

DECLARATION OF CONFORMITY

For the following equipment:

Type of pressure equipment: Vessel


Product Name: Pressure Vessel for Autoclaves

Chamber size(mm):

Cylinder type: $\phi 200$, $\phi 230$, $\phi 260$, $\phi 300$, $\phi 380$, $\phi 400$, $\phi 450$, $\phi 500$, $\phi 610$, $\phi 700$, and $\phi 900$

Rectangle type: 410x410, 510x510, 610x610, 610x915, and 774x774

is herewith confirmed to comply with the requirements set out in the Council Directive on the harmonization of the Laws of the Member States concerning **Pressure Equipment Directive 2014/68/EU** with the compliance of conformity assessment of **Module H**. The notified body is **DNV GL Business Assurance**

Italia S.r.l. (number –  0496).

The following **standards** are applied:

ASME BPVC VIII.1:2015	ASME Boiler & Pressure Vessel Code 2015 Edition VIII RULES FOR CONSTRUCTION OF PRESSURE VESSELS Division 1
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Fluid group: 2

Category: I, II, and III

Number of certificate: 4028-2013-CE-RGC-ACCREDIA

The following European Authorized Representative is stated to the declaration:

APEX MEDICAL S.L.

Elcano 9, 6^a planta 48008 Bilbao. Vizcaya SPAIN

The EU declaration of conformity is issued under the sole responsibility of the manufacturer.

The following person is responsible for the compliance of declaration:

STURDY INDUSTRIAL CO., LTD.

Mr. Wolfgang Huang

Quality Manager

Taipei, Taiwan



23 February, 2018

Name and position

Legally signature

Date