

ATTESTATION CE / EC CERTIFICATE

Approbation du Système Complet d'assurance Qualité / Approval of full Quality Assurance System

ANNEXE II excluant le point 4 Directive 93/42/CEE relative aux dispositifs médicaux

ANNEX II excluding section 4 Directive 93/42/EEC concerning medical devices

Pour les dispositifs de classe III, un certificat CE de conception est requis

For class III devices, a EC design certificate is required

Fabricant / Manufacturer

ECA Medical Instruments

1107 Tourmaline Drive

NEWBURY PARK, CA 91320 UNITED STATES

Catégorie du(des) dispositif(s) / Device(s) category

Instruments à usage unique stériles ou non stériles comprenant : limiteurs de couple brevetés et instruments chirurgicaux destinés à la chirurgie crano-maxillo-faciale, de management du rythme cardiaque, de neuromodulation, cardiovasculaire et orthopédique

Single use proprietary torque-limiting devices and surgical instruments supplied sterile or non-sterile and intended to be used for crano-maxillofacial (CMF), cardiac rhythm management (CRM), neuromodulation, cardiovascular, and orthopedic surgeries

Voir détails sur addendum / See attachment for additional information

Le LNE/G-MED atteste qu'à l'examen des résultats figurant dans le rapport référencé P129290-3 & P129290-R, le système d'assurance qualité - pour la conception, la production et le contrôle final - des dispositifs médicaux énumérés ci-dessus est conforme aux exigences de l'annexe II excluant le point 4 de la Directive 93/42/CEE.

LNE/G-MED certifies that, on the basis of the results contained in the file referenced P129290-3 & P129290-R, the quality system - for design, manufacturing, and final inspection - of medical devices listed here above complies with the requirements of the Directive 93/42/EEC, annex II excluding section 4

La validité du présent certificat est soumise à une vérification périodique ou imprévue

The validity of the certificate is subject to periodic or unexpected verification

Début de validité / Effective date : March 17th, 2016 (included)

Valable jusqu'au / Expiry date : March 16th, 2019 (included)



On behalf of the Certification Director

Cécile VAUGELADE

G-MED Certification Division Manager

Identification des dispositifs / Identification of devices

Nom du dispositif médical <i>Medical device name</i>	Dénomination commerciale <i>Commercial designation</i>	Classe du DM <i>MD Class</i>
Devices Provided Non-sterile		
Axial Handle Unidirectional or bi-directional torque Driver (non-sterile)	212	IIa
Axial Handle Unidirectional or bi-directional torque Driver (non-sterile)	214	
Axial Handle Unidirectional or bi-directional torque Driver (non-sterile)	215	
Devices Provided Sterile and Non-sterile		
Micro Axial Fixed Driver	216	IIa
Triangular Long and Short handle Torque Driver	218	
Medium Axial and Palm Handle Torque Driver	219	
Large Axial and T-Handle Torque Driver	220	
Axial and T-Handle with 1/4 inch square, AO or other coupling Torque Driver	225	
SpeedEcap™ Standard Spin Top Medium Axial Fingertip Swivel Fixed Driver with WickAway® Technology	400	
SpeedEcap™ Crown Top Precision Palm Control and Actuation with WickAway® Technology	401	
Mini Axial Fixed Driver Series	412/414	
T-Handle Fixed Driver	420	
Standard Medium Palm Fixed Driver with WickAway® Technology	430	
Standard Medium Palm Axial awl with WickAway® Technology	440	
Standard Medium Palm Axial Fixed Driver with WickAway® Technology	450	

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On behalf of the Certification Director
Cécile VAUGELADE
G-MED Certification Division Manager

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Ce certificat couvre les activités et les sites suivants :
This certificate covers the following activities and sites:

ECA Medical Instruments
1107 Tourmaline Drive
Newbury Park, CA 91320 - USA

Fabrication et contrôle final / Manufacturing and final Control

ECA Medical Instruments
2193 Anchor Court
Thousand Oaks, CA 91320 - USA

Conception et fabrication / Design and manufacturing

2 sites / 2 sites

LNE/G-MED

0459



On behalf of the Certification Director
Cécile VAUGELADE
G-MED Certification Division Manager

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