



Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

DeviceLab, Inc.
3002 Dow Avenue, Unit 124
Tustin
California
92780
USA

Holds Certificate No:

FM 576996

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 for the following scope:

Contract design and product development for medical devices.

For and on behalf of BSI:

Stewart Brain, Head of Compliance & Risk - Medical Devices

Original Registration Date: 2011-08-26

Latest Revision Date: 2019-01-17

Effective Date: 2017-08-26

Expiry Date: 2020-08-25

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