



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.*



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:
July 10, 2022

Expiration Date:
July 9, 2025

Certificate No.:
C2022-02822