

Gemäß EU-Verordnung 2017/745 MDR

Carl Martin GmbH
Neuenkamper Straße 80-86
42657 Solingen
Germany
SRN: DE-MF-000005066



erklärt in alleiniger Verantwortung, dass die folgenden Produktgruppen, auf die sich diese Konformitätserklärung bezieht, unter Beachtung der EU 2017/745 MDR und der daraus resultierenden regulatorischen Anforderungen hergestellt wurden:

Produktgruppe	Gruppen.-Nr.	Risikoklasse und Regel	Basis-UDI-DI
Zahnzangen	001	Ir (Regel 6)	++ECMSBUDI0016L
Mund- / Wangen- / Lippenhalter / Mundsperrer	006	Ir (Regel 6)	++ECMSBUDI0066W
Zungenspatel / Zungenzangen	007	Ir (Regel 6)	++ECMSBUDI0076Y
Wurzelheber / Syndesmotome / Drehmeißel / Periotome	008	Ir (Regel 6)	++ECMSBUDI00872
Chirurgische / Anatomische Pinzetten	015	Ir (Regel 6)	++ECMSBUDI0156X
Scheren	022	Ir (Regel 6)	++ECMSBUDI0226U
Skalpelle / Skalpellklingenhalter	027	Ir (Regel 6)	++ECMSBUDI02776
Scaler / Küretten	031	Ir (Regel 6)	++ECMSBUDI0316V
Endodontieinstrumente	034	Ir (Regel 6)	++ECMSBUDI03473
Parodontalsonden	035	Ir (Regel 6)	++ECMSBUDI03575
Excavatoren	038	Ir (Regel 6)	++ECMSBUDI0387B
Tamponstopfer / Watterollenhalter	040	Ir (Regel 6)	++ECMSBUDI0406W
Wundhaken	042	Ir (Regel 6)	++ECMSBUDI04272
Scharfe Löffel	043	Ir (Regel 6)	++ECMSBUDI04374
Nadelhalter	045	Ir (Regel 6)	++ECMSBUDI04578
Korn- / Tupferzange / Tuchklemmen	058	Ir (Regel 6)	++ECMSBUDI0587H
Hohlmeißelzangen / Gewebebezangen	059	Ir (Regel 6)	++ECMSBUDI0597K
Fistel-/Myrthenblattsonden	060	Ir (Regel 6)	++ECMSBUDI06074
Flach- / Hohlmeißel / Osteotome / Sinutome	062	Ir (Regel 6)	++ECMSBUDI06278

Konformitätserklärung

Knochenschaber /Knochenfeilen / Gingivalrandschräger	063	Ir (Regel 6)	++ECMSBUDI0637A
Spritzen	064	Ir (Regel 6)	++ECMSBUDI0647C
Tunnelierungsinstrumente	071	Ir (Regel 6)	++ECMSBUDI07179
Raspatorien / Papillenheber	072	Ir (Regel 6)	++ECMSBUDI0727B
Sinuslifttelevatoren / Membraninstrumente	074	Ir (Regel 6)	++ECMSBUDI0747F
Knochenmaterielinstrumente	080	Ir (Regel 6)	++ECMSBUDI0807A
Arterienklemme / Blutstillungsmeißel	085	Ir (Regel 6)	++ECMSBUDI0857L

Benannte Stelle und Kennnummer:

TÜV Rheinland LGA Products GmbH (0197), Tillystraße 2, 90431 Nürnberg

Durchgeführtes Konformitätsbewertungsverfahren:

MDR Anhang IX Kapitel I

EU-Zertifikat:

Registration No.: HZ 1594091-1

Report No.: 1206333-10

Gültig ab: 29.06.2026

Gültig bis: 31.12.2028

Solingen, 29.06.2026


Peter Holzknecht
 Geschäftsführer

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Declaration of Conformity

According to EU Regulation 2017/745 MDR

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declares under sole responsibility that the following product groups, to which this declaration of conformity relates, have been manufactured in compliance with the EU Regulation 2017/745 MDR and the resulting regulatory requirements:

Product Group	Group No.	Risk Class and Rule	Basic-UDI-DI
Extraction forceps	001	Ir (Rule 6)	++ECMSBUDI0016L
Mouth- / cheek- / lip retractor / mouth gag	006	Ir (Rule 6)	++ECMSBUDI0066W
Tongue depressors / tongue forceps	007	Ir (Rule 6)	++ECMSBUDI0076Y
Root elevator / syndesmotome / lathe chisel / periostome	008	Ir (Rule 6)	++ECMSBUDI00872
Tissue forceps	015	Ir (Rule 6)	++ECMSBUDI0156X
Scissors	022	Ir (Rule 6)	++ECMSBUDI0226U
Scalpels / scalpel blade holders	027	Ir (Rule 6)	++ECMSBUDI02776
Scaler / curettes	031	Ir (Rule 6)	++ECMSBUDI0316V
Endodontic instruments	034	Ir (Rule 6)	++ECMSBUDI03473
Periodontal probes	035	Ir (Rule 6)	++ECMSBUDI03575
Excavators	038	Ir (Rule 6)	++ECMSBUDI0387B
Tongue depressors / cotton roll holders	040	Ir (Rule 6)	++ECMSBUDI0406W
Tissue retractors	042	Ir (Rule 6)	++ECMSBUDI04272
Bone curettes	043	Ir (Rule 6)	++ECMSBUDI04374
Needle holder	045	Ir (Rule 6)	++ECMSBUDI04578
Dressing and sponge forceps / Towel clips	058	Ir (Rule 6)	++ECMSBUDI0587H
Bone rongeur forceps / tissue nipper	059	Ir (Rule 6)	++ECMSBUDI0597K
Antrum probes	060	Ir (Rule 6)	++ECMSBUDI06074
Chisels / gouges / osteotomes / sinutomes	062	Ir (Rule 6)	++ECMSBUDI06278
Bone scraper / bone files / gingival margin trimmer	063	Ir (Rule 6)	++ECMSBUDI0637A
Syringes	064	Ir (Rule 6)	++ECMSBUDI0647C

Declaration of Conformity

Tunneling instruments	071	Ir (Rule 6)	++ECMSBUDI07179
Periosteal elevators / papilla elevator	072	Ir (Rule 6)	++ECMSBUDI0727B
Sinus-lifting instruments / membrane instruments	074	Ir (Rule 6)	++ECMSBUDI0747F
Bone material instruments	080	Ir (Rule 6)	++ECMSBUDI0807A
Hemostatic forceps / hemostatic chisel	085	Ir (Rule 6)	++ECMSBUDI0857L

Notified Body and number:

TÜV Rheinland LGA Products GmbH (0197), Tillystraße 2, 90431 Nürnberg

Conformity assessment based on:

MDR Annex IX chapter I

EU Certificate:

Registration No.: HZ 1594091-1

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Solingen, 29.06.2026


Peter Holzknecht
Managing Director