# CERTIFICATE OF REGISTRATION

**This is to certify that the management system of:**

## LKC Technologies, Inc.

(FIN F003655)

Main Site: 2 Professional Drive, Suite 222

Gaithersburg, MD 20879 USA

**has been registered by Intertek, an MDSAP recognized auditing organization, as conforming to the requirements of:**

## ISO 13485:2016

**Australia**: Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6)

**Brazil**: Federal Law n. 6360/76; RDC ANVISA n. 16/2013; RDC ANVISA n. 23/2012; RDC ANVISA n. 67/2009; RDC ANVISA n. 56/2001

**Canada**: Medical Devices Regulations – Part 1- SOR 98/282

**United States**: 21 CFR 820, 21 CFR 803, 21 CFR 806, 21 CFR 807 (Subparts A to D)

**Japan**: MHLW Ministerial Ordinance 169, Article 4 to Article 68; PMD Act

### The management system is applicable to:

*The design, development, production, installation, distribution and servicing of visual electrodiagnostic equipment.*

**Certificate Number:**
0089398-01

**Initial Certification Date:**
2019-04-10

**Date of Certification Decision:**
2022-04-28

**Certification Effective Date:**
2022-04-28

**Certification Expiry Date:**
2025-04-09

ISO 13485:2016 CERTIFICATION™

MDSAP MEDICAL DEVICE SINGLE AUDIT PROGRAM

intertek

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