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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-12012453 24 09 069430

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

**Product:** **Biotestcell-O**

**Testplan:** **TP-BRM-12012453**

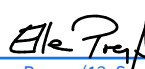
**Batch:** **3436011-00 (REF 12012453)**

**Basic UDI-DI:** **361052A003878Q**

**Expiry Date:** **2024-11-04**

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 0001 Rev. 00.

**Date,** 2024-09-12

  
Ellen Preuß (12. September 2024 15:06 GMT+2)  
Adobe Acrobat Sign-Transaktionsnummer: CBJCHBCAABAAr7Wu9PwxUVspD1zJNr66BLcX6Lqmiq3p  
pp Dr. Ellen Preuß  
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:

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