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: LymphoTrack® Dx

Device Trade Name: ***IGH* FR3 Assay Kits – S5/PGM**

Catalog #	Device	Quantity
9-121-0047	LymphoTrack® Dx *IGH* FR3 Assay – S5/PGM	60 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: 13 Feb 2019

By: _____

Jason Gerhold
Global Director of Regulatory Affairs & Quality Assurance
Invivoscribe Inc.
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