PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Analytical Testing Laboratories, Inc

345 Trapelo Road, Belmont, MA 02478

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:*
July 19, 2014	October 16, 2026	February 28, 2029

Accreditation No.:	*Certificate No.:*
78387	L26-924

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

Analytical Testing Laboratories, Inc

345 Trapelo Road, Belmont, MA 02478
Contact Name: Ashley Hurst Phone: 617-484-7400

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/Beverage Environmental	Gluten	AOAC OMA 991.19 AOAC PTM 061201	Neogen Veratox® for Gliadin Kit	F1, F2	F
Biological	Food/Beverage Environmental	Aerobic Plate Count	AOAC OMA 990.12 AOAC OMA 2015.13 CMMEF Ch. 8	Neogen Petrifilms Neogen Agar Petridishes	F1, F2	F
Biological	Food/Beverage Environmental	*Bacillus cereus*	CMMEF Ch. 31	Biomerieux Petridishes	F1, F2	F
Biological	Food/Beverage Environmental	*Enterobacteriaceae*	AOAC OMA 2003.01	Neogen Petrifilms	F1, F2	F
Biological	Food/Beverage Environmental	E. Coli, Coliforms	AOAC OMA 991.14 AOAC OMA 996.02 CMMEF Ch. 9	Neogen Petrifilms Neogen Agar Petridishes Neogen Broth	F1, F2	F
Biological	Food/Beverage Environmental	Lactic Acid Bacteria	AOAC PTM 041701 CMMEF Ch. 19	Neogen Petrifilms	F1, F2	F
Biological	Food/Beverage Environmental	*Staphylococcus aureus*	AOAC OMA 2003.08 CMMEF Ch. 39	Neogen Petrifilms	F1, F2	F
Biological	Food/Beverage Environmental	Yeast and Mold	AOAC OMA 997.02 AOAC OMA 2014.05 CMMEF Ch. 21	Neogen Petrifilms Neogen Agar Petridishes	F1, F2	F
Biological	Food/Beverage Environmental	*Listeria monocytogenes*	AOAC 2003.12 AOAC 2013.11 CMMEF Ch. 35	Biomerieux Vidas® Neogen Agar Petridishes Biomerieux Agar Petridishes	F1, F2	F
Biological	Food/Beverage Environmental	*Listeria* Spp.	AOAC 2013.10 CMMEF Ch. 35	Biomerieux Vidas®	F1, F2	F
Biological	Food/Beverage Environmental	*Salmonella* Spp.	AOAC 2013.01 AOAC 2013.02 CMMEF Ch. 36	Biomerieux Vidas® Biomerieux Agar Petridishes	F1, F2	F
Biological	Food/Beverage Environmental	*E. coli* O157:H7	AOAC 996.09 CMMEF Ch. 34	Millipore VIP® Gold Lateral Flow	F1, F2	F

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Biological	Water/Waste Water	Total Coliforms	SM 9222 A SM 9222 B SM 9222 D	Neogen Agar Petridishes Millipore Agar Petridishes Membrane Filtration	F1, F2	F
Biological	Water/Waste Water	Heterotrophic Plate Count	SM-9215 A	Neogen Petrifilms Neogen Agar Petridishes	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