DECLARATION OF CONFORMITY
(Certificate of Quality)

Manufacturer:	**Innovision ApS** Skovvænget 2 DK-5620 Glamsbjerg Denmark Tel.: +45 65 95 91 00 Fax: +45 65 95 78 00
Conformity assessment procedure:	Annex II of the Medical Device Directive 93/42/EEC
Certificate(s):	Presafe Denmark A/S, EC Certificate Number DGM-859 Design, manufacture and final inspection of cardiopulmonary function test equipment in class IIa Initial date of issue: 2015-07-06 Notified Body, Identification No. 0543
Product:	Innocor Cardiopulmonary function test equipment Class: IIa Models: Innocor with options SpO2 + O2 + BbB + LCI (INN00010,INN00050,INN00100,INN00400)

We, the manufacturer hereby declare that this product is in conformity with the requirements in the Council Directive 93/42/EEC concerning medical devices as amended and transposed into Danish law.

Signed in Glamsbjerg:	2015-07-06
Name and authority:	Gert Røge, Quality Assurance Manager
Signature: