illumina®

DECLARATION OF CONFORMITY

Manufacturer's Name:	Illumina, Inc.
European Authorized Representative:	Illumina Cambridge Limited Chesterford Research Park, Little Chesterford Saffron Walden, Essex, CB10 1XL UNITED KINGDOM
Device Name:	VeriSeq NIPT Analysis Software (16 samples)
Device Model/Catalogue Number:	RH-400-1001
Classification:	Non List A, List B
Conformity Assessment Procedure:	Annex III, Self-declared
Year of first CE Marking:	2016

I, the undersigned declare that the above mentioned in vitro diagnostic medical devices conform(s) with the requirements of the Directive, 98/79/EC of the Council of October 27, 1998 concerning *in vitro* diagnostic medical devices.

Authorized by:

Mya Thomae
Vice President, Regulatory Affairs

1-25-16
Date