MEDICARE UITGEEST B.V.

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Uitgeest, December 8 2016

Declaration of Conformity medical devices

We hereby declare that the distributed CE-marked products, specified in the product list, are covered by the 'CE Marking of Conformity Certificate' with reference 2110398CN re-issued on February 1 2016 by the notified body: DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem, Netherlands, with notified body identification number 0344.

The products conform to the required technical documentation in accordance with annex VII of the Dutch law "Besluit Medische Hulpmiddelen" transposing the European Community Directive as the Council Directive 93/42/EEC of 14th June 1993 and Directive 2007/43/EG on medical devices.

We declare that the CE marked products mentioned in the certificate meet the applicable provisions of the European Community Directive for Class IIa devices.

This declaration is based on the application of the Quality System approved for the manufacturing and final inspection of the products listed in accordance with Annex V of the European Community Directive for medical devices. Conformity of Product quality assurance stated in Annex V is confirmed by the CE Marking of Conformity certificate as issued by DEKRA Certification B.V.

The following standards have been used for the products below:
NEN-EN-ISO 9170-1:2008 Terminal units for medical gas pipeline systems - Part 1: Terminal units for use with compressed medical gases and vacuum

Product list:

Name	Model	Serialnumbers and higher
Medicare gasafnamepunten voor medische gassen	Starting with model number 0200xxxx	0209720

F. Wilms
Managing Director