



EU Declaration of Conformity

Basic UDI-DI: 038074SLI0002T5
Basic UDI-DI Name: ARCHITECT Concentrated Wash Buffer
Risk Class: Class A

List Number and Size Code	Product and Trade Name	GMDN Code	EMDN Code
6C54-58	ARCHITECT Concentrated Wash Buffer	58236	W0201020185
6C54-82	ARCHITECT Concentrated Wash Buffer	58236	W0201020185
6C54-88	ARCHITECT ARM Concentrated Wash Buffer	58236	W0201020185

Manufacturer (Name and Address)	Abbott Ireland Diagnostics Division Finisklin Business Park Sligo, Ireland
Manufacturer SRN	IE-MF-000009849
Authorized Representative (Name and Address)	N/A
Authorized Representative SRN	N/A
Produced by (Site of Manufacture) (Name and Address)	Abbott Ireland Diagnostics Division Finisklin Business Park Sligo, Ireland
Conformity Assessment Procedure	Annex II and III

We, the undersigned, hereby declare that the in vitro diagnostic medical device(s) described above conform with the applicable provisions of the Regulation (EU) 2017/746 of the European Parliament and of the Council of 5 April 2017 on In Vitro Diagnostic Medical Devices. This declaration is made in accordance with Annex IV of the IVD Regulation and is issued under the sole responsibility of the manufacturer.

Full Name: <u>Noel Haren</u>	Full Name: <u>Joe Murray</u>
Function: <u>Manager Regulatory Affairs</u>	Function: <u>Director Quality Assurance</u>
Signature: <u>[Signature]</u>	Signature: <u>[Signature]</u>
Date of Approval: <u>15 Jul 2022</u>	Date of Approval: <u>15 Jul 2022</u>
Signed for, and on behalf of: <u>Abbott Ireland Diagnostics Division, Sligo</u>	
Date Issued: <u>15 Jul 2022</u>	Place Issued: <u>Sligo, Ireland</u>
Supersedes: <u>23 May 2022</u>	Effective (Date or Lot Number): <u>15 Jul 2022</u>