DECLARATION OF CONFORMITY

Manufacturer: Illumina
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San Diego, CA 92122
United States

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

Device Name: TruSight Cystic Fibrosis Library Prep

*Note: See device components for each of the device model/catalogue number on page 2 of this declaration of conformity*

Device Model/Catalogue Number: 20036925
Basic UDI-DI: 0081627002CYSTFIB8C
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 within this Declaration of Conformity and distributed onwards from the signature date below.

Authorized by:

Bryan Schneider
Associate Director, Regulatory Affairs - HQ

Date: 17-APR-2020
## Device Component List

**Device Name**
TruSight Cystic Fibrosis Library Prep 20036925

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