

The management system of

MEDIMARK EUROPE

11 rue Emile Zola, BP 2332
38100 GRENOBLE Cedex 2
France



has been assessed and certified as meeting the requirements of

ISO 9001 : 2015

For the following activities

**European authorized representative activities
for medical devices, active implantable devices
and in vitro diagnostic medical devices.**

This certificate is valid from 8 May 2017 until 7 May 2020
and remains valid subject to satisfactory surveillance audits.
Issue 8. First Certification date December 1998

Authorised by

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