



DECLARATION OF CONFORMITY

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

Device Trade Name: **IGL Gene Clonality Assay**

| Catalog # | Device | Quantity | Basic UDI-DI | GTIN |
|------------|---|---------------|---------------------|----------------|
| 9-103-0011 | IGL Gene Clonality Assay for ABI Fluorescence Detection | 33 Reactions | 08100227391030011PV | 00850052003234 |
| 9-103-0021 | IGL Gene Clonality Assay MegaKit for ABI Fluorescence Detection | 330 Reactions | 08100227391030021PY | 00850052003265 |

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
Invivoscribe, Inc.
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