PERRY JOHNSON REGISTRARS, INC.

Certificate of Registration

Perry Johnson Registrars, Inc., has audited
the Quality Management System for Medical Devices:

Leinco Technologies, Inc.
410 Axminister Drive, Fenton, MO 63026 United States

(Hereinafter called the Organization) and hereby declares that
Organization is in conformance with:

ISO 13485:2016

This Registration is in respect to the following scope:

Design/Development and Manufacture of Human Diagnostic Components and Reagents for use in the Medical Device Industry

(Product Category: Reagents and Reagent Products, Calibrators, and Control Materials for: Clinical Chemistry, Immunochemistry (Immunology), Haematology/Haemostasis/Immunohematology, Microbiology, Infectious Immunology, Histology/Cytology, Genetic Testing)

This Registration is granted subject to the system rules governing the Registration referred to above, and the Organization hereby covenants with the Assessment body duty to observe and comply with the said rules.

PERRY JOHNSON REGISTRARS, INC.

ANAB
ANSI National Accreditation Board
ACCREDITED
ISO/IEC 17021-1
MANAGEMENT SYSTEMS CERTIFICATION BODY

Terry Boboige

Terry Boboige, President
Perry Johnson Registrars, Inc. (PJR)
755 West Big Beaver Road, Suite 1340
Troy, Michigan 48084
(248) 358-3388

The validity of this certificate is dependent upon ongoing surveillance.

Effective Date:	Expiration Date:	Certificate No.:
July 10, 2022	July 9, 2025	C2022-02822