FEDERAL COMMUNICATIONS COMMISSION

DECLARATION OF CONFORMITY (DoC)

Equipment:	Elecronic stethoscope
Trademark(s) and Model(s):	iMED+ / DS101 , Omni-Steth
Manufacturer:	IMEDIPLUS INC.
FCC ID in case other parts of this equipment are subject to certification:	2AM7NDS101

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:
MW/2018/10049	**Feb. 21, 2018**

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.602
Fax:	+886-3-6589535
E-mail:	elio.chen@imediplus.com
Date:	2018/03/05

Signature:

