InVivoScribe Technologies
www.invivoscribe.com

DECLARATION OF CONFORMITY

Manufacturer:

***In Vivo*Scribe Technologies, Inc.**
6330 Nancy Ridge Drive, Suite 106
San Diego, California 92121
United States of America

Authorized Representative:

***In Vivo*Scribe Technologies, SARL**
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Family Name:

LeukoStrat™

Device Trade Name:

***FLT3* Mutation Assay**

Catalog #	Device	Quantity
9-412-0010	*FLT3* Mutation Assay for Gel Detection	33 Reactions
9-412-0011	*FLT3* Mutation Assay for ABI Fluorescence Detection	33 Reactions
9-412-0012	*FLT3* Mutation Assay for Beckman Coulter CE / NAT	33 Reactions
9-412-0020	*FLT3* Mutation Assay MegaKit for Gel Detection	330 Reactions
9-412-0021	*FLT3* Mutation Assay MegaKit for ABI Fluorescence Detection	330 Reactions
9-412-0022	*FLT3* Mutation Assay MegaKit for 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: Jan 27, 2009

By: _____

Jeffrey E. Miller, PhD
CSO and CEO
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USA