



DECLARATION OF CONFORMITY

Manufacturer: **Invivoscribe, Inc.**
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Family Name: **IdentiClone™**

Device Trade Name: **IGH+IGK B-Cell Clonality Assay**

Catalog #	Device	Quantity	Basic UDI-DI	GTIN
9-100-0031	IGH+IGK B-Cell Clonality Assay for ABI Fluorescence Detection	33 Reactions	08100227391000031P2	00850052003067
9-100-0041	IGH+IGK B-Cell Clonality Assay MegaKit for ABI Fluorescence Detection	330 Reactions	08100227391000041P5	00850052003074

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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