

**Certificate of Verification of Manufactured Products**  
**Directive 98/79/EC Annex IV, Section 6**  
**In Vitro Diagnostic Medical Devices**

**Registration No.:** LL 60031592 0039

**Report No.:** 21110677 061

**Manufacturer:**

Bio-Rad Medical Diagnostics GmbH  
Industriestr. 1  
63303 Dreieich  
Deutschland

**Product Identification:**

Seraclone® Anti-CDE  
Lot No.: 3320080-00  
Article No.: 802080

The Notified Body hereby declares that a verification relating to the listed products has been performed according to Annex IV, section 6 of the directive 98/79/EC and that the devices conform to the requirements of the abovementioned directive.

Notified Body



Date: 2023-09-07

  
Katja Mierisch

**TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg**

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 98/79/EC concerning in vitro diagnostic medical devices with the identification number 0197.