

Declaration of Conformity

Certificate Identification: DOC-08P6101-SD DLK TPM
Legal Manufacturer's Name: Abbott GmbH
Legal Manufacturer's Address: Max-Planck-Ring 2, 65205 Wiesbaden, Germany

List Numbers and Size Code of Devices	GMDN Code	Names and Description of Devices	Classification
08P6101	41830	Alinity c Bilirubin Calibrator Kit	Self-declared

Authorized European Representative (name and address)	N/A
Storage site of technical documentation (name and address)	Abbott Laboratories, 1921 Hurd Drive, Irving, Texas 75038, USA
Harmonized Standards	Listed in the Technical Documentation

We, the undersigned, hereby declare that the in vitro diagnostic medical devices described above and bearing the CE marking, conform with the applicable provisions of the EC Directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on In Vitro Diagnostic Medical Devices as they are transposed into the laws of the member states.

This declaration is made in accordance with Annex III of the IVD Directive and is issued under the sole responsibility of the manufacturer.

<p>Signature: <u> <i>C. Becker</i> </u></p> <p>Full Name: Claudia Becker</p> <p>Position: Director Quality Assurance</p> <p>Date of Approval: <u> 02 Feb 2022 </u></p>	<p>Signature: <u> <i>Tiffini Jenkins</i> </u></p> <p>Full Name: Tiffini Jenkins</p> <p>Position: Manager Regulatory Affairs</p> <p>Date of Approval: <u> 1-Feb-2022 </u></p> <p>Date Issued: <u> 02 Feb 2022 </u></p> <p>Place Issued: 65205 Wiesbaden, Germany</p> <p>Supersedes: 05-April-2017</p> <p>Effective (Date or Lot Number): <u> 02 Feb 2022 </u></p>
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