EC Declaration of Conformity

Manufacturer:	*Innovation Technologies doo*
SRN:	SI-MF-000041369
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Postcode:	*8290*
City:	*Sevnica*
Country:	*Slovenia*
Telephone number	*+(389)70205737*

We declare that the EC Declaration of Conformity is issued under our sole responsibility and belongs to the following product:

Product name:	*ViewECG (Version 1.2.0)* *Medical device software*
Product device models (chapters):	*ViewECG medical device software*
Basic UDI-DI:	*383008599001Z8*
MDR code:	*MDA 0315; MDS 1010*
EMDN code:	*Z12050392 for ViewECG*
Classification:	*IIa by Annex VIII, Rule 11 of MDR 2017/745*

The product was subject to the conformity assessment procedure outlined in Annex IX — Conformity assessment based on a quality management system and evaluation of technical documentation, as per MDR 2017/745.
The company has implemented a quality management system in accordance with the requirements of the standard EN ISO 13485:2016 and A11:2021.

This product has fulfilled the requirements of the Medical Device Regulation (EU) MDR 2017/745 of the European Parliament and of the Council.

Standards applied:
EN ISO 13485:2016 + A11:2021; EN ISO 14971:2019 + A11:2021; EN ISO 15223-1:2021 + A1:2025; EN 60601-1-6:2010 + A1:2015 + A2:2021; EN 60601-2-47:2015; EN 62304:2006 + A1:2015; EN 62366-1:2015; IEC 81001-5-1:2021; IEC/TR 80002-1:2009; ISO/TR 80002-2:2017

Notified Body:
SIQ, Mašera – Spasićeva 10, SI-1000, Ljubljana, CE 1304

EC Certificate:
Nr: MDR-0045, Issued: 30.06.2026, Valid until: 29.06.2031

Signed for and on behalf of:

Sevnica, Slovenia	**30.06.2026**	
Place of issue	*Date of issue*	**Marjan Gushev, CEO**