

**EC Declaration of Conformity**  
**EG Konformitäts-Erklärung**  
**Déclaration CE de Conformité**  
**Declaración de Conformidad tipo EC**

We, Datrend Systems Inc, declare in exclusive responsibility that the product:  
Wir, Datrend Systems Inc, erklären in alleiniger Verantwortung, daß das Produkt:  
Nous, Datrend Systems Inc, déclarons sous notre seule responsabilité que le produit:  
Datrend Systems Inc declaramos en la responsabilidad exclusiva que el producto:

Model / Modell / Modèle / Modelo: **vPad-A1 Modular Patient Simulator<sup>1</sup>**

to which this declaration relates is in conformity with the following Directives:  
auf die sich diese Erklärung bezieht, steht im Einklang mit den folgenden Richtlinien:  
auquel se réfère cette déclaration est conforme aux Directives suivantes:  
a cuál se relaciona este declaración es conforme a las siguientes Directivas:

**2014/30/EU - Electromagnetic Compatibility (EMC)**  
**2014/35/EU - Low Voltage Directive (LVD)**

Harmonized standards used:  
Verwendete harmonisierte Standards:  
Normes harmonisées utilisées:  
Normas armonizados utilizados:

**EN 61326-1:2013** *Electrical equipment for measurement, control and laboratory use - EMC requirements*  
**EN 61010-1:2010** *Safety requirements for electrical equipment for measurement, control and laboratory use*

Signed for, and on behalf of, Datrend Systems Inc.



Lloyd Van't Haaff  
Quality Manager

Date of Declaration: 2016 August 22

Place of Issue: Richmond BC Canada

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<sup>1</sup> Comprising in combination the A1 Base Module and any one or more of the following simulation modules: vPad-PS Patient Simulator; vPad-O2 Pulse Oximeter Tester; and/or vPad-NIBP Non-Invasive Blood Pressure Simulator