

Title: NovaSeq 6000Dx IVDR EU Declaration of Conformity

Document Number: 200027832 Ver. 00

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EU DECLARATION OF CONFORMITY

Product Name(s) / Trade Name(s) NovaSeq[™] 6000Dx Instrument

> **Intended Purpose** The NovaSeq 6000Dx instrument is intended for sequencing

of DNA libraries when used with in vitro diagnostic (IVD) assays. The NovaSeq 6000Dx instrument is intended for use with specific registered, certified or approved IVD reagents

and analytical software.

REF 20068232

Basic UDI-DI (BUDI-DI) 0081627002NSQX2

NovaSeq 6000Dx S2 Reagent v1.5 Kit (300 cycles) Product Name(s) / Trade Name(s)

NovaSeq 6000Dx S4 Reagent v1.5 Kit (300 cycles)

NovaSeq 6000Dx S2 Buffer Cartridge NovaSeq 6000Dx S4 Buffer Cartridge

Intended Purpose The NovaSeg 6000Dx Reagent Kit (300 cycles) (S2 or S4)

> and NovaSeq 6000Dx Buffer Cartridge (S2 or S4) are reagents and consumables intended for sequencing of sample libraries when used with validated assays. The kit is intended for use with the NovaSeq 6000Dx instrument and

analytical software.

20046931 REF

20046933 20062292 20062293

Basic UDI-DI (BUDI-DI) 0081627002NSQKTTN

NovaSeq 6000Dx Library Tube

Product Name(s) / Trade Name(s) NovaSeq 6000Dx Library Tube, 24 pack

> **Intended Purpose** The NovaSeq 6000 Dx Library Tubes (single tube or 24

pack) are intended to hold sample libraries for sequencing when used with NovaSeq 6000Dx S2 or S4 reagent kits and validated assays. The library tube is intended for use with

NovaSeq 6000Dx instrument.

20062290 20062291

Basic UDI-DI (BUDI-DI) 0081627002NSQKTTN



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SRN: US-MF-000013476



Illumina Netherlands B.V. Steenoven 19 5626 DK Eindhoven The Netherlands

SRN: NL-AR-000012614

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 (Instrument and Reagents)
- Radio Equipment Directive 2014/53/EU (Instrument)
- RoHS Directive 2011/65/EU as amended by (EU) 2015/863 (Instrument) Annex III exemptions apply

RISK CLASS				
⊠A	□В	\Box C	\Box D	
CONFORMIT	Y ROUTE:			
Annex I & II+	III of Regulati	on EU 2017/74	6; Self-Declar	ation
Common Specification (CS): N/A				
Joe Mcmull	M Reason	Electronically signed by: Joe Mcmullen Reason: Approver Date: Sep 27, 2022 10:29 PDT		September 27, 2022
E. Joseph Mcmullen Date Sr. Director, Regulatory Affairs Illumina, Inc.				
San Diego, CA				
Issued in				

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