Medical Devices Regulation (EU) 2017/745 (MDR)

Declaration of Conformity acc. Annex IV

For

Class 1 Medical Devices

Manufacturer's name:	Van Raam
Manufacturer's address:	Guldenweg 23 7051 HT Varsseveld The Netherlands
Product:	Therapy tandem
Device Name:	Kivo-Plus
Part number:	377-XXXX

I, the undersigned, hereby declare that the equipment specified above, complies with the Essential Requirements of Medical Devices Regulation (EU) 2017/745 (MDR). The Products are labelled with the CE-Mark according to the above Directive.

This product also complies with the Machine Directive 2006/42/EG.

Varsseveld, 25-03-2020

Place, Date of Issue

JW Boezel

Signature, Name, Position