

**MANUFACTURER'S DECLARATION: MEDICAL DEVICES REGULATION 2017/745**

As this component is not a complete "Medical Device" under the definition provided in the Regulation but a part with no function on its own, a Declaration of Conformity or CE mark is not appropriate until the component is incorporated. The following Declaration is issued in order to aid the installation of the component below into a finished appliance, for use by, or sale to, an end-user.

The following declares the status of these products with regard to Union Regulation 2017/745:

Product Description	Part Numbers
VR2 Lighting Module	D51234
VSI Lighting Module	D50242
VR2 LED Lighting Module	D51475

The object of the declaration described above is in conformity with the relevant Union harmonisation legislation to the extent possible as a component to be integrated into a complete product.

**APPLICABLE STANDARDS**

The following standards (and those called by them) have been used in order to assess a presumption of conformity with the essential requirements of the above regulation as far as the component allows:

EN 14971:2019 + A11:2021	Medical devices — Application of risk management to medical devices
EN 62304:2006 + A1:2015	Medical device software — Software life-cycle processes
EN 10993-1:2021	Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process
EN 12184:2022	Electrically powered wheelchairs, scooters and their chargers. Requirements and test methods
ISO 7176-9:2009	Wheelchairs -- Part 9: Climatic tests for electric wheelchairs
ISO 7176-14:2022	Wheelchairs -- Part 14: Power and control systems for electrically powered wheelchairs and scooters -- Requirements and test methods
ISO 7176-21:2009	Wheelchairs -- Part 21: Requirements and test methods for electromagnetic compatibility of electrically powered wheelchairs and scooters, and battery chargers

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Nigel Mills, Senior Manager, Engineering

Signed at, for and on behalf of:

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