



## 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:** **TRG Assay Kits – MiSeq**

Catalog #	Device	Quantity
9-227-0019	LymphoTrack® Dx TRG Assay Kit A – MiSeq	40 Reactions
9-227-0009	LymphoTrack® Dx TRG 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: 21 Mar 2019

By: 

Jason Gerhold  
Global Director of Regulatory Affairs & Quality Assurance  
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