

DECLARATION OF CONFORMITY

Manufacturer:

Illumina Inc.

5200 Illumina Way San Diego, CA 92122

USA

European Authorized Representative:

Illumina Cambridge Limited

Chesterford Research Park

Little Chesterford Saffron Walden CB10 1XL

United Kingdom

Device Name:

VeriSeq NIPT Solution

Device Model/Catalogue Number:

15066801, 15066802, 15076164

Classification:

Annex II, List B

Conformity Assessment Procedure:

Annex IV

Notified Body:

BSI

Notified Body Number 0086

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:

Mya Thomae

Vice President, Regulatory, Clinical and Medical Affairs

Date

pn/ 2017