MOTOmed®

# Annex to EC-Declaration of Conformity
# MOTOmed viva2 of 3 August 2018

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|---|---|
| Manufacturer: | **RECK-Technik GmbH & Co. KG** Medical Devices<br>Reckstraße 1–5, 88422 Betzenweiler, GERMANY<br>Email: vigilance@motomed.com |
| Name of Product: | **MOTOmed viva2**<br>Basic UDI-DI: 4260193710017M<br>REF 200.003, 200.012, 200.004, 200.021, 200.008 |

We, as the manufacturer declare under our sole responsibility in reference to Regulation (EU) 2017/745 in conjunction with Regulation (EU) 2023/607 of 15 March 2023 that our Declaration of Conformity according to Annex II of Council Directive 93/42/EEC (2007/47/EC) for our medical devices **MOTOmed viva2 of 3 August 2018** is valid until 26 May 2024 due to the extended transition period for class IIa medical devices.

In this context, we reference our manufacturer's declaration according to Regulation (EU) 2023/607 of 24 July 2023 (P03D56rev02).

RECK-Technik GmbH & Co. KG
Betzenweiler, 25 July 2023

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RECK