PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Alfa Laboratorio / Martha Elidia Lujano Sarabia

Montes Cantábricos # 19642, Fracc. Lomas Dorada
Tijuana, Baja California, México C.P. 22214

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 Environmental Analysis 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

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

Initial Accreditation Date:	*Issue Date:*	*Expiration Date:*
April 09, 2014	June 26, 2026	August 31, 2028

Accreditation No.:	*Certificate No.:*
71351	L26-609

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.pjlabs.com

Certificate of Accreditation: Supplement

Alfa Laboratorio / Martha Elidia Lujano Sarabia

Montes Cantábricos # 19642, Fracc. Lomas Dorada
Tijuana, Baja California, México C.P. 22214
Contact Name: Martha Lujano. Phone: 664-903-4190

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
Biological	Medical Devices	Bioburden	ISO 11737-1	Membrane Filtration/Plate Count	F1, F2	F
Biological	Medical Devices and Water	Bacterial Endotoxin (Limulus Amebocyte Lysate-LAL)	USP <85 > USP <161>	LAL Gel Clot	F1, F2	F
Biological	Food, Water, Surface and Air	Aerobic Bacteria	NOM 092-SSA1	Plate Count/ Pour Plate Method	F1, F2	F
Biological	Food, Water, Surface and Air	Yeast and Mold	NOM 111-SSA1	Plate Count/ Pour Plate Method	F1, F2	F
Biological	Food, Water	Total coliforms, Fecal coliforms and *E. coli*	NOM 210-SSA1 (APPENDIX H)	MPN	F1, F2	F
Biological	Food, Water	Total Coliforms	NOM 113-SSA1	Plate Count/ Pour Plate Method	F1, F2	F
Biological	Clean Room and Controlled Environments Microbial	Viable Particle Count	ISO 14698-1 and 2	Air Sampler/ Plate Count	F1, F2	F, O
Environmental Analysis	Clean Room and Controlled Environments	Particle Count in Air	ISO 14644-1, 2 and 3	Particle Counter	F1, F2	O
Environmental Analysis	Clean Room and Controlled Environments	Air Velocity	ISO 14644-1, 2 and 3	Balometer	F1, F2	O
Environmental Analysis	Clean Room and Controlled Environments	Temperature and Humidity	ISO 14644-1, 2 and 3	Particle Counter	F1, F2	O

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

1. Location of activity:

Location Code	Location
F	Conformity assessment activity is performed at the CAB's fixed facility
O	Conformity assessment activity is performed onsite at the CAB's customer location

2. Flex Code:

F0: When no flexibility is identified. There are no changes to items tested, characteristics identified or versions of methods except for updating to the most recent version of a standard method after verification.
F1: The 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: The laboratory has the capability to introduce the newest revision of an accredited authoritative standard method (with no modifications) identified on the scope
F3: The laboratory has the capability to introduce a parameter/component/analyte to an accredited test method identified on the scope
F4: The 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: The 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 for the same parameter, component, or analyte identified on the line item of the scope.