



EU DECLARATION OF CONFORMITY

Manufacturer: Duckworth & Kent Ltd
Terence House
7 Marquis Business Centre
Royston Road
Baldock, Hertfordshire
SG7 6XL, United Kingdom

Authorised Representative: Emergo Europe
Prinsessegracht 20
2514 AP The Hague
The Netherlands

Duckworth & Kent Ltd declares that the medical devices listed in Table 1 and described hereafter,

Duckworth & Kent range of Non-Invasive Markers

are in conformity with the essential requirements and provisions of the Council Directive 93/42/EEC as amended by Directive 2007/47/EC and as transposed into UK legislation (UK Medical Devices Regulations SI 2002 No. 618, as amended).

In accordance with Annex VII of the Council Directive 93/42/EEC as amended by Directive 2007/47/EC and as transposed into UK legislation (UK Medical Devices Regulations SI 2002 No. 618, as amended),

These devices are designated as Class I

Classification Route

These devices are all transient use, reusable, non-sterile, non-invasive medical devices.

The classification is determined under Annex IX, Rule 1.

This declaration is issued under the sole responsibility of Duckworth & Kent Ltd.

Martin Lock
Head of Regulatory Affairs



Table 1. Medical Devices covered under this Declaration of Conformity.

Markers, Non-Invasive	
REF	Device Title
6-112	Hook
6-945	Trephine Guide
6-946	Trephine Guide
9-503	Fixation Ring
9-509	Fixation Ring
9-510	Fixation Ring
9-510R	Fixation Ring
9-515R	Fixation Ring
9-518-1	CCC Marker
9-518-2	CCC Marker
9-518-2R	CCC Marker
9-518-3	CCC Marker
9-518-4	CCC Marker
9-544	Injection Guide
9-544-2	Injection Guide
9-544-4	Injection Guide
9-649	Marking Caliper
9-650	Marking Caliper
9-692	Marking Caliper
9-700	Degree Gauge
9-705R	Degree Gauge
9-705R-1	Degree Gauge
9-729	Blade Marker
9-729-1	Blade Marker
9-730	Blade Marker
9-732	Blade Marker
9-733	Blade Marker
9-734	Blade Marker
9-749	Incision Marker
9-778	Ring Marker
9-779	Ring Marker
9-781	Ring Marker
9-781W	Ring Marker
9-788	Ring Marker
9-789W-1	Ring Marker
9-830	S Marker
9-831	F Marker
9-840	Toric Marker
9-840-1	Toric Marker
9-840-3	Toric Marker
9-841	Toric Marker
9-841-1	Toric Marker
9-841-2	Toric Marker
9-841-3	Toric Marker
9-841-4	Toric Marker
9-842-1	Blade Marker
9-845-2	Incision Marker
9-846	Incision Marker
9-854R	LASIK Marker



Markers, Non-Invasive	
REF	Device Title
9-855	LASIK Marker
9-864-1	Barrel Marker

Table 2. Relevant standards and common specifications.

Applicable Standards	
Standard Number	Title
EN 1041:2008	Information supplied by the manufacturer of medical devices
EN ISO 10993-1:2009	Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process (ISO 10993-1:2009) EN ISO 10993-1:2009/AC:2010
EN ISO 13485:2016	Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2016) EN ISO 13485:2016/AC:2018
EN ISO 14971:2012	Medical Devices - Application of risk management to medical devices (ISO 14971:2007, Corrected version 2007-10-01)
EN ISO 15223-1:2016	Medical devices - Symbols to be used with medical device labels, labelling and information to be supplied - Part 1: General requirements (ISO 15223-1:2016, Corrected version 2017-03)
EN ISO 17664:2017	Processing of health care products - Information to be provided by the medical device manufacturer for the processing of medical devices (ISO 17664:2017)
EN 62366-1:2015+A1:2020	Medical devices - Part 1: Application of usability engineering to medical devices
93/42/EEC	Council Directive concerning medical devices
2007/47/EC	Council Directive amending 93/42/EEC
SI 2002 No. 618	Medical Devices Regulations (UK)
MEDDEV 2.7/1 rev.4 June 2016	Clinical evaluation: A guide for manufacturers and notified bodies
MEDDEV 2.12/1 rev.8 Jan 2013	Guidelines on a medical device's vigilance system



**ADDENDUM TO THE ORIGINAL
DECLARATION OF CONFORMITY**

Per 31 January 2023, the address of our EU Authorised Representative as listed on the original Declaration of Conformity has changed

Old Address

Authorised Representative:

Emergo Europe
Prinsessegracht 20
2514 AP The Hague
The Netherlands

New Address

Authorised Representative:

Emergo Europe
Westervoortsedijk 60
6827 AT Arnhem
The Netherlands

.....
Martin Lock
Head of Regulatory Affairs

..... 25/01/23

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Date