DECLARATION OF CONFORMITY

Manufacturer: Illumina
5200 Illumina Way
San Diego, CA 92122
United States

European Authorized Representative: Illumina Netherlands B. V.
Freddy van Riemsdijkweg 15
5657 EE Eindhoven
The Netherlands

Device Name: *TruSeq™ Custom Amplicon Dx Kit – FFPE QC

Device Model/Catalogue Number: 20006259

Basic UDI-DI: 0081627002TSCAS6

Classification: General IVD


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:

[Signature]

Bryan Schneider
Associate Director, Regulatory Affairs - HQ

[22-APR-2020]
Date (DD-MMM-YYYY)
### Device Component List

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<td>Device Components</td>
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