EC DECLARATION OF CONFORMITY

Sequent Medical Inc.
11A Columbia
Aliso Viejo, California 92656 USA

Sequent Medical Inc. declares that according to Directive 93/42/EEC Annex II under our sole responsibility that the products to which this declaration relates are in conformity with 93/42/EEC and fulfill the Essential Requirements as described in Directive 93/42/EEC Annex I.

Conformity Assessment Route: Annex II, Section 3 – Full Quality Assurance System
Annex II, Section 4 – EC Design Examination
Annex II, Section 5 – Surveillance

Product Category and Name	Product Part Number – FG17154-01, FG21154-01, FG27154-01, FG33133-01	MDD Class, Annex and Rule
VIA™ Microcatheter	**VIA™ Microcatheter Model Numbers**	Class III, Annex 9, Rule 7
	VIA-17-154-01 (154cm in length, 0.0175" lumen Diameter)	
	VIA-21-154-01 (154cm in length, 0.021" lumen Diameter)	
	VIA-27-154-01 (154cm in length, 0.027" lumen Diameter)	
	VIA-33-133-01 (133 cm in length, 0.033" lumen diameter)	

Manufacturer Name and Address:
Sequent Medical, Inc.
11A Columbia
Aliso Viejo, CA 92656

Notified Body Design Certificate #: 507073 MRA:
DQS Medizinprodukte GmbH
D-60433 Frankfurt am Main,
Germany
Notified Body Number: 0297

European Representative:
MediMark Europe SARL
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Curtis Hanson
Director Quality Assurance

Aliso Viejo, CA 92656; USA	November 10, 2017	November 6, 2022
Place of Issue	Date of Issue	Expiration Date

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