

Certificate of Verification of Manufactured Products

Directive 98/79/EC Annex IV, Section 6
In Vitro Diagnostic Medical Devices

Registration No.: LL 60033857 0016

Report No.: 21112948 027

Manufacturer:

Bio-Rad Medical Diagnostics GmbH

Industriestr. 1 63303 Dreieich Deutschland

Product Identification:

Date: 2022-05-16

Seraclone® Control ABO+Rh

Lot No.: 3209040-00 Article No.: 805171

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

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.