

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

Registration No.: LL 60032396 0032

Report No.: 21110693 072

Manufacturer:

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

Product Identification:

Erytype® ABO+D
Lot No.: 3247010
Article No.: 806010

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: 2022-12-14


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.