



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 13 08 85349 002

Manufacturer: A&D Company, Limited
1-243 Asahi, Kitamoto-shi
Saitama-ken
364-8585 JAPAN

EC-Representative: A&D INSTRUMENTS LIMITED
Unit 24/26 Blacklands Way
Abingdon Business Park
Abingdon, Oxon
OX14 1DY
UNITED KINGDOM

Product Category(ies): Blood Pressure Monitor,
Blood Pressure Analyzing Software

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: TAQ235011283A

Valid from: 2013-11-01

Valid until: 2018-10-31

Hans-Heiner Junker

Date, 2013-10-23

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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Facility(ies):

A&D Company, Limited
1-243 Asahi, Kitamoto-shi, Saitama-ken, 364-8585
JAPAN

KENSEI KOGYO Co., Ltd
4210-15 Takasai, Shimotsuma-shi, Ibaraki-ken,
304-0031 JAPAN

A&D Company, Limited Tokai Division
9-19 Takaramachi, Tajimi-shi, Gifu-ken, 507-0054
JAPAN