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: 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  
Illumina, Inc.

01-SEP-2021

Date (dd-mmm-yyyy)
### Device Component List

#### Device Name

- Device Components

**TruSight Cystic Fibrosis Library Prep 20036925**

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