PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Microchem Laboratory, LLC

1700 Chisholm Trail Road, Round Rock, TX 78681

and hereby declares that the Organization is accredited in accordance with the recognized International Standard:

ISO/IEC 17025:2017

& Meets the Requirements of Good Laboratory Practice for Nonclinical Laboratory Studies, Title 21 CFR Part 58 Accreditation Program

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:*
Febraury 26, 2025	July 17, 2025	October 31, 2027

Accreditation No.:	*Certificate No.:*
128395	L25-536-1

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: www.pjlabs.com

Certificate of Accreditation: Supplement

Microchem Laboratory, LLC

1700 Chisholm Trail Road, Round Rock, TX 78681
Contact Name: Dr. Benjamin Tanner Phone: 512-310-8378

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	Reusable Medical Devices	Validation of Cleaning Efficacy	ST98	Spectrophotometer	F1, F2	F
Biological	Reusable Medical Devices	Validation of Sterilization/Reprocessing	AAMI TIR 12	Plating	F1, F2	F
Biological	Cosmetics	Evaluation of Antimicrobial Protection of Cosmetic Products	ISO 11930	Plating	F1, F2	F
Biological	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Nutraceuticals, Dietary Supplements and Raw Materials for the above product types.	Aerobic Plate Count Enrichment	USP <61> USP <62> Quantification and Enrichment: MET-TFO-132	Plating	F1, F2	F
Biological	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Nutraceuticals, Dietary Supplements and Raw Materials for the above product types.	Determination of a population of microorganisms on products	ISO 11737-1	Plating	F1, F2	F
Biological	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Nutraceuticals, Dietary Supplements and Raw Materials for the above product types.	Sterility Testing	USP <71>	Plating, ISO Class 7 Cleanroom, ISO Class 5 Hoods, Incubators	F1, F2	F

Certificate of Accreditation: Supplement

Microchem Laboratory, LLC
1700 Chisholm Trail Road, Round Rock, TX 78681
Contact Name: Dr. Benjamin Tanner Phone: 512-310-8378

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

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

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