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CE

Declaration of Conformity

according to annex VII of the council directive 93/42/EEC

We, the company VAKUFORM GMBH, Weiterstädter Landstrasse 13, 64291 Darmstadt, hereby declare on our own responsibility that the

Class 1 medical devices (Rule N of annex IX-93/42/EEC)

VAKUFORM PRO Seat (Order no. 11000)
VAKUFORM PRO Seat XXL (Order no. 11001)
VAKUFORM PRO Back (Order no. 11010)
VAKUFORM PRO Back XXL (Order no. 11011)
VAKUFORM PRO Anatomical seat (Order no. 11100)
VAKUFORM PRO Anatomical back (Order no. 11110)

(Products according to product description in annex A of our technical documentation)

are manufactured in accordance with the technical documentation and comply with the Essential Requirements of Medical Devices Directive 93/42/EEC (annex I). The products are labeled with the CE-Mark according to the above directive.

Darmstadt, 11.05.2016

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