Certificate of Verification of Manufactured Products

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

Registration No.: LL 60034087 0028

Report No.: 21138220 031

Manufacturer:

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

Product Identification:

Solidscreen II Anti-D (RH1) Blend
Lot-Nr.: 3232051-00
Article No.: 806530

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

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

Date: 2022-10-10

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.