

CERTIFICATE



Management System as per EN ISO 13485 : 2016

In accordance with TÜV AUSTRIA procedures, it is hereby certified that

LERIVA PHARMA S.A.

HEAD OFFICE:33, Pigis Avε.& 6, Anapiron Polemou Str., GR-151 27 MELISSIA, GREECE BRANCH 1: Lakkos Kamaterou, GR-193 00 ASPROPIRGOS, GREECE BRANCH 2: 12th KM. Thessalonikis-Moudanion Ave., GR- 570 01 THESSALONIKI, GREECE

Applies a Quality Management System for Medical Devices in line with the above Standard for the following Scope

IMPORT, TRADING AND DISTRIBUTION OF:

 ACTIVE MEDICAL DEVICES AND ASSOCIATED CONSUMABLES IN THE FIELD OF RADIOLOGY AND CARDIOLOGY

• GENERAL, NON-ACTIVE, NON-STERILE SELF-CARE MEDICAL DEVICES

TECHNICAL SUPPORT OF ACTIVE MEDICAL DEVICES IN THE FIELD OF RADIOLOGY AND CARDIOLOGY

Certificate Registration No.: 20302190002308

Myannoi

Maria Agapitou Head of Management Systems & Products Certification Division

Certification Body at TÜV AUSTRIA

Valid until: 2025-10-16 Initial certification: 2019-10-17

Athens, 2025-04-29

This certification was conducted in accordance with TÜV AUSTRIA auditing and certification procedures and is subject to regular surveillance audits.

TÜV AUSTRIA HELLAS 429, Mesogeion Ave. GR-153 43 Athens, Greece <u>www.tuvaustriahellas.gr</u> GEMI No: 1650201000



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Headquarters in Athens bear the responsibility of the Certification decision



approval by TÜV AUSTRIA

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