



## 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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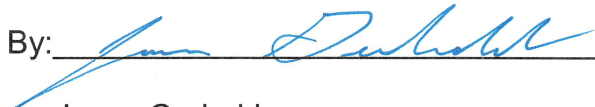
**Family Name:** LymphoTrack® Dx

**Device Trade Name:** **IGH 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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