Certificate of Verification of Manufactured Products

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

Registration No.: LL 60031012 0046

Report No.: 21114637 065

Manufacturer:

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

Product Identification:

Erytype S Rh+K reagent I
Lot No.: 3307010
Article No.: 806180

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: 2023-03-17

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.