



Declaration of Conformity

EUROIMMUN Medizinische Labordiagnostika AG
Seekamp 31, 23560 Lübeck, Germany

declares under its sole responsibility as manufacturer that the ELISA product

Anti-LC-1 ELISA (IgG)

EA 1307-9601 G

(product name, order number)

meets the following demands of:

Directive 98/79/EC on in vitro diagnostic medical devices of 27 October 1998 and its transpositions in national laws which apply to it.

Conformity assessment procedure: Annex III

This Declaration of Conformity is valid based on the respective currently valid version of technical documentation.

Lübeck, May 19 2022

(Place and date of issue)

Dr. Ewald Müller-Kunert
- Head of Quality Management -

Susanne Aleksandrowicz
- Member of the Executive Board -