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	January 21, 2025	February 28, 2027

Accreditation No.:	*Certificate No.:*
78387	L25-49

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 991.19 AOAC-RI 061201	Neogen Veratox® for Gliadin Kit	F1, F2	F
		Aerobic Plate Count	AOAC 990.12 AOAC 2015.13 CMMEF Ch. 8	Neogen Petrifilms Neogen Agar Petridishes	F1, F2	
		Bacillus cereus	CMMEF Ch. 31	Biomerieux Petridishes	F1, F2	
		Enterobacteriaceae	AOAC 2003.01	Neogen Petrifilms	F1, F2	
		Coliforms	AOAC 991.14 AOAC 996.02 AOAC 2000.15 CMMEF Ch. 9	Neogen Petrifilms Neogen Agar Petridishes Neogen Broth	F1, F2	
		Lactic Acid Bacteria	AOAC RI 041701 CMMEF Ch. 19	Neogen Petrifilms	F1, F2	
		Staphylococcus aureus	AOAC 2003.08 CMMEF Ch. 39		F1, F2	
		Yeast and Mold	AOAC 997.02 AOAC 2014.05 CMMEF Ch. 21	Neogen Petrifilms Neogen Agar Petridishes	F1, F2	
		Listeria monocytogenes	AOAC 2003.12 AOAC 2013.11 AOAC RI PTM 070702 AOAC RI PTM 080901 AOAC RI PTM 121402 CMMEF Ch. 35	Biomerieux Vidas® Neogen Agar Petridishes	F1, F2	
		Listeria Spp.	AOAC 2013.10 AOAC RI 030502 AOAC RI 050903 AOAC RI 081401 CMMEF Ch. 35	Biomerieux Vidas®	F1, F2	

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Biological	Food/Beverage Environmental	*Salmonella* Spp.	AOAC 2003.09 AOAC 2011.03 AOAC 2013.01 AOAC 2013.02 CMMEF Ch. 36	Biomerieux Vidas® Biomerieux Agar Petridishes	F1, F2	F
		E. coli O157:H7	AOAC 996.09 AOAC RI 050501 AOAC RI 031002 AOAC RI 091301 CMMEF Ch. 34	Millipore VIP® Gold Lateral Flow	F1, F2	
	Water	Total Coliforms	SMEWW 9222 A, B, D	Neogen Agar Petridishes Millipore Agar Petridishes Membrane Filtration	F1, F2	F
		Heterotrophic Plate Count	SMEWW 9215 A	Neogen Petrifilms Neogen Agar Petridishes	F1, F2	

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