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* FR2 Assay Kit – MiSeq

Catalog #	Device	Quantity	UDI-DI	GTIN
9-121-0089	LymphoTrack® Dx IGH FR2 Assay Kit A – MiSeq®	40 Reactions	08100227391210089R6	850052003838
9-121-0099	LymphoTrack® Dx IGH FR2 Assay Panel – MiSeq®	120 Reactions	08100227391210099R9	850052003845

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
Global Director of Quality, Regulatory and Clinical Affairs
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