Global Inter Certification

Management System Certification Body

CERTIFICATE

No. 25-B-0075 Rev.1

This is to certify that the Medical Devices-Quality Management Systems of

KMI IMI Group

4 Autry Irvine, California 92618, USA

Company Reg. No.: 33-0896073

has documented and implemented system in compliance with the requirements of

ISO 13485:2016
Medical Devices-Quality Management Systems

for

Design, Development & Manufacture & Sales of Surgery Equipment, Surgical Instruments, Cannulas, Medical Canister, Surgical Supplies, Medical LED Lamps & PRP Extraction Kit

Technical Area:

Non-active Medical Devices- General non-active, non-implantable medical devices, Non-active medical devices other than specified above

The certificate is issued on the basis of the results mentioned in the pertinent audit report.
Validity of the certificate is conditionally limited by positive results of surveillance audits, which the certified company is committed to undergo.

The validity of this certificate can be checked on the main page of the GIC website www.gicert.org.

This certificate can be invalid if the certificate holder does not fulfill the conditions set out in the certification agreement.

ISO 13485 CERTIFIED

CB-MS-12731

Initial issue date:	Jan. 24. 2019
Re-issue date:	Apr. 22. 2026
Expire date:	Jan. 23. 2028

Tyrone Dyse
Head of Certification Body