IVD Technical File Declaration of Conformity for LumiraDx SARS-CoV-2 Ab Test Strips

S-RA-TEC-0009

Document Number:	S-RA-REP-00128	Revision:	2
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Contents

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Information Classification: Public

LumiraDx

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 products to which this declaration relates,

Identification	
Product Name	LumiraDx SARS-CoV-2 Ab Test Strips
Catalogue Numbers	L0170001nnxxx (where nn represents two digits corresponding to language variants, and where xxx represents three digits corresponding to number of units the catalogue number contains).
GMDN Code	66470, SARS-CoV-2 immunoglobulin G (IgG)/IgM antibody IVD, kit, rapid microfluidic immunoassay, clinical
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:	
Original Declaration of Conformity signed (S-RA-REP-00128 Rev 1) by David Scott		18 September 2020	
DS.		10 Nov 2022	

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

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