

DECLARATION OF CONFORMITY (DoC)

Equipment: Electronic Stethoscope
Trademark(s) and Model(s): iMED+ / CaRDiaRT
Manufacturer: IMEDIPLUS INC.
FCC ID in case other parts of this equipment are subject to certification: 2AM7NDS3011A

This device complies with Part 15 of the FCC Rules.

Operation is subject to the following two conditions:

- (1) **this device may not cause harmful interference, and**
- (2) **this device must accept any interference received, including interference that may cause undesired operation.**

The following test reports are subject to this declaration:

Test report number: Issue date:
EM/2017/40061 2017/08/01

The following manufacturer/importer/entity is responsible for this declaration:

Company name: IMEDIPLUS INC.
Name/Title (legal representative): CHEN,CHIH-HAO/Special Assistant
Address: 2F, 12 Sheng Yi Rd. Sec. 2, Chupei City, Hsinchu
County 302, Taiwan (R.O.C).
Phone: +886-3-6587700 ext.102
Fax: +886-3-6589535
E-mail: elio.chen@imediplus.com
Date: 2017/08/10
Signature: *Chen Chih-Hao*