



EC Certificate

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

Certificate Number: 1984-MDD-11-102

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:

AYGÜN CERRAHİ ALETLER SANAYİ VE TİCARET ANONİM ŞİRKETİ

Kerimbey OSB Mahallesi Yaşardoğu Cad. No:76/1 Tekkeköy, Samsun, Turkey

Products: Electrical and battery operated surgical motor system, High speed surgical motor system, Electrical and pneumatical surgical motor systems, Surgical power system

Products defined at the enclosure which is the part of this certificate and contains one page. The certificate is valid till expiration date, subject to successful completion of periodical surveillance audits. Please contact Kiwa for details.

Report Number:

M.3608.10

Date of first issue:

05 July 2011

Date of last issue:

17 May 2021

Revision Number:

09

Expiry Date:

27 May 2024

17 May 2021, Istanbul, Turkey

Muhteşemm Gökhan Yücel Head of Notified Body





Enclosure of the EC Certificate Page 1/1
Full Quality Assurance System according to
Medical Devices Directive 93/42/EEC Annex-II Section 3
Certificate Number: 1984-MDD-11-102, Revision Number: 09

Concerned medical devices:

Product: Electrical and battery operated surgical motor system

Model Name: Praxis 3, BES Dermatome, Dermatome ST-5, Karınca 206, Karınca

208, Praxis 3B, Raptor SPS 4

Product: High speed surgical motor system **Model Name:** HSM-2, HSM-2 Mini Bone

Product: Electrical and pneumatical surgical motor systems

Model Name: Electrical Control Unit (M6-034), Pneumatic Surgical Motor

Product: Surgical power system

Model Name: Scoren

Kiwa Belgelendirme Hizmetleri A.Ş. is Notified Body under Council Directive

93/42/EEC concerning medical devices with identification number: 1984

17 May 2021, Istanbul, Turkey

Muhteşemm Gökhan Yücel Head of Notified Body

MDCG 2020-3 Change Verification Form



Date:	30.11.2021	
Reference No:	MY-21-002021	

Dear Sir/Madam,

Your following proposed change is reviewed according to the MDCG 2020-3 Guidance on significant changes regarding the transitional provision under Article 120 of the MDR with regard to devices covered by certificates according to MDD or AIMDD.

Information on EC Ce	ertificate	
EC Certificate No:	1984-MDD-11-102	
Revision:	1984-MDD-11-102 Rev. 09 (17 Mayıs 2021)	
Manufacturer:	AYGÜN CERRAHİ ALETLER SANAYİ VE TİCARET ANONİM ŞİRKETİ	
Expiry Date:	27.05.2024	
Information on Propo	sed Change	
Subject of Change:	Electrical and pneumatic surgical motor systems products are removed the from certificate.	

The implementation of the above change proposed by AYGÜN CERRAHİ ALETLER SANAYİ VE TİCARET ANONİM ŞİRKETİ does not represent a significant change in the design and usage purpose within the scope of MDR Article 120(3).

The above-mentioned EC certificate is valid until its expiration date and is subject to the successful completion of periodic surveillance audits.

This letter is valid when presented together with the EC Certificate mentioned above.

Sincerely

Orhan Karakuş Quality Manager