

## Declaration of Conformity

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<b>Manufacturer</b>	Penlon Limited, Abingdon Science Park, Barton Lane, Abingdon, OX14 3NB, UK
<b>Product Name</b>	Suction Controllers <ul style="list-style-type: none"><li>• SC760 High, Low, Twin Suction Controllers</li><li>• Intermittent Suction Controller</li><li>• East Range High Suction; Low Suction or Twin Suction Controllers</li><li>• Suction Controller Kit</li></ul> Accessories <ul style="list-style-type: none"><li>• East Receiver Jar Range (jar and / or cradle)</li><li>• V Socket</li><li>• Bactitrap</li></ul>
<b>Classification</b>	Class I Rule 2
<b>Conformity Assessment Route</b>	Annex VII
<b>GMDN Code</b>	36778
<b>Quality Management System</b>	ISO 13485:2016
<b>Start of CE Marking</b>	1999

We hereby declare that the above mentioned products meet the provisions of the Council Directive 93/42/EEC for Medical Devices, as amended by Directive 2007/47/EC. All supporting documentation is retained at the premises of the manufacturer.

**Place and Date of Issue** Abingdon, 5<sup>th</sup> March 2020

**Signed by:**



SIGNATURE

**Mary Ryan**

PRINT NAME

**Director of Quality Assurance and Regulatory Affairs**

POSITION