

EU Declaration of Conformity

Manufacturer name: **Kommanditgesellschaft Behnk Elektronik GmbH & Co.**
Manufacturer address: **Hans-Böckler-Ring 27, 22851 Norderstedt, GERMANY**
SRN: **DE-MF-000005304**
Tel: +49 (40) 529 861 0
Email: regulatory@behnk.de
Internet: www.behnk.de

Basic UDI-DI: ++EUBECOASETRTMC4A
Device Name: **Cuvette racks and balls**
REFERENCE: **050210**

UDI-DI: +EUBE0502100W
EMDN: W0202020185
GIVD / EDMA: 23.02
GMDN: 56690

Classification: A in accordance with rule 5a of Annex VIII of (EU) 2017/746
Technical file date: 2024-11-21

This declaration of conformity is issued under the sole responsibility of **Kommanditgesellschaft Behnk Elektronik GmbH & Co.**

We hereby declare that the in vitro diagnostic device specified above meets the provision of the Regulation (EU) 2017/746 on in vitro diagnostic medical devices. This declaration is supported by the Quality System approval to ISO 13485 issued by "Deutsche Akkreditierungsstelle" D-ZM-19630-04-00.

Norderstedt, 2025-02-20



Kommanditgesellschaft
Behnk Elektronik GmbH & Co. 
Hans-Böckler-Ring 27 || 22851 Norderstedt || DE
+49 40 529 861 0 +49 40 524 10 94
www.behnk.de



Peter Letsinger
Person responsible for regulatory compliance