TÜVRheinland®

APPROVAL

EC Directive 93/42/EEC; Annex V, Article 3

Quality Assurance System Production

Registration No.: OD 69241029 0001

Report No.: 28212078 001

Manufacturer: TensioMed Tudományos Informatikai és Orvos-Elektronikai Kft.
Kőér u. 2/e.; H-1103 Budapest,
Hungary

Scope: Manufacturing of blood pressure measuring devices and non-invasive haemodinamical diagnostic devices
Products: TensioClinic arteriograph
TensioDay blood pressure monitor
TensioDayPlus blood pressure monitor
Arteriograph24 blood pressure and arterial function monitor
Arteriograph software
TensioWin software

Date of expiry: 2016-07-11

The Notified Body hereby authorizes the quality management system established and applied by the company mentioned above. The requirements of Annex V, Article 3 of the directive have been met. This approval is subject to periodic surveillance, defined by Annex V, Article 4 of the aforementioned EC Directive, and can be used by the company with the manufacturer's Declaration of Conformity.

Notified Body

TÜV Rheinland InterCert Kft. TÜVRheinland 5 Notified Body 1008

Balázs Bozsik
Certifier

Budapest, 2011-07-12

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Notified under No. 1008 to the EC Commission.

The CE marking may be used if all relevant and effective EC Directives are complied with.