



PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

Life Science Outsourcing, Inc.

830 Challenger Street, Brea, CA 92821

*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, Environmental Simulation, Mechanical, and Non-Destructive
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:

July 07, 2026

Issue Date:

July 07, 2026

Expiration Date:

August 31, 2028

Accreditation No.:

120500

Certificate No.:

L26-623

*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.pjllabs.com*



Certificate of Accreditation: Supplement

Life Science Outsourcing, Inc.

830 Challenger Street, Brea, CA 92821

Contact Name: Matthew Emrick Phone: 714-406-2085

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
Mechanical	Medical Device Sterile Barrier Packaging	Measuring the Force Necessary to Pull the Seal apart	ASTM F88	Calibrated Tensile Tester	F0	F
Mechanical	Medical Device Sterile Barrier Packaging	Observation of Bubbles Emitting from an Internally Pressurized Sterile Barrier Packaging	ASTM F2096	Pressure induced Packaging using Calibrated gauge	F0	F
Mechanical	Medical Device Sterile Barrier Packaging	Observation of Dye Penetrating the Seal on Porous Packaging	ASTM F1929	Sterile Barrier seal exposed to formulated dye	F0	F
Mechanical	Medical Device Sterile Barrier Packaging	Observation of dye penetrating the seal on Non-porous Packaging	ASTM F3039	Sterile Barrier seal exposed to formulated dye	F0	F
Mechanical	Medical Device Sterile Barrier Packaging	Observation of delamination once the Seal has been pulled apart	EN 868-5	Observation Only	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Observation of Shipping Container Damage	ASTM D4169, ASTM D5276	Shock Producing Drop Tester Verified With Calibrated Ruler for Height	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Measuring force received from vertical compression	ASTM D4169, ASTM D642	Calibrated Tensile/Compression Tester	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Exposing Shipping Container to Repetitive Shock	ASTM D4169, ASTM D999	Calibrated vibration table	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Exposing Shipping Container to Low pressure representing High altitudes	ASTM D4169, ASTM D6653	Vacuum pressure Container and pump With Calibrated gauge	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Exposing Shipping Container to Concentrated Impact	ASTM D4169, ASTM D6344	Calibrated Weight Dropped from specific Height to Attempt to Penetrate the Shipping Container	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Exposing Shipping Container to Concentrated Impact (for Long Packages Only)	ASTM D4169, ASTM D5265	Calibrated Weight Dropped from specific Height to Attempt to Penetrate the Shipping Container	F0	F



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Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Exposing Shipping Container to Repetitive Shock	ISTA 1A	Calibrated vibration table	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Observation of Shipping Container Damage	ISTA 1A	Shock Producing Drop Tester Verified With Calibrated Ruler for Height	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Observation of Shipping Container Damage	ISTA 2A	Shock Producing Drop Tester Verified With Calibrated Ruler for Height	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Measuring Force received from Vertical Compression	ISTA 2A	Calibrated Tensile/Compression Tester	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Observation of Shipping Container Damage	ISTA 3A	Shock Producing Drop Tester Verified With Calibrated Ruler for Height	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Observation of Shipping Container Damage	ISTA 3A	Shock produced by letting Shipping Container fall from standing Position (on edge)	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Observation of Shipping Container Damage	ISTA 3A	Shock Produced by Letting Shipping Container fall from Standing Position (on block to Rotate when Falling)	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Exposing Shipping Container to Concentrated Impact (for Long Packages Only)	ISTA 3A	Calibrated Weight Dropped from Specific Height to Attempt to Penetrate the Shipping Container	F0	F
Mechanical	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Observation of Shipping Container Damage	ISTA 1G	Shock Producing Drop Tester Verified With Calibrated Ruler for Height	F0	F
Non-destructive	Medical Device Sterile Barrier Packaging	Measuring the seal width	ASTM F2203	Measurements through Calibrated Calipers or Calibrated Ruler	F0	F



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Non-destructive	Medical Device Sterile Barrier Packaging	Measuring the seal width	EN 868-5	Measurements through Calibrated Calipers or Calibrated Ruler	F0	F
Non-destructive	Medical Device Sterile Barrier Packaging	Observing Primary Packaging for Liquid Leakage	ISTA 3A	Visual Observation Only	F0	F
Non-destructive	Medical Device Sterile Barrier Packaging	Observation for Defects Found on the Seal or Throughout the Sterile Barrier Packaging	ASTM F1886	Observation under 540 Lux With Calibrated Light Meter	F0	F
Environmental Simulation	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Exposing Shipping Container to Extreme Temperatures and Relative Humidities	ISTA 3A	Controlled Environment With Temperature and Relative Humidity Probes	F0	F
Environmental Simulation	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Exposing Shipping Container to Extreme Temperatures and Relative Humidities	ASTM D4332	Controlled Environment With Temperature and Relative Humidity Probes	F0	F
Environmental Simulation	Shipping Container With Sterile Barrier Medical Device Packaging and Devices	Exposing Shipping Container to Elevated Temperatures and Moderate to Low Humidities	ASTM F1980	Controlled Environment With Temperature and Relative Humidity Probes	F0	F
Biological	Biological Indicator Inoculated With Microorganisms	Exposing Biological Indicator to Biological Temperatures for Microbial Growth After Sterilization	ANSI/AAMI/ISO 11138-3	Calibrated Incubator	F0	F
Biological	Biological Indicator Inoculated With Microorganisms	Exposing Biological Indicator to Biological Temperatures for Microbial Growth After Sterilization	ANSI/AAMI/ISO 11138-1	Calibrated Incubator	F0	F
Biological	Biological Indicator Inoculated With Microorganisms	Exposing Biological Indicator to Biological Temperatures for Microbial Growth After Sterilization	ANSI/AAMI/ISO 11138-2	Calibrated Incubator	F0	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

