IVD Technical File Declaration of Conformity for LumiraDx INR Quality Controls (S-RA-TEC-0003)

Document Number:	S-RA-REP-00211	Revision:	2

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EC Declaration of Conformity
(EN)

We, the legal manufacturer, as stated below herby declares under our sole responsibility,

Legal Manufacturer:	LumiraDx UK Ltd
Address:	Dumyat Business Park Alloa FK10 2PB United Kingdom
EC Authorized Representative:	LumiraDx AB Västra Vägen 5A 169 61 Solna Sweden

that the identified product to which this declaration relates,

Identification	
Product Name	LumiraDx INR Quality Controls
Catalogue Number	L003080101003
GMDN Code	55985 – Prothrombin time (PT), IVD, control
Classification	General IVD
Conformity Assessment Route	Annex III (excluding Section 6) of 98/79/EC

is in conformity with the following European Regulations and Directives as transposed into the national laws of the member states:

Directives and Regulations
Directive 98/79/EC of the European Parliament and of the Council of 27 October 1998 on in vitro diagnostic medical devices

Signed for and on the behalf of Manufacturer LumiraDx UK Limited:

Name	David Scott	**Position:**	Chief Technology Officer
Signature		**Date:**	
		8th June 2021	

With approval of this Declaration of Conformity, we hereby affix the CE Mark to the product.