

UK Declaration of Conformity

130 - 4020 Viking Way Richmond, BC V6V 2L4 CANADA

We,

Datrend Systems Inc.

declare in exclusive responsibility that the product:

Model: vPad-A1 Modular Patient Simulator1

is in conformity with the essential requirements and other relevant requirements of:

Restriction of the Use of Certain Hazardous Substances in Electrical and Electronic Equipment Regulations 2012

Electromagnetic Compatibility Regulations 2016

Electrical Equipment (Safety) Regulations 2016

Designated standards:

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

EN 63000:2018 Technical documentation for assessment of electrical and electronic

products with respect to restriction of hazardous substances

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

Keith Ching

Director of Manufacturing Technology

Date of Original Declaration: 2022-Jul-21

Date of Declaration: 2025-May-1 Place of Issue: Richmond BC Canada

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