invivoscribe

DECLARATION OF CONFORMITY

Manufacturer: **Invivoscribe, Inc.**
10222 Barnes Canyon Rd. Bldg 1
San Diego, California 92121
United States of America

Authorized Representative: **Invivoscribe Technologies, SARL**
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Family Name: IdentiClone™

Device Trade Name: **T Cell Receptor Gama Gene Rearrangement**

Catalog #	Device	Quantity
9-207-0101	T Cell Receptor Gamma Gene Rearrangement Assay 2.0	33 Reactions
9-207-0111	T Cell Receptor Gamma Gene Rearrangement Assay 2.0 Megakit	330 Reactions

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: 19 June 2020

By: _____

Jason Gerhold
Global Director of Regulatory, Quality and Clinical Affairs
Invivoscribe, Inc.
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