invivoscribe™

DECLARATION OF CONFORMITY

Manufacturer:

Invivoscribe Technologies, Inc.
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Authorized Representative:

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Family Name: IdentiClone™

Device Trade Name: **IGH + IGK B-Cell Clonality Assay**

Catalog #	Device	Quantity
9-100-0010	IGH + IGK B-Cell Clonality Assay - Gel Detection	33 Reactions
9-100-0031	IGH + IGK B-Cell Clonality Assay - ABI Fluorescence Detection	33 Reactions
9-100-0012	IGH + IGK B-Cell Clonality Assay - Beckman-Coulter CE/NAT	33 Reactions
9-100-0020	IGH + IGK B-Cell Clonality Assay MegaKit - Gel Detection	330 Reactions
9-100-0041	IGH + IGK B-Cell Clonality Assay MegaKit - ABI Fluorescence Detection	330 Reactions
9-100-0022	IGH + IGK B-Cell Clonality Assay MegaKit - Beckman-Coulter CE/NAT	330 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: Nov 2, 2010

By: _____

Jeffrey E. Miller, PhD
CSO and CEO
Invivoscribe Technologies, Inc.
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