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Verification Report

according to Regulation (EU) 2017/746 on In vitro Diagnostic Medical Devices – Annex IX, sec. 4.12 or Annex XI sec. 5

No. BRM-808188 25 04 072457

Manufacturer: **Bio-Rad Medical Diagnostics GmbH**
Industriestraße 1
63303 Dreieich

Product: **Seraclone Anti-Fya (FY1)**

Testplan: **TP-BRM-808188**

Batch: **3512041-00**

Basic UDI-DI: **361052A004037V**

Expiry Date: **2027-03-11**

The above mentioned batch meets the batch release criteria established during technical documentation assessment and may be placed on the market. The EU Technical Documentation Assessment Certificate (IVDR) issued for this product is V70 118577 0007 Rev. 00.

This verification report is valid for REF 808188.

Date, 2025-04-29

Adobe Acrobat Sign-Transaktionsnummer: CBJCHBCAABAAR2mGI6QC26Lq_vaQfQFP0W8J-jEVnrga
pp Dr. Carsten Krumbholz
In-vitro Diagnostics

TÜV SÜD Product Service GmbH is Notified Body according to Council Vitro Diagnostic Regulation 2017/746 concerning In-vitro Diagnostic Medical Devices with Identification No. 0123.

To:
Bio-Rad Medical Diagnostics GmbH

cc: