

PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Analytical Resource Laboratories

520 South 850 East, Suite B3, Lehi, UT 84043

(Hereinafter called the Organization) and hereby declares that Organization is accredited in accordance with the recognized International Standard:

ISO/IEC 17025:2017

This accreditation demonstrates technical competence for a defined scope and the operation of a laboratory quality management system (as outlined by the joint ISO-ILAC-IAF Communiqué dated April 2017):

Chemical and Microbiological Testing (As detailed in the supplement)

Accreditation claims for such testing and/or calibration services shall only be made from addresses referenced within this certificate. This Accreditation is granted subject to the system rules governing the Accreditation referred to above, and the Organization hereby covenants with the Accreditation body's duty to observe and comply with the said rules.

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:
February 18, 2015	July 17, 2023	August 31, 2025
Accreditation N	lo.: Certific	eate No.:
77504	L23-54	48-1

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: <u>www.pjlabs.com</u>



Certificate of Accreditation: Supplement

Analytical Resource Laboratories 520 South 850 East, Suite B3, Lehi, UT 84043

520 South 850 East, Suite B3, Lehi, UT 84043 Contact Name: Jacob Teller Phone: 801-331-8293

Accreditation is granted to the facility to perform the following testing:

FIELD OF TEST	ITEMS, MATERIALS OR PRODUCTS TESTED	SPECIFIC TESTS OR PROPERTIES MEASURED	SPECIFICATION, STANDARD METHOD OR TECHNIQUE USED	RANGE (WHERE APPROPRIATE) AND DETECTION LIMIT
Chemical ^F Nutraceuti	Nutraceutical,	Aluminum	ARL 8.016	D.L. = 0.007 mg/kg
	Cosmetics, and	Antimony		D.L. = 0.000 1 mg/kg
	Food	Arsenic	-	D.L. = 0.001 mg/kg
		Cadmium	-	D.L. = 0.000 1 mg/kg
		Chromium		D.L. = 0.002 mg/kg
		Lead		D.L. = 0.001 mg/kg
		Manganese		D.L. = 0.001 mg/kg
		Mercury		D.L. = 0.000 1 mg/kg
		Nickel		D.L. = 0.002 mg/kg
		Tin		D.L. = 0.001 mg/kg
		Zinc		D.L. = 0.001 mg/kg
		Ash	AOAC 942.05	D.L. = 0.01 mg
		Loss on Drying (gravimetric)	USP <731>	D.L. = 1 mg
		Gluten Allergens (as Gliadin)	AOAC 991.19	N/A
		Water Content (Karl Fischer)	ASTM 203-01	
		рН	AOAC 981.12-E	
	Cannabinoids	AOAC 2018.11 (modified)	D.L. = 3 mcg/mL	
	Caffeine	HPLC-DAD ARL 2.013	D.L. = 1 mcg/mL	
	Total, Soluble and Insoluble Dietary Fiber	AOAC 991.43 ARL 2.034	N/A	
		Mitragynine, Paynantheine, 7-hydroxymitragynine, and Speciogynine	HPLC-DAD ARL 2.046	D.L. = 3 mcg/mL
		FTIR	ARL 8.012, USP <1854>	Characterization test
Microbiological ^F	Nutraceutical, Cosmetics, and	Yeast and Mold Rapid Analysis (Enumeration)	AOAC 2014.05 (modified)	D.L = 10 CFU's/g
For Nu Co	Food	Staphylococcus aureus (Detection and Confirmation Testing)	Gene-Up, PCR ARL 1.030	Absence or Presence per 10 g
		<i>Listeria</i> spp. (Detection and Confirmation Testing)	Gene-Up, PCR ARL 1.030	Absence or Presence per 10 g
	Nutraceuticals, Cosmetics, and	Total Aerobic Plate Count (Detection)	USP <61>	D.L. = 10 CFU/g
	Drug Products	Total Combined Yeast and Mold (Detection)	USP <61>	
		Salmonella spp. (Detection)	USP <62>	Absence or Presence per 10 g
		Staphylococcus aureus (Detection)	USP <62>	

This supplement is in conjunction with certificate #L23-548-1



Certificate of Accreditation: Supplement

Analytical Resource Laboratories

520 South 850 East, Suite B3, Lehi, UT 84043 Contact Name: Jacob Teller Phone: 801-331-8293

Accreditation	is granted	l to the faci	litv to perforn	1 the following	testing:
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FIELD OF TEST	ITEMS, MATERIALS OR PRODUCTS TESTED	SPECIFIC TESTS OR PROPERTIES MEASURED	SPECIFICATION, STANDARD METHOD OR TECHNIQUE USED	RANGE (WHERE APPROPRIATE) AND DETECTION LIMIT
Microbiological ^F	Nutraceuticals,	Escherichia coli	USP <62>	Absence or Presence per 10 g
	Cosmetics, and	(Detection)		
	Drug Products	Bile Tolerant Gram-	USP <62>	Absence or Presence per 1 g
		Negative Bacteria		
		(Detection)		
	Nutraceutical,	Pathogenic E. coli	Gene-up, PCR	Absence or Presence per 10 g
	Cosmetics, and	(Detection and	ARL 1.030	
	Food	Confirmation Testing)		
		Escherichia coli	Gene-up, PCR	Absence or Presence per 10 g
		(Detection and	ARL 1.030	
		Confirmation Testing)		
		Listeria (Detection and	Gene-up, PCR	Absence or Presence per 25 g
		Confirmation Testing)	ARL 1.030	
		Salmonella (Detection and	Gene-Up, PCR	Absence or Presence per 10 g
		Confirmation Testing)	ARL 1.030	
		Total Aerobic Plate Count	USP <2021>	D.L. = 10 CFU/g
		(Enumeration)	(modified)	
		Yeast Count	USP <2021>	
		(Enumeration)	(modified)	
		Mold Count	USP <2021>	
		(Enumeration)	(modified)	
		Escherichia coli	USP <2022>	Absence or Presence per 10 g
		(Detection)	(modified)	
		Staphylococcus aureus	USP <2022>	
		(Detection)	(modified)	
		Salmonella spp.	USP <2022>	
		(Detection)	(modified)	
		Enterobacteriaceae	AOAC 2003.01	D.L. = 10 CFU/g
		(Enumeration)	(modified)	
		Coliforms (Enumeration)	AOAC 991.14	
			(modified)	

1. The presence of a superscript F means that the laboratory performs testing of the indicated parameter at its fixed location. Example: Outside Micrometer ^F would mean that the laboratory performs this testing at its fixed location.

2. This is the primary site for all quality management system activities.