

EU DECLARATION OF CONFORMITY

Product Name(s) / Trade Name(s) VeriSeq NIPT Workflow Manager v2

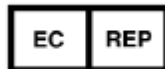


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Basic UDI-DI (BUDI-DI) 0081627002NIPTWFM9Q



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We, Illumina, as the manufacturer of the device(s) take sole responsibility for and hereby declare that the above mentioned product(s) meet(s) the provisions of the following Regulation(s)/Directives:

- Regulation EU 2017/746 on *In vitro* Diagnostic Medical Devices

RISK CLASS:

A B C D

CONFORMITY ROUTE:

Annex I & II+III of Regulation EU 2017/746; Self-Declaration

Common Specification (CS): N/A

E. Joseph McMullen
Illumina Regulatory Affairs

San Diego, CA _____

Issued in

08-FEB-2022

Date

Signature: *Joe McMullen*

Electronically signed
by: Joe McMullen
Reason: Approver
Date: Feb 8, 2022
13:55 PST

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Request Signature for VeriSeq NIPTv2 WFM DoC

Final Audit Report

2022-02-08

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"Request Signature for VeriSeq NIPTv2 WFM DoC" History

-  Document created by Debleena Sinha (dsinha@illumina.com)
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