TÜVRheinland®

EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60089448 0001

Report No.: 31291105 001

Manufacturer:
W.A. Baum Co., Inc.
620 Oak Street
Copiague, NY 11726
USA

Products:
Sphygmomanometer - manual aneroid type
Sphygmomanometer - manual mercury gravity type for professional use only

Expiry Date: 2018-10-15

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2013-10-16

Date: 2013-10-16

Notified Body

TÜV Rheinland LGA Products GmbH Zertifizierstelle TÜVRheinland

Jürgen Welte

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.