INTEGRA Biosciences AG – 7205 Zizers, Switzerland

declares on its own responsibility that the devices

<table>
<thead>
<tr>
<th>Description</th>
<th>Models</th>
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<tbody>
<tr>
<td>MEDIAJET</td>
<td>103005, 103006</td>
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<tr>
<td>MEDIAJET vario</td>
<td>113000, 113001, 113002</td>
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</tbody>
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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 laboratory use 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 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 UL 61010-1
- Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes. 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

Zizers, February 20, 2020

Urs Hartmann  
CEO

Thomas Neher  
Quality Manager