



CERTIFICATE



## EC Certificate

### Full Quality Assurance System according to Medical Devices Directive 93/42/EEC Annex-II Section 3

Certificate Number: 1984-MDD-19-602

We hereby declare that an examination of the under mentioned full quality assurance system has been carried out following the requirements of the national legislation to which the undersigned is subjected, transposing annex II (with the exemption of section 4) of the Directive 93/42/EEC on medical devices. We certify that the full quality assurance system conforms with the relevant provisions of the aforementioned directive.

**Organization:**

**RİTİM GRUP TIBBİ CİHAZLAR ELEKTRONİK  
SANAYİ VE TİCARET ANONİM ŞİRKETİ ANKARA ŞUBESİ**

Yeni Batı Mahallesi 2402 Cad. No:2/2 Yenimahalle, Ankara, Turkey

**Product:** Fully Automatic / Semi Automatic External Defibrillator  
**Model Number:** Aselsan Heartline AED, Ritimport RP3002

The certificate is valid till expiration date, subject to successful completion of periodical surveillance audits. Please contact Kiwa for details.

**Report Number:** M.5447.02  
**Date of first issue:** 25 June 2019  
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**Revision Number:** 01  
**Expiry Date:** 27 May 2024

Kiwa Belgelendirme Hizmetleri A.Ş. is Notified Body under Council Directive 93/42/EEC concerning medical devices with identification number: 1984

Muhteşem Gökhan Yücel  
Head of Notified Body

16 September 2020, Istanbul, Turkey