PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Laboratorio Valgio, S.A. de C.V.

Marco Tulio # 3204, Col. Camino Real
Guadalupe, Nuevo Leon, México. C.P. 67170

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 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.

Initial Accreditation Date:	*Issue Date:*	*Expiration Date:*
June 19, 2019	July 22, 2025	August 31, 2027

Accreditation No.:	*Certificate No.:*
103751	L25-571

For PJLA:

Tracy Szerszen
President

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

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

Laboratorio Valgio, S.A. de C.V.

Marco Tulio # 3204, Col. Camino Real
Guadalupe, Nuevo Leon, México. C.P. 67170
Contact Name: Ulises Valdes Barajas Phone: 818-647-3694

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	Food, Water, Surfaces and Environmental	Aerobic Platter Count	NOM-092-SSA1 NOM-110-SSA1 AOAC 2015.13	Petrifilm Direct Method, Balance Count Colony	F1, F2	F
Biological	Food, Water, Surfaces and Environmental	Total and Fecal Coliform and Escherichia Coli Counts (NMP/100 mL)	NOM-113-SSA NOM-110-SSA1 AOAC 991.14 NOM-210-SSA1 Regulatory Appendix H	Petrifilm Direct Method, Balance Count Colony	F1, F2	F
Biological	Food, Environmental	Enumeration of Yeast and Mold	NOM-111-SSA-1 AOAC 2014.05	Petrifilm Direct Method, Balance Count Colony	F1, F2	F
Biological	Food, Water, Surfaces and Environmental	*Salmonella spp.*	AOAC 2014.01	Petrifilm	F1, F2	F
Biological	Environment and Surface	Listeria spp.	AOAC RI #030601	Petrifilm	F1, F2	F
Biological	Food	Staphylococcus aureus	AOAC 2003.07	Petrifilm	F1, F2	F
Biological	Food	Listeria monocytogenes	AOAC PTM #060801	Petrifilm	F1, F2	F
Biological	Fecal Stool	Entamoeba Coli Giardia Lamblia Entamoeba Histolytic	Manual Parasitology Copro Parasitoscopic Microscope	Frame of Reference	F1, F2	F
Biological	Fecal Stool	*Salmonella spp.*	CHROMagar Salmonella AOAC RI 020502	Plate Culture and Visual Determination by Color Turn	F1, F2	F

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Biological	Pharyngeal Exudate	Streptococcus Beta Hemolytic Group A	Technical Guide (Medical Magazine Clinical Microbiology) Blood Agar Chocolate Agar (CHOC)	Plate Culture and Observational Determination, Characteristic Hemolytic Colonies	F1, F2	F
Biological	Food/ Water	Vibrio cholerae and Vibrio parahemoliticus	ISO 21872-1	Plate Culture and Visual Determination by Color turn	F1, F2	F
Biological	Blood	Febrile Reactions *Salmonella* Typhi o Antigen, *Salmonella* Typhi H Antigen, *Salmonella* Paratyphi A and Paratyphi B Antigen, Proteus OX-19 Brucella Abortus	Manufacturer Procedure (Licon)	Direct Method Plate Agglutination Method, with Licon Reagents. Visualization and Confirmation by Optical microscopy	F1, F2	F
Biological	Blood	Determination VDRL	Manufacturer Procedure (Licon)	Plate Agglutination Method, with Licon Reagents. Visualization and confirmation by Optical Microscopy	F1, F2	F

1. Location of activity:

Location Code	Location
F	Conformity assessment activity is performed at the CAB's fixed facility

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

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.