



PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Alfa Laboratorio/ Martha Elidia Lujano Sarabia

***Montes Cantábricos # 19642, Fracc. Lomas Dorada
Tijuana, Baja California, México. CP. 22214***

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

ISO/IEC 17025:2017

This accreditation demonstrates technical competence for a defined scope and the
operation of a laboratory quality management system
(as outlined by the joint ISO-ILAC-IAF Communiqué dated April 2017):

Environmental and Biological Testing *(As detailed in the supplement)*

Accreditation claims for such testing and/or calibration services shall only be made from addresses referenced within this certificate. This Accreditation is granted subject to the system rules governing the Accreditation referred to above, and the Organization hereby covenants with the Accreditation body's duty to observe and comply with the said rules.

For PJLA:

Tracy Szerszen
President

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

Initial Accreditation Date:

April 09, 2014

Issue Date:

June 26, 2024

Expiration Date:

August 31, 2026

Accreditation No.:

71351

Certificate No.:

L24-483

*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*



Certificate of Accreditation: Supplement

Alfa Laboratorio / Martha Elidía Lujano Sarabia

Montes Cantábricos # 19642, Fracc. Lomas Dorada

Tijuana, Baja California, México. CP. 22214

Contact Name: Martha E. Lujano Sarabia Phone: 664-903-4190

Accreditation is granted to the facility to perform the following testing:

FLEX CODE	FIELD OF TEST	ITEMS, MATERIALS, OR PRODUCTS TESTED	COMPONENT, CHARACTERISTIC, PARAMETER TESTED	SPECIFICATION OR STANDARD METHOD	TECHNOLOGY OR TECHNIQUE USED
F1, F2	Biological ^F	Medical Devices	Bioburden	ISO 11737-1 Part 1: Determination of a Population or Microorganisms on Products	Membrane Filtration/Plate Count
F1, F2		Medical Devices and Water	Bacterial Endotoxin (<u>Limulus Amebocyte Lysate-LAL</u>)	USP <85 > Bacterial Endotoxin Test USP <161> Medical Device-Bacterial Endotoxin and Pyrogen, Test ANS I/ AAMI ST72/(R)	LAL Gel Clot
F1, F2		Food, Water, Surface and Air	Aerobic Bacteria	NOM 092-SSA1 Aerobic Plate Count in Foods AOAC Official Method 990.12	Plate Count/ Pour Plate Method
F1, F2			Yeast and Mold	NOM 111-SSA1 Yeast and Mold Count in Foods	Plate Count/ Pour Plate Method
F1, F2		Food, Water	Coliforms Bacteria	NOM 210-SSA1 (APENDICE H) Estimating Total Density, Fecal, Coliforms and E. coli	MPN
F1, F2				NOM 113-SSA1 AOAC Official Method 991.14 Coliform and Escherichia Coli Counts in Food	Plate Count/ Pour Plate Method
F1, F2	Biological ^O	Clean Room and Controlled Environments Microbial	Viable Particle Count	ISO 14698-1 and 2, Clean Rooms and Associated Controlled Environments - Bio Contamination Control	Air Sampler/ Plate Count
F1, F2	Environmental ^O	Clean room and Controlled Environments Particle	Particle Count in Air	ISO 14644 1, 2 and 3, Cleanrooms and Associated Controlled Environments	Particle Counter
F1, F2		Clean room and Controlled Environments Air Flow	Air Velocity	ISO 14644 1, 2 and 3, Clean Rooms and Associated Controlled Environments	Balometer
F1, F2		Clean Room and Controlled Environments Temperature and Humidity	Temperature and Humidity	ISO 14644 1, 2 and 3, Cleanrooms and Associated Controlled Environments	Particle Counter



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Accreditation is granted to the facility to perform the following testing:

1. The presence of a superscript F means that the laboratory performs testing of the indicated parameter at its fixed location.
2. The presence of a superscript O means that the laboratory performs testing of the indicated parameter onsite at customer locations.
3. Flex Code:
 - F1-Introduction of the testing of a new item, material, matrix, or product for an accredited test method
 - F2-Introduction of a new version of an accredited standard method (with no modifications)
 - F3-Introduction of a new parameter/component/analyte to an accredited test method
 - F4- Introduction of a new version or modifications of an accredited non-standard method
 - F5-Introduction of a new method that is equivalent to an accredited method (using same technology or technique)

