

## Declaration of Conformity

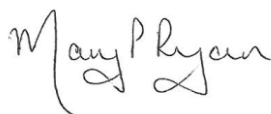
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<b>Manufacturer</b>	Penlon Limited, Abingdon Science Park, Barton Lane, Abingdon, OX14 3NB, UK
<b>Product Name</b>	Prima 300 Series Anaesthetic Machines
<b>Models</b>	Prima 320 Anaesthetic Machines Prima 320 Advance Anaesthetic Machines
<b>Classification</b>	Class IIb, Rule 11
<b>Conformity Assessment Route</b>	Annex II
<b>GMDN Code</b>	37710
<b>Quality Management System</b>	ISO 13485:2016
<b>Notified Body</b>	SGS Belgium NV, Noorderlaan 87, BE-2030 Antwerpen, Belgium (ID Number: 1639)
<b>CE Certificate Number</b>	GB20/965349
<b>Start of CE Marking</b>	4 <sup>th</sup> July 2017
<b>European Authorised Representative</b>	Obelis s.a. Bd. Général Wahis 53, 1030-Brussels, Belgium

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, 13<sup>th</sup> February 2023

**Signed by:**



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SIGNATURE

**Mary Ryan**

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PRINT NAME

**Director of Innovation, Technology & Regulatory Affairs  
EU PRRC**

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POSITION