

Federal Communication Commission
 Equipment Authorization Division, Application Processing Branch
 7435 Oakland Mills Road
 Columbia, MD 21048

Certification and Engineering Bureau
 Innovation, Science and Economic Development Canada
 Spectrum Engineering Branch
 3701 Carling Avenue, Building 94
 Ottawa, Ontario K2H 8S2

Philips Medizin Systeme Böblingen GmbH
 Regulatory and Clinical Affairs
 Monitoring & Analytics and Therapeutic Care

+49 151 / 1961 5514
 +49 7031 / 463-2442
 Stefan.Breuer@philips.com

Subject:

MODULAR APPROVAL STATEMENT

Date: 01 February 2019

FCC Certification Number: PQC-NFCBV1

Only applicable for ISED certification:

ISED Company Number: 3549C

UPN: NFCBV1

HVIN: NFCBV1-A

PMN: NFCBV1-A

(Hardware Version Identification Number)

(Product Marketing Name)

HMN: N/A

FVIN: N/A

(Host Marketing Name)

(Firmware Version Identification Number)

To whom it may concern:

Pursuant to Annex D in RSP-100 Issue 11 January 2016 and CFR § 15.212, we herewith declare for our module:

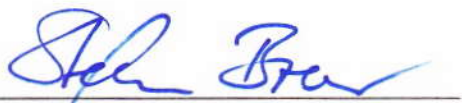
Modular approval requirement	Yes	No *
(a) The radio elements must have the radio frequency circuitry shielded. Physical/discrete and tuning capacitors may be located external to the shield, but must be on the module assembly.	✓	
* Please provide a detailed explanation if the answer is "No": N/A		
(b) The module shall have buffered modulation/data input(s) (if such inputs are provided) to ensure that the module will comply with the requirements set out in the applicable RSS / part 15 under conditions of excessive data rates or over-modulation.	✓	
* Please provide a detailed explanation if the answer is "No": N/A		



Modular approval requirement	Yes	No *
(c) The module shall have its own power supply regulation on the module. This is to ensure that the module will comply with the requirements set out in the applicable standard regardless of the design of the power supplying circuitry in the host device which houses the module.	✓	
* Please provide a detailed explanation if the answer is "No": N/A		
(d) The module shall comply with the provisions for external power amplifiers and antennas detailed in this standard. The equipment certification submission shall contain a detailed description of the configuration of highest antenna gain for each type of antenna.	✓	
* Please provide a detailed explanation if the answer is "No": N/A		
(e) The module shall be tested for compliance with the applicable standard in a stand-alone configuration, i.e. the module must not be inside another device during testing.	✓	
* Please provide a detailed explanation if the answer is "No": N/A		
(f) The module shall comply with applicable RSS-102 exposure requirements and any applicable FCC RF exposure requirement which are based on the intended configuration/integration in a host.	✓	
* Please provide a detailed explanation if the answer is "No": N/A		
Only applicable for FCC certification:		
(g) The module must be equipped with either a permanently affixed label or must be capable of electronically displaying its FCC identification number.	✓	
* Please provide a detailed explanation if the answer is "No": N/A		
(h) The modular transmitter complies with all applicable FCC rules. Instructions for maintaining compliance are given in the user instructions.	✓	

If you have any questions, please feel free to contact us at the address shown below.

Best Regards,



Stefan Breuer
Senior Regulatory Affairs Engineer

Philips Medizin Systeme Böblingen GmbH
Hewlett-Packard-Str. 2, 71034 Boeblingen, Germany
Tel. +49 151 1961 5514

INFO for applicant: LMA may be granted when **one or more** of the requirements in the table above cannot be demonstrated. LMA will also be issued in those instances where applicants can demonstrate that they will retain control over the final installation of the device, such that compliance of the end product is assured. In such cases, an operating condition on the LMA for the module must state that the module is only approved for use when installed in devices produced by a specific manufacturer.

When LMA is sought, the application for equipment certification must specifically state **how control of the end product**, into which the module will be installed, will be maintained, such that full compliance of the end product is always ensured.