PERRY JOHNSON LABORATORY ACCREDITATION, INC.

Certificate of Accreditation

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

RMS Company

8600 Evergreen Blvd, Minneapolis, Coon Rapids, MN 55433

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:	*Issue Date:*	*Expiration Date:*
August 03, 2026	August 03, 2026	November 30, 2028

Accreditation No.:	*Certificate No.:*
136871	L26-697

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

RMS Company

8600 Evergreen Blvd, Minneapolis, Coon Rapids, MN 55433
Contact Name: Tom Casey Phone: 763-783-6009

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	Implantable medical Devices, USP 1231 Purified Water	Bioburden	ISO 11737-1, ISO 19227, USP <1231>, MP-03376	Microbiological Vacuum Filtration, Incubators, Biological Hood, Micro Pipette	F2	F
Biological	Implantable medical Devices	Cytotoxicity	ISO 10993-5, ISO 19227, MP-03381	Cell Culturing, Microplate Reader Spectrophotometer, Micro Pipette	F2	F
Biological	Implantable medical Devices, USP 1231 Purified Water	Endotoxin	ISO 11737-3, USP <85>, ISO 19227, USP <1231>, MP-03380	Microplate reader, Spectrophotometer, Micro Pipette	F2	F
Chemical	Implantable medical Devices	Gravimetric-Nonpolar Residues	ASTM F2459-18, ISO 19227, MP-03383	Weight Scale, Accelerated Solvent Extractor	F2	F
Chemical	Implantable medical Devices	Gravimetric-Polar Residues	ASTM F2459-18, ISO 19227, MP-03383	Weight Scale, Accelerated Solvent Extractor	F2	F
Chemical	Implantable medical Devices	Total Hydrocarbon	ASTM F2459-18, ISO 19227, MP-03382	GC, Micro pipette	F2	F
Chemical	Implantable medical Devices	Particulates	USP <788>, ISO 19227, MP-03379	Beckman Coulter HIAC 9703+	F2	F
Chemical	Implantable medical Devices, USP 1231 Purified Water	Total Organic Carbon	USP <643>, ISO 19227, USP <1231>, MP-03378	TOC Analyzer	F2	F

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Chemical	Implantable medical Devices	Inorganic Contaminants	USP <232>, USP <233>, ISO 19227, MP-03412	ICP-MS	F2	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