Keeler

Ophthalmic Instruments

CE

Declaration of Conformity

We declare that the following product(s) conform to the requirements of the Medical Device Directive 93/42/EEC and are classified as **2a** device(s) in accordance with Annex IX of the Directive. Conformance is assured according to the requirements stated in Annex II of the Directive.

Product Description
Tonometer, line-powered GMDN Code 35399 Type P

Product Description	Part No	Variant
Tonometer	2414-P-2001	Pulsair Intellipuff

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Nickie Power
Quality & Regulatory Compliance Engineer

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