PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Paragon Laboratories, Inc.

12649 Richfield Court, Livonia, MI 48150

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.

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:*
January 20, 2025	January 20, 2025	Febraury 28, 2027
Revision Date:	*Accreditation No.:*	*Certificate No.:*
November 24, 2025	78772	L25-50-R1

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

Paragon Laboratories, Inc.

12649 Richfield Court, Livonia, MI 48150
Contact Name: Paul Kijek Phone: 734-469-5626

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 Products/Swab/Sponge Testing	Salmonella genus	AOAC 2020.02	PCR - GENE-UP	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Listeria species	AOAC 2019.10	PCR - GENE-UP	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Listeria monocytogenes	AOAC 2019.11	PCR - GENE-UP	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	E. coli O157:H7	AOAC RI 031002	PCR - GENE-UP	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Enterohemorrhagic E. coli (EHEC)	AOAC RI 121806	PCR - GENE-UP	F1, F2	F
Biological	Water Testing	Legionella (Potable Water)	Legiolert Test Kit Instructions ASTM D8429	IDEXX Quanti-tray	F1, F2	F
Biological	Water Testing	Legionella (Non-Potable Water)	Legiolert Test Kit Instructions ASTM D8429	IDEXX Quanti-tray	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Total Aerobic Bacteria Count	AOAC 121204	TEMPO	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Lactic Acid Bacteria (LAB)	TEMPO 80071 Test Kit Instructions	TEMPO	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Total Yeast & Mold	AOAC RI 041001	TEMPO	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Total Coliforms	AOAC RI 060702	TEMPO	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Escherichia coli	AOAC 2009.02	TEMPO	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Enterobacteriaceae	AOAC 050801	TEMPO	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Staphylococcus aureus	AOAC RI 120901	TEMPO	F1, F2	F
Biological	Food Products/Swab/Sponge Testing	Salmonella species	AOAC 2013.01	VIDAS	F1, F2	F

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Biological	Food Products/Swab/Sponge Testing	Listeria species	AOAC 2013.10	VIDAS	F1, F2	F

1. Location of activity:

Location	Location
F	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