



PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

Perry Johnson Laboratory Accreditation, Inc. has assessed the Organization of:

Natural Vitamins Laboratory Corp.
4400 NW 133rd Street, Opa Locka, FL 33054

*and hereby declares that the Organization is accredited in accordance with
the recognized International Standard:*

ISO/IEC 17025:2017

Whereby, technical competence has been confirmed for the associated scope supplement, in the fields of:

Biological and Chemical Testing
(As detailed in the supplement)

Accreditation claims for conformity assessment activities shall only be made from the addresses referenced within this certificate and shall apply solely to those activities identified in the related scope. This Accreditation is granted subject to the Accreditation Body rules governing the Accreditation referred to above, and the Organization hereby commits to observing and complying with those rules in their entirety.

For PJLA:

Tracy Szerszen
President

Initial Accreditation Date:

March 05, 2021

Issue Date:

October 31, 2025

Expiration Date:

November 30, 2027

Accreditation No.:

111506

Certificate No.:

L25-801

*The validity of this certificate is maintained through ongoing assessments based
on a continuous accreditation cycle. The validity of this certificate should be
confirmed through the PJLA website: www.pjilabs.com*

Perry Johnson Laboratory
Accreditation, Inc. (PJLA)
755 W. Big Beaver, Suite 1325
Troy, Michigan 48084



Certificate of Accreditation: Supplement

Natural Vitamins Laboratory Corp.

4400 NW 133rd Street, Opa Locka, FL 33054

Contact Name: Ashish Patel Phone: 305-265-1660

Accreditation is granted to the facility to perform the following conformity assessment activities:

FIELD OF TEST	ITEMS, MATERIALS, OR PRODUCTS TESTED	COMPONENT, CHARACTERISTIC, PARAMETER TESTED	SPECIFICATION OR STANDARD METHOD	TECHNOLOGY OR TECHNIQUE USED	FLEX CODE	LOCATION OF ACTIVITY
Chemical	Raw Materials and Manufactured Products	Compound Identification	USP <43> / NF 38	HPLC-UV	F1, F2, F3	F
Chemical	Raw Materials and Manufactured Products	Compound Identification	USP <621>; NVL SOP #QC-037-01	GC-FID	F1, F2, F3, F4	F
Chemical	Raw Materials and Manufactured Products	Compound Identification	NVL SOP #QC-016-02	FT-IR	F1, F3, F4	F
Chemical	Raw Materials and Manufactured Products	Compound Identification	USP <43>/NF <38>	HPTLC	F1, F2, F3	F
Chemical	Raw Materials and Manufactured Products	Protein (as Nitrogen)	NVL SOP #STM-123	Kjeldahl Nitrogen	F1, F4	F
Chemical	Raw Materials and Manufactured Products	Arsenic Cadmium Lead Mercury	USP <43> / NF38	ICP/MS	F1, F2	F
Chemical	Manufactured Products	Stability/Shelf Life	NVL-SOP#QC-065-00	Stability Chamber	F1, F4	F
Biological	Raw Materials and Manufactured Products	Total Plate Count	USP <43> / NF38	BIOLUMIX	F1, F2	F
Biological	Raw Materials and Manufactured Products	Yeast and Mold	USP <43> / NF38	BIOLUMIX	F1, F2	F
Biological	Raw Materials and Manufactured Products	<i>E. coli</i>	USP <43> / NF38	BIOLUMIX	F1, F2	F
Biological	Raw Materials and Manufactured Products	<i>Salmonella</i>	USP <43> / NF38	BIOLUMIX	F1, F2	F



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Accreditation is granted to the facility to perform the following conformity assessment activities:

1. Location of activity:

Location

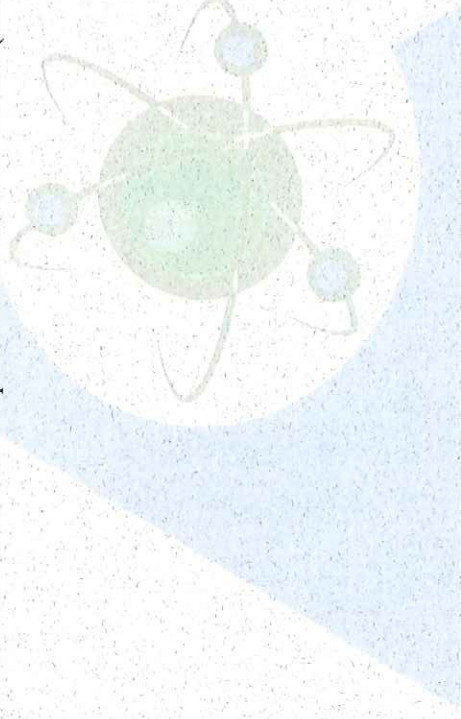
F

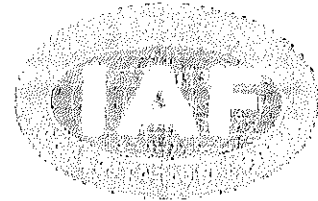
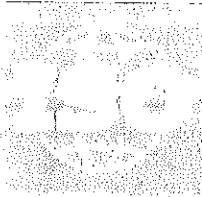
Location

Conformity assessment activity is performed at the CABs fixed facility

2. Flex Code:

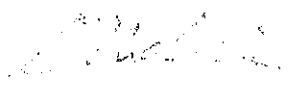
- F0- Fixed scope item. No deviations allowed to the line item as identified, except for updating to the most recent version of an accredited standard method after verification.
- F1- Laboratory has the capability to test a new item, material, matrix, or product similar in composition to item, material, matrix, or product identified on the scope
- F2- Laboratory has the capability to introduce the newest revision of an accredited authoritative standard method (with no modifications) identified on the scope
- F3- Laboratory has the capability to introduce a parameter/component/analyte to an accredited test method identified on the scope
- F4- Laboratory has the capability to introduce a new revision of an accredited non-standard method using the same technology or technique identified on the scope
- F5- Laboratory has the capability to introduce a validated method that is equivalent to an accredited method (using same technology or technique) identified on the scope



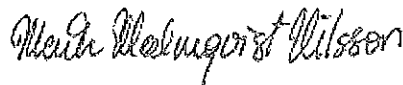


*Joint ISO-ILAC-IAF
Communique on the
Management Systems Requirements of ISO/IEC 17025,
General Requirements for the competence of testing and
calibration laboratories*

*A laboratory's fulfillment of the requirements of ISO/IEC 17025 means the laboratory meets both the technical competence requirements and **management system requirements** that are necessary for it to consistently deliver technically valid test results and calibrations. The **management system requirements** in ISO/IEC 17025 are written in language relevant to laboratory operations and operate generally in accordance with the principles of ISO 9001.*



ISO Acting Secretary General



ILAC Chair



IAF Chair

April 2017