

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

**Registration No.:** LL 60030333 0032

**Report No.:** 21110629 078

**Manufacturer:**

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

**Product Identification:**

Seraclone® Anti-D (RH1) 226  
Lot No.: 3306081-00  
Article No.: 802039, 802042

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: 2023-04-12

  
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.