









ISO9001:2015 and ISO13485:2016 Certified

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 Invasive Scissors

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, invasive medical devices.

The classification is determined under Annex IX, Rule 5.

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

Martin Lock

Head of Regulatory Affairs

Date 14-MAY-2021

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

Scissors, Invasive		
REF	Device Title	
1-110	Vannas Scissors	
1-111	Vannas Scissors	
1-111B	Vannas Scissors	
1-112	Vannas Scissors	
1-116	Capsule Scissors	
1-118	Corneal Scissors	
1-120	Vannas Scissors	
1-121	Vannas Scissors	
1-122	Vannas Scissors	
1-211	Iris Scissors	
1-211B	Iris Scissors	
1-218	Corneal Scissors	
1-219	Corneal Scissors	
1-227	Conjunctival Scissors	
1-312	Vannas Scissors	
1-400	Corneal Scissors	
1-401	Corneal Scissors	
1-410	Corneal Scissors	
1-411	Corneal Scissors	
1-500	Westcott Scissors	
1-500B	Westcott Scissors	
1-501	Westcott Scissors	
1-510	Westcott Scissors	
1-625	Capsule Scissors	
1-630	Iris Scissors	
1-695	IOL Scissors	
1-700	IOL Scissors	
1-705	IOL Scissors	
1-841	Vitreoretinal Scissors	
1-841N	Vitreoretinal Scissors Head	
1-842	Vitreoretinal Scissors	
1-842N	Vitreoretinal Scissors Head	
7-101	Descemet's Punch	
7-102	Descemet's Punch	
7-105	Descemet's Punch	
P5464	IOL Scissors	

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+44 (0) 1462 893254





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Table 2. Relevant standards and common specifications.

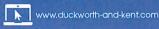
Applicable Standards			
Standard Number	Title		
EN 1041:2008	Information supplied by the manufacturer of medical devices		
EN 1707:1996	Conical fittings with a 6% (Luer) taper for syringes, needles and certain other medical equipment - Lock fittings		
EN ISO 10993-1:2009	Biological evaluation of medical 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 20594-1:1993+A1:1997	Conical fittings with a 6% (Luer) taper for syringes, needles and certain other medical equipment - Part 1: General requirements (ISO 594-1:1986)		
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		

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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 .

Date

Date 25-JAN-2023

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