



PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Jordi Labs, LLC

200 Gilbert Street, Mansfield, MA 02048

*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:

Initial Accreditation Date:

Issue Date:

Expiration Date:

June 26, 2020

June 08, 2026

August 31, 2028

Accreditation No.:

Certificate No.:

96418

L26-491

Tracy Szerszen
President

*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.pjlab.com*

Perry Johnson Laboratory
Accreditation, Inc. (PJLA)
755 W. Big Beaver, Suite 1325
Troy, Michigan 48084



Certificate of Accreditation: Supplement

Jordi Labs, LLC

200 Gilbert Street, Mansfield, MA 02048
Contact Name: Tyler Stewart Phone: 508-966-1301

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	Food Contact Surfaces, Finished Medical Devices, Medical Device Components, Drugs and Packaging	Chemical Characterization, Extractables and Leachables Analysis	SOP P7.1.1.104 Based on: ISO 10993-12 BS/EN/ISO 10993-12 ISO 10993-18 BS/EN/ISO 10993-18	High-Performance Liquid Chromatography (HPLC/MS/UV/CAD)	F1, F2, F4	F
Chemical	Food Contact Surfaces, Finished Medical Devices, Medical Device Components, Drugs and Packaging	Chemical Characterization, Extractables and Leachables Analysis	SOP P7.1.1.271 Based on: ISO 10993-12 BS/EN/ISO 10993-12 ISO 10993-18 BS/EN/ISO 10993-18	Inductively Coupled Plasma Mass Spectrometry (ICPMS)	F1, F2, F4	F
Chemical	Food Contact Surfaces, Finished Medical Devices, Medical Device Components, Drugs and Packaging	Chemical Characterization, Extractables and Leachables Analysis	SOP P7.1.1.272 Based on: ISO 10993-12 BS/EN/ISO 10993-12 ISO 10993-18 BS/EN/ISO 10993-18	Gas Chromatography Mass Spectrometry (GCMS)	F1, F2, F4	F
Chemical	Food Contact Surfaces, Finished Medical Devices, Medical Device Components, Drugs and Packaging	Chemical Characterization, Extractables and Leachables Analysis	SOP P7.1.1.273 Based on: ISO 10993-12 BS/EN/ISO 10993-12 ISO 10993-18 BS/EN/ISO 10993-18	Head Space Gas Chromatography Mass Spectrometry (HS-GCMS)	F1, F2, F4	F
Chemical	Food Contact Surfaces, Finished Medical Devices, Medical Device Components, Drugs and Packaging	Mass of Nonvolatile Residue (Gravimetric Analysis), Extractables and Leachables Analysis	SOP P7.1.1.72 Based on: ISO 10993-18 BS/EN/ISO 10993-18	Gravimetric Analysis	F1, F2, F4	F



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

1. Location of activity:

Location

F

Location

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

