



PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

BioChroma Analytical Laboratory, LLC
1309 Record Crossing Road, Dallas, TX 75235

*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 and Chemical 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: **April 30, 2020**

Issue Date: **July 30, 2024**

Expiration Date: **July 31, 2026**

Revision Date: **October 21, 2025**

Accreditation No.: **105837**

Certificate No.: **L24-580-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.pilabs.com*



Certificate of Accreditation: Supplement

BioChroma Analytical Laboratory, LLC

1309 Record Crossing Road, Dallas, TX 75235

Contact Name: Benny McKee Phone: 972-454-9166

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	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Aerobic Plate Count	USP <61> USP <2021>	Standard Plate Count	F1, F2	F
Biological	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Yeast and Mold Count	USP <61> USP <2021>	Standard Plate Count	F1, F2	F
Biological	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Escherichia coli	USP <62> USP <2022>	Selective Media	F1, F2	F
Biological	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Staphylococcus aureus	USP <62> USP <2022>	Selective Media	F1, F2	F
Biological	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Salmonella sp.	USP <62> USP <2022>	Selective Media	F1, F2	F
Biological	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Enterobacteriaceae	USP <62> USP <2022>	Selective Media	F1, F2	F



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Biological	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Total Coliforms	USP <62> USP <2022>	Selective Media	F1, F2	F
Chemical	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Heavy Metals: Lead Mercury Arsenic Cadmium	BCAL-DV-AMP030	ICPMS	F1, F4	F
Chemical	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Minerals	BCAL-DV-AMP051	IC	F1, F4	F
Chemical	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Identification	USP <197>	FTIR	F1, F2	F
Chemical	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Concentration	USP <621>	HPLC	F1, F2	F
Chemical	Pharmaceutical Ingredients, OTC Drug Products, Pet Products, Personal Care Products, Cosmetics, Medical Devices, Dietary Supplements and Raw Materials for the above product types, Hemp and Hemp Products	Concentration	USP <621>	GC	F1, F2	F



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Chemical	Flower Distillate Creams Tinctures Oils Edibles	Cannabinoids: CBD $\Delta 9$ -THC CBC CBG CBN CBDV $\Delta 8$ -THC CBGA THCV THCA CBDA	BCAL-DV-AMP026	LCMS	F1, F4	F

1. Location of activity:

Location

F

Conformity assessment activity is performed at the CABs fixed facility

Location

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