



# EC Declaration of Conformity



according to the Medical Devices Directive 93/42/EEC

*Class I Medical Device*

*(non-sterile, without measuring function)*

**Manufacturer:** POD ACTIVE Pty. Ltd.

**Address:** Level 3, 115 Myers Street  
Geelong, Victoria 3220  
AUSTRALIA

**EC Representative:** Wellkang Ltd ([www.CE-marking.eu](http://www.CE-marking.eu))  
Enterprise Hub, NW Business Complex,  
1 Beraghmore Rd., Derry, BT48 8SE, Northern Ireland, UK.

EC Rep SRN in EUDAMED: XI-AR-000001836.

**We, the manufacturer, declare under our sole responsibility that**

	Product Name	Type/model, identification of product allowing traceability (Where applicable)
the medical device(s)	Please refer to attached list	
of class	according to annex IX of directive 93/42/EEC	Class I Medical Device (non-sterile, without measuring function)

is/are in conformity with the relevant provisions and essential requirements of directive 93/42/EEC, as amended by Directive 2007/47/EC.

Conformity assessment procedure	Module A (EC Declaration of Conformity (Annex VII) + Technical Files)
Notified Body (name & number)	NOT applicable

Certificate & number

Signed on: **Friday, 14 May 2021.** Place: **Geelong, Victoria, AUSTRALIA**

Signature (on behalf of the manufacturer)

Name of authorized signatory: **Brett Nicholas**

Position held in the company: **CEO**

**POD Active Pty. Ltd.**  
**Medical Device Registration | May 2021**

**Product List**

<b>Product Name (Device Make)</b>	<b>Model</b>	<b>Class**</b>	<b>Directive* (Regulation)</b>
POD K4 2.0 KNEE BRACE	POD K4 2.0 KNEE BRACE	1	MDR
POD K8 2.0 KNEE BRACE	POD K8 2.0 KNEE BRACE	1	MDR
CTi MISSION BRACE	CTI-3001 CTI-2001 CTI-3002 CTI-2002 CTI-3003 CTI-2003 CTI-3004 CTI-2004 CTI-3005 CTI-2005 CTI-3001-PL CTI-2001-PL CTI-3002-PL CTI-2002-PL CTI-3003-PL CTI-2003-PL CTI-3004-PL CTI-2004-PL CTI-3005-PL CTI-2005-PL	1	MDR
POD A3 ANKLE BRACE	POD A3 ANKLE BRACE	1	MDD
DonJoy Performance POD® Ankle Support Brace	DonJoy Performance POD® Ankle Support Brace	1	MDD

\* The applicable directive must be either **MDD** or **IVDD**.

\*\* Under the **MDD** (i.e. directive 93/42/EEC), the class must be either **I** (Class I, non-sterile, without measuring function), **Is** (Class I, sterile), **Im** (Class I, with measuring function), **Ila**, **Ilb** or **III**, and/or **future UK relevant legislation**

\*\* Under the **IVDD** (i.e. directive 98/79/EC), the class must be either **List A**, **List B**, **Self-Test**, or **Others**, and/or **future UK relevant legislation**

The classification is based on the claim of the manufacturer and is under the sole responsibility of the manufacturer.