

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

Registration No.: LL 60034125 0034

Report No.: 21111096 063

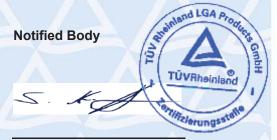
Manufacturer:

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

Product Identification:

Seraclone® Anti-e (RH5) Lot No.: 3228081-00 Article No.: 802370, 802372

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.



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TÜV, TUEV and TUV are registered trademarks. Utilisation and application requires prior approva

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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.