DECLARATION OF CONFORMITY

Manufacturer:	Illumina 5200 Illumina Way San Diego, CA 92122 United States
European Authorized Representative:	Illumina Netherlands B.V. Steenoven 19 5626 DK Eindhoven The Netherlands
Device Name:	TruSeq Custom Amplicon Dx Kit – FFPE QC
Device Model/Catalogue Number:	20006259
Basic UDI-DI	0081627002TSCAS6
Classification:	General IVD
Conformity Assessment Procedure:	Annex III of IVDD 98/79/EC Council Directive; Self-Declaration

We, Illumina, declare under our sole responsibility that the *in vitro* Diagnostic Device(s) categories specified above and provided with the CE marking, meet the provisions of the EC Directive 98/79/EC (including amendments issued in the years following) which apply to them.

This declaration is supported by the EC Quality System Certificate(s) according to the provisions of relevant Annex(es) of this Directive. This declaration applies to all devices specified above distributed from the signature date forward.

Authorized by:

Joe Mcmullen

*Electronically signed by: Joe Mcmullen
Reason: Approver
Date: Sep 21, 2021 09:13 PDT*

21-Sep-2021

E. Joseph McMullen
Sr. Director, Regulatory Affairs
Illumina, Inc.

Date

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Final Audit Report 2021-09-21

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