date

January 07, 2025

Certification Number

C0312779-HSCDS-3

Initial Certification

March 17, 2023

Expiration Date

January 02, 2026

NSF International has assessed and confirmed compliance of

Elevate Health Sciences

Facility:3421 Sierra Vista Way, Provo, UT, 84606, United States

Scope: NSF/ANSI 455-2 - 2021

which includes 21CFR Part 111, 21 CFR Part 117, 21 CFR Part 11, 21 CFR Part 1.5 Subpart L & 21 CFR Part 1.9 Subpart O

Product Technologies:

Dry Formulation, Encapsulation, Mixing, Packaging/Labeling Operation, Packaging/Labeling Operation - Bulk Packaging, Packaging/Labeling Operation - Dispensing, Packaging/Labeling Operation - Primary Packaging, Packaging/Labeling Operation - Secondary Packaging, Quality Unit Operations, Warehousing

Product Categories:

Capsule, Powder, Soft Gel

Signed on behalf of
NSF International

David Trosin
Senior Director Global Certification,
Health Sciences

ANAB
ANSI National Accreditation Board
ACCREDITED
ISO/IEC 17065
PROCESS CERTIFICATION BODY

NSF International
789 N. Dixboro Road, Ann Arbor, MI 48105 USA

GMP CERTIFIED
NSF/ANSI 455-2
Dietary Supplements

NSF INTERNATIONAL

789 N. Dixboro Road, Ann Arbor, Michigan 48105 USA
+1 800 673 6275

GMP for Sport™

NSF International has assessed and confirmed compliance of

CSB Nutrition Corporation

Facility: 2600 North Main Street, Spanish Fork, UT, 84660, United States

NSF GMP For Sport Program Requirements

Print Date:	December 02, 2024
Certificate Number:	C0175333-CS-7
Initial Certification:	February 06, 2014
Expiration Date:	December 01, 2025

David Trosin
Senior Director Global Certification,
Health Sciences