EC Declaration of Conformity

Qlife Aps
Borupvang 3,
DK-2765 Ballerup,
Denmark

We hereby declare that the product(s) described below meets the appliable requirements of Directive 98/79/EC of the European Parliament and of the Council of October 27, 1998, on *in vitro diagnostic* medical devices (IVDD) as specified in Annex III:

Class: ☒General ☐ Annex II/List A ☐Annex II/ List B

☐ Self-testing ☐ Performance Evaluation

Product:

Model name	Article No.	GMDN Code*
Q200 Egoo system,	e78852	61039

*According to the nomenclature provided in ISO/TS-20225

Notified Body:
As Specified in the Directive and Annex mentioned above, the conformity assessment procedure in this class does not require the involvement of a Notified Body.

Issuance:
Name: Kristina Christensen
Title: VP QA/RA

Place: Qlife Aps
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Kristina Christensen

Signature:

2021-01-24

Date: