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: ***IGHV* Leader SHM Assay Kits – MiSeq®**

Catalog #	Device	Quantity
9-121-0059	LymphoTrack® Dx *IGHV* Leader Somatic Hypermutation Assay Kit A – MiSeq®	40 Reactions
9-121-0069	LymphoTrack® Dx *IGHV* Leader Somatic Hypermutation Assay Panel – MiSeq®	120 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: 20 Mar 2019

By: _____

Jason Gerhold
Global Director of Regulatory Affairs & Quality Assurance
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