Doc Title:	Rythmic™ PN.PN+.Perf.Perf+_Declaration Of Conformity MDD							
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Item Name	Rythmic™ PN.PN+.Perf.Perf+ Model						
Item Code:	12000	Item Version:	5	Item ERP No.:	N/A	Item Level:	Model

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1 Document Approval - Electronic Signatures

Christos Freris;Document Control Review;Friday, June 5, 2020 11:41:28 AM EEST
Achilleas Tsoukalis;Review;Wednesday, June 17, 2020 6:35:02 PM EEST
Sasa Karpeti;Final Approval;Thursday, June 18, 2020 11:41:56 AM EEST
Christos Freris;Final Approval;Monday, June 22, 2020 12:31:52 PM EEST

2 Declaration of Conformity

We,

the medical device manufacturer,

Micrel Medical Devices S.A.
42 Konstantinoupoleos Str.
Koropi/Athens GR-19441
GREECE

declare under our sole responsibility

that the products listed in the document

"**ML-PR-12018_Rythmic™ PN.PN+.Perf.Perf+_Registration_Item List**"

- are in conformity with the essential requirements and principles for safety and performance of the

  Medical Devices Directive 93/42/EEC ("MDD") amended by Council Directive 2007/47/EC

- are classified Class IIb according to the Annex IX, rule 11, of the above directive

- are CE certified according to the conformity assessment and supervision of the

  Notified Body SGS Belgium NV (CE1639),
  SGS House, Noorderlaan 87, 2030 Antwerp, Belgium

This declaration is signed electronically and valid until 2024-05-24,
the expiry date of the EC Full Quality Assurance System Certificate BG19/871877.