Declaration of conformity

INTEGRA Biosciences AG – 7205 Zizers, Switzerland

declares on its own responsibility that the devices

Description	Models
VIAFLO 96	**6000, 6001**
VIAFLO 384	**6030, 6031**

comply with:

EU Directives (DoW: Date of Withdrawal)	Before DoW	DoW	After DoW
Low Voltage Equipment	**2006/95/EC**	20.04.2016	**2014/35/EU**
Electromagnetic Compatibility	**2004/108/EC**	20.04.2016	**2014/30/EU**
Restriction of Hazardous Substances	**2011/65/EU**		
Waste Electrical and Electronic Equipment	**2012/19/EU**		

EU Regulations	
Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH)	**1907/2006**

Standards for EU	
Safety requirements for electrical equipment for measurement, control and laboratory use - General requirements.	**EN 61010-1: 2010**
Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes.	**EN 61010-2-81: 2015**
Electrical equipment for measurement, control and laboratory use - EMC requirements.	**EN 61326-1: 2013**

Standards for Canada and USA	
Safety requirements for electrical equipment for measurement, control and laboratory use - General requirements[a].	**CAN/CSA-C22.2 No. 61010-1**
Safety requirements for electrical equipment for measurement, control and laboratory use - General requirements[a].	**UL 61010-1**
Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes[a].	**UL 61010-2-81**
Operation is subject to the following two conditions: (1) this device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.	**Part 15 of the FCC Rules Class A**

[a]NRTL certificate number (TÜV Süd): U8 17 05 42035 007

Zizers, June 19, 2017

E. Morscher

Elmar Morscher
CEO

T. Neher

Thomas Neher
Quality Manager