invivoscribe

DECLARATION OF CONFORMITY

Manufacturer:

Invivoscribe, Inc.
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United States of America

Authorized Representative:

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Family Name: LymphoTrack® Dx

Device Trade Name: *IGH* FR1/2/3 Assay Kit – MiSeq®

Catalog #	Device	Quantity	UDI-DI	GTIN
9-121-0129	LymphoTrack® Dx IGH FR1/2/3 Assay Kit A – MiSeq®	40 + 40+ 40 Reactions	08100227391210129QR	00850052003876
9-121-0139	LymphoTrack® Dx IGH FR1/2/3 Assay Panel – MiSeq®	120 + 120 + 120 Reactions	08100227391210139QU	00850052003883

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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