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
Le Forum – Bât B
515 Avenue de la Tramontane
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Family Name: LymphoTrack® Dx

Device Trade Name: ***IGH* FR1 Assay Kit – S5/PGM**

Catalog #	Device	Quantity
9-121-0007	LymphoTrack® Dx *IGH* FR 1 Assay Kit – S5/PGM	60 Reactions

I, the undersigned, hereby declare that the *in-vitro* diagnostic medical device specified above conforms 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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