PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Medipharm Laboratories, Inc.

6736 NW 72nd Avenue, Miami, FL 33166

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:

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:	*Issue Date:*	*Expiration Date:*
June 01, 2026	June 01, 2026	July 31, 2028

Accreditation No.:	*Certificate No.:*
122271	L26-477

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

Medipharm Laboratories, Inc.

6736 NW 72nd Avenue, Miami, FL 33166
Contact Name: Carlos Lopetegui Phone: 305-889-8301

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
Chemical	Raw Materials, Finished Pharmaceutical Products	pH	USP-NF Standards	ISE	F1, F4	F
Chemical	Raw Materials, Finished Pharmaceutical Products	HPLC Identifiable Compounds	USP-NF Standards	HPLC / UV	F1, F3, F4	F
Chemical	Raw Materials, Finished Pharmaceutical Products	FTIR Identifiable Compounds	USP-NF Standards	FTIR	F1, F3, F4	F
Chemical	Raw Materials, Finished Pharmaceutical Products	UV/VIS Spec Identifiable Compounds	USP-NF Standards	UV/VIS Spec	F1, F3, F4	F
Chemical	Raw Materials, Finished Pharmaceutical Products	GC-FID Identifiable Compounds	USP-NF Standards	Gas Chromatograph / FID	F1, F3, F4	F
Chemical	Raw Materials, Finished Pharmaceutical Products	Fluoride	USP-NF Standards	ISE	F1, F4	F
Chemical	Raw Materials, Finished Pharmaceutical Products	Product stability / Shelf life	USP-NF Standards	Stability Chamber	F1, F4	F
Chemical	Raw Materials, Finished Pharmaceutical Products	Dissolution	USP-NF Standards	Dissolution Apparatus	F1, F4	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