UKCA Declaration of Conformity for LumiraDx SARS-CoV-2 & Flu A/B and Quality Controls (S-RA-TEC-0012)

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Information Classification		Public	

Contents

UK Declaration of Conformity for LumiraDx SARS-CoV-2 & Flu A/B (EN) and LumiraDx SARS-CoV-2 & Flu A/B Quality Controls (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

that the identified products to which this declaration relates,

Identification	
Product Name	LumiraDx SARS-CoV-2 & Flu A/B
Catalogue Numbers	L019000101012, L019000101024, L019000101048, L019000102012, L019000102024, L019000102048, L019000104012,L019000104024, L019000104048, L019000105012, L019000105024, L019000105048, L019000108012, L019000108024, L019000108048
GMDN Code	66045 - SARS-CoV-2/influenza A/B antigen IVD, kit, fluorescent immunoassay, rapid
Classification	General IVD
Conformity Assessment Route	The Medical Devices Regulations 2002 (SI 2002 No 618, as amended) (UK MDR 2002) through Annex III (excluding Section 6) of 98/79/EC
Product Name	LumiraDx SARS-CoV-2 & Flu A/B Quality Controls
Catalogue Numbers	L019080101002, L019080102002
GMDN Code	65646 - SARS-CoV-2/influenza A/B antigen IVD, control
Classification	General IVD
Conformity Assessment Route	The Medical Devices Regulations 2002 (SI 2002 No 618, as amended) (UK MDR 2002) through Annex III (excluding Section 6) of 98/79/EC

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

Directives and Regulations
SI 2002 No 618 (As Amended) Medical Devices Regulations 2002

This declaration is also supported by the following applied standards:

Standards/Common Specifications	
EN ISO 13485:2016	Medical Devices – Quality Management Systems – Requirements for Regulatory Purposes
EN ISO 14971:2019	Medical Devices – Applications of Risk Management to Medical Devices

Full responsibility and compliance of the product is signed for and on the behalf of Manufacturer LumiraDx UK Limited:

Name	David Scott	**Position:**	Chief Technology Officer
Signature		**Date:**	
Previously signed (S-RA-REP-00358 Rev 1) by David Scott under GMDN code for LumiraDx SARS-CoV-2 & Flu A/B test strips: 61885-Multiple respiratory pathogen antigen IVD, kit, fluorescent immunoassay		22 December 2021	
DScott.		04 MARCH 2022	

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