Certificate of CE-Registration

This is to certify that, in accordance with either medical device Directive 93/42/EEC as amended by 2007/47/EC or Directive 98/79/EC, mdi Europa GmbH agree to perform all duties and responsibilities as the Authorized Representative for

Cue Inc.
4980 Carroll Canyon Rd.
Suite 100
CA 92121 San Diego
USA

as stipulated and demanded by the afore-mentioned Directives. The German competent authorities have allocated the medical devices of the manufacturer the following registration numbers:

EDMA Code	Description	Classification	Registration Number
1504409000	OTHER VIROLOGY - NA REAGENTS Cue COVID-19 Test	other IVD	DE/CA09/0760/C18/IVD/001-01

The manufacturer has provided mdi Europa with all necessary documentation, together with an appropriate Declaration of Conformity confirming that the medical devices fulfill the essential requirements of either Directive 93/42/EEC as amended by 2007/47/EC or 98/79/EC. A safety officer has been appointed for Germany and therefore is in full compliance with § 31 MPG.

Signed on 11 December 2020

Werner Sander
President