Declaration of Conformity

Manufacturer: **ViVest Medical Technology Co., Ltd.**
Unit 205, 206, 207, B1 Building, SIP BioBay, No.218, Xinghu Street, Suzhou Industrial Park, Suzhou Area, Suzhou Free Trade Pilot Zone, 215213, Jiangsu, P.R.China

European Representative: **Shanghai International Holding Corp. GmbH (Europe)**
Eiffestrasse 80, 20537 Hamburg, Germany

Product Name: Automatic External Defibrillator
Model Number: PowerBeat X1 & PowerBeat X3
UMDNS/GMDN Code: 37805
Classification (MDD, Annex IX): IIb, rule 9
Conformity Assessment Route: Annex II.3

We hereby declare that the above mentioned products meet the transposition into national law, the provisions of the following EC Council Directives and Standards. All supporting documentations are retained under the premises of the manufacturer. We are exclusively responsible for the Declaration of Conformity.

DIRECTIVES

General applicable directives:

Medical Device Directive: COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices (MDD 93/42/EEC). Amended by DIRECTIVE 2007/14/EC

Notified Body: **TÜV SÜD Product Service GmbH, Ridlerstr. 65, 80339 MÜnchen, Germany**
Identification number: **CE0123**
(EC) Certificate(s): /
Expire date of the Certificate: /
Start of CE Marking: /
Place, Date of Issue: /

Signature: _____

Name: 纪悦玲/Ji Yueling

Position: Management Representative