invivoscribe

DECLARATION OF CONFORMITY

Manufacturer:

Invivoscribe, Inc.
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Authorized Representative:

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Family Name: LymphoTrack® Dx

Device Trade Name: IGH FR1 Assay Kits – MiSeq

Catalog #	Device	Quantity	Basic UDI-DI	GTIN
9-121-0009	LymphoTrack® Dx *IGH* FR1 Assay Kit A – MiSeq®	40 Reactions	08100227391210009QE	00850052003647
9-121-0039	LymphoTrack® Dx *IGH* FR1 Assay Panel – MiSeq®	120 Reactions	08100227391210039QP	00850052003654

I, the undersigned, hereby declare that the *in-vitro* diagnostic medical devices specified above conform to the European Directive 98/79/EC, *In vitro* Diagnostic Medical Device Directive, Annex III.

Date of Validity: 17 May 2022

By: Jason Gerhold

Jason Gerhold
Global Director of Quality, Regulatory and Clinical Affairs
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