BeamMed
Focused On Bone Health

CE DECLARATION OF CONFORMITY
European Medical Device Directive 93/42/EEC

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I, the undersigned, hereby declare that the distributed CE marked products, specified in the annexed product list, conform to the product(s) covered by the "CE Marking of Conformity Certificate", reference number 93741CE01, firstly issued on July 1, 2001 and delivered by DEKRA Certification Meander 1051, 6825 MJ Arnhem, the Netherlands, Notified Body Identification Number 0344, in accordance with Annex II of the Council Directive 93/42/EEC of 14 June 1993, concerning medical devices,

This declaration is supported by the Quality System certificate for the products concerned, in accordance with Annex II of the EC-Directive. The conformity to quality assurance sets out in the said ISO 13485:2016 Conformity Certificate number 2078798, issued and delivered by DEKRA Certification B.V. Arnhem, the Netherlands.

This declaration is valid until: 1 March 2024 Expiration date of CE Marking of Conformity Certificate

Date: 24-Jan-2022

Signature: _____

Tal Marom,
CEO
BeamMed Ltd.

Annex : Product List (document identification)

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ANNEX
PRODUCT LIST

This product list belongs to the Declaration of Conformity identified by BeamMed Ltd. products and specifies the CE marked products concerned that BeamMed Ltd. products intends to distribute in conformity with the provisions of the Council Directive 93/42/EEC of 14 June 1993 concerning medical devices. The following list identifies the products by Model No. and type.

Category: Ultrasound diagnostic system:

Ultrasonic Bone Sonometer units and Ultrasound Probes:

Product Name	Models	Class	Placed on the market
Sunlight MiniOmni	S / P	IIa	2010
Ultrasound probe CM		IIa	09/2006
Ultrasound probe CS		IIa	09/2006
Ultrasound probe CR		IIa	09/2006