Declaration of conformity

INTEGRA Biosciences AG – 7205 Zizers, Switzerland

declares on its own responsibility that the devices

<table>
<thead>
<tr>
<th>Description</th>
<th>Models</th>
</tr>
</thead>
<tbody>
<tr>
<td>VIAFLO 96</td>
<td>6000, 6001</td>
</tr>
<tr>
<td>VIAFLO 384</td>
<td>6030, 6031</td>
</tr>
</tbody>
</table>

comply with:

**EU Directives**

- Low Voltage Equipment 2014/35/EU
- Electromagnetic Compatibility 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 requirementsa. CAN/CSA-C22.2 No. 61010-1
- Safety requirements for electrical equipment for measurement, control and laboratory use - General requirementsa. UL 61010-1
- Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposesa. 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

aNRTL certificate number (TÜV Süd): U8 17 05 42035 007

Zizers, March 2, 2020

Urs Hartmann
CEO

Thomas Neher
Quality Manager