



EU Declaration of Conformity

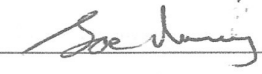
Basic UDI-DI: 038074SLI0002T5
Basic UDI-DI Name: ARCHITECT Pre-Trigger Solution
Risk Class: Class A

List Number and Size Code	Product and Trade Name	GMDN Code	EMDN Code
6E23-68	ARCHITECT Pre-Trigger Solution	61163	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: Noel Haren
Function: Manager Regulatory Affairs
Signature: 
Date of Approval: 15 Jul 2022
Signed for, and on behalf of: Abbott Ireland Diagnostics Division, Sligo
Date Issued: 15 Jul 2022
Supersedes: 23 May 2022

Full Name: Joe Murray
Function: Director Quality Assurance
Signature: 
Date of Approval: 15 Jul 2022
Place Issued: Sligo, Ireland
Effective (Date or Lot Number): 15 Jul 2022