8 Appendix

8.1 EC Declaration of Conformity

EC Declaration of Conformity

Document No. / Month.Year: QKE 12.20.002-09 / 12.08

Manufacturer: Medset Medizintechnik GmbH
Curslacker Neuer Deich 66
D-21029 Hamburg

We hereby declare that the product:

Long-term blood pressure measurement system
SCANLIGHT

consisting of:

ABPM monitor
SCANLIGHT III Recorder
from version A

and the ABPM evaluation software
SCANLIGHT
from version 1.7c

with the following accessories:
recorder bag, SCANLIGHT II/III
PC interface L, SCANLIGHT II/III
PC interface U, SCANLIGHT II/III
blood pressure cuff S, small, SCANLIGHT II/III
blood pressure cuff M, medium, SCANLIGHT II/III
blood pressure cuff X, large, SCANLIGHT II/III
blood pressure cuff XL, extra large, SCANLIGHT II/III

complies with European Directive 93/42/EEC of the Council of 14th June 1993 concerning medical products.

It bears the mark:

CE 0124

Hamburg, 1st December 2008

Medset Medizintechnik GmbH

Legally binding signature:

Klaus Kophstahl
- Managing director -