Penlon

Declaration of Conformity

Manufacturer	Penlon Limited, Abingdon Science Park, Barton Lane, Abingdon, OX14 3NB, UK
Product Name	Penlon Patient Monitor Range
Models	SP M5 SP M8 AnaVue 4000
Classification	Class IIb, Rule 10
Conformity Assessment Route	Annex II
GMDN Code	33586
Quality Management System	ISO 13485:2016
Notified Body	SGS Belgium NV, Noorderlaan 87, BE-2030 Antwerpen, Belgium (**ID Number:** 1639)
CE Certificate Number	GB20/965349
Start of CE Marking	2006

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, 5th March 2020

Signed by:

Mary P Ryan

SIGNATURE

Mary Ryan

PRINT NAME

Director of Quality Assurance and Regulatory Affairs

POSITION