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

Tarsons Products Ltd.
902, Martin Burn Business Park, BP-3, Saltlake, Sector-V
West Bengal
700091 Kolkata
India

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
29.01.10	other Hardware + accessories + consumables + software	other IVD	DE/CA09/0760/T12/IVD/001-03

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 02 August 2021

Werner Sander
President