Qarad BV

ISO 13485:2016

this approval are listed.

Marta Escudero

Regional Director, Europe Issued by: LRQA Limited

Groenenborgerlaan 16, 2610 Wilrijk, Belgium

Approval number(s): ISO 13485 - 0019831

The scope of this approval is applicable to:

and provision of e-labeling services for IVD and MD industry.

This certificate is valid only in association with the certificate schedule bearing the same number on which the locations applicable to

The sales and provision of regulatory and quality consulting for in vitro diagnostics (IVD), authorized representative services for IVD's & medical devices (MD), medical writing for IVD's and performance evaluations studies for IVD's. The design and development, sales

Certificate of Approval

This is to certify that the Management System of:

has been approved by LRQA to the following standards:

14 September 2024 13 September 2027 10628557

Original approval(s): ISO 13485 - 14 September 2012

assumes no responsibility and shall not be liable to any person for any loss, damage or expense caused by reliance on the information or advice in this document or howsoever provided, unless that person has signed a contract with the relevant LRQA entity for the provision of this information or advice and in that case any responsibility or liability is exclusively on the terms and conditions set out in that contract.

LRQA Group Limited, its affiliates and subsidiaries and their respective officers, employees or agents are, individually and collectively, referred to in this clause as 'LRQA'. LRQA





Certificate Schedule

Location Activities

Qarad BV

Cipalstraat 3, 2440 Geel, Belgium

ISO 13485:2016

The sales and provision of regulatory and quality consulting for in vitro diagnostics (IVD), authorized representative services for IVD's & medical devices (MD), medical writing for IVD's and performance evaluations studies for IVD's. The design and development, sales and provision of e-labeling services for IVD and MD industry.

Qarad EC-REP BV

Pas 257, 2440 Geel, Belgium

ISO 13485:2016

The sales and provision of regulatory and quality consulting for in vitro diagnostics (IVD), authorized representative services for IVD's & medical devices (MD), medical writing for IVD's and performance evaluations studies for IVD's. The design and development, sales and provision of e-labeling services for IVD and MD industry.

QbD Flanders NV

Groenenbrogerlaan 16, 2610 Wilrijk, Belgium

ISO 13485:2016

The sales and provision of regulatory and quality consulting for in vitro diagnostics (IVD), authorized representative services for IVD's & medical devices (MD), medical writing for IVD's and performance evaluations studies for IVD's. The design and development, sales and provision of e-labeling services for IVD and MD industry.

Qarad Suisse S.A.

World Trade Center Av. de Gratta-Paille 2, CH 1018 Lausanne, Switzerland

ISO 13485:2016

The sales and provision of regulatory and quality consulting for in vitro diagnostics (IVD), authorized representative services for IVD's & medical devices (MD), medical writing for IVD's and performance evaluations studies for IVD's. The design and development, sales and provision of e-labeling services for IVD and MD industry.



LRQA

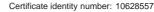
LRQ/

LRQA Group Limited, its affiliates and subsidiaries and their respective officers, employees or agents are, individually and collectively, referred to in this clause as 'LRQA'. LRQA assumes no responsibility and shall not be liable to any person for any loss, damage or expense caused by reliance on the information or advice in this document or howsoever provided, unless that person has signed a contract with the relevant LRQA entity for the provision of this information or advice and in that case any responsibility or liability is exclusively on the terms and conditions set out in that contract.

Page 2 of 3

2 of 3

Issued by: LRQA Limited, 1 Trinity Park, Bickenhill Lane, Birmingham B37 7ES, United Kingdom





Certificate Schedule

Location **Activities**

Qarad UK Ltd.

8 Northumberland Avenue, London, WC2N 5BY, United Kingdom

ISO 13485:2016

The sales and provision of regulatory and quality consulting for in vitro diagnostics (IVD), authorized representative services for IVD's & medical devices (MD), medical writing for IVD's and performance evaluations studies for IVD's. The design and development, sales and provision of e-labeling services for IVD and MD industry.

LRQA Group Limited, its affiliates and subsidiaries and their respective officers, employees or agents are, individually and collectively, referred to in this clause as 'LRQA'. LRQA assumes no responsibility and shall not be liable to any person for any loss, damage or expense caused by reliance on the information or advice in this document or howsoever provided, unless that person has signed a contract with the relevant LRQA entity for the provision of this information or advice and in that case any responsibility or liability is exclusively on the terms and conditions set out in that contract.