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QMD™ Degania™ Biometrix™ Arthesys™

Biometrix™ PERFORMANCE FOR LIFE

To whom it may concern:

Date: October 14, 2024

EXTENSION OF DECLARATIONS OF CONFORMITY

We, Biometrix Ltd located at Kyriat Mada 4, 9145000 Jerusalem, Israel, hereby declare that all our products are marketed in compliance with Annex II excluding 4 (CE certificate G10023740005 Rev.00, devices in class IIa, IIb and III) and Annex V (CE certificate G2S 002374 0006 Rev.00, devices in class I sterile) of Medical Device Directive, MDD 93/42/EEC as amended by 2007/47/EC

Due to the confirmation letters CL 002374 0010 Rev. 01, CL 002374 0011 Rev. 00, CL 002374 0012 Rev. 00 received from the Notified body TÜV SÜD Product Service GmbH (NB#0123), regarding the formal application of Biometrix Ltd products in the transition period, all Declarations of conformity (DoC) are valid till the transition timelines shown below:

- 31 December 2027 for Class III devices and Class IIb implantable devices (except sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- 31 December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition, measuring function
- 31 December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition, measuring function

Karina Janekova,

RA Manager,

Biometrix Ltd

DocuSigned by:
Karina Janekova
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