

According to EU Regulation 2017/745 MDR

Carl Martin GmbH
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Germany
SRN: DE-MF-000005066



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	Basic-UDI-DI
Extraction forceps	001	Ir	++ECMSBUDI0016L
Mouth- / cheek- / lip retractor / mouth gag	006	Ir	++ECMSBUDI0066W
Tongue depressors / tongue forceps	007	Ir	++ECMSBUDI0076Y
Root elevator / syndesmotome / lathe chisel / periotome	008	Ir	++ECMSBUDI00872
Tissue forceps	015	Ir	++ECMSBUDI0156X
Scissors	022	Ir	++ECMSBUDI0226U
Scalpels / scalpel blade holders	027	Ir	++ECMSBUDI02776
Scaler / curettes	031	Ir	++ECMSBUDI0316V
Endodontic instruments	034	Ir	++ECMSBUDI03473
Periodontal probes	035	Ir	++ECMSBUDI03575
Excavators	038	Ir	++ECMSBUDI0387B
Tongue depressors / cotton roll holders	040	Ir	++ECMSBUDI0406W
Tissue retractors	042	Ir	++ECMSBUDI04272
Bone curettes	043	Ir	++ECMSBUDI04374
Needle holder	045	Ir	++ECMSBUDI04578
Dressing and sponge forceps / Towel clips	058	Ir	++ECMSBUDI0587H
Bone rongeur forceps / tissue nipper	059	Ir	++ECMSBUDI0597K
Antrum probes	060	Ir	++ECMSBUDI06074
Chisels / gouges / osteotomes / sinutomes	062	Ir	++ECMSBUDI06278
Bone scraper / bone files / gingival margin trimmer	063	Ir	++ECMSBUDI0637A
Syringes	064	Ir	++ECMSBUDI0647C

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

Notified Body and number:

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

Conformity assessment based on:

MDR Annex IX chapter I

Valid from: 13.10.2023

Valid to: 31.12.2024

Solingen, 13.10.2023



Peter Holzknecht
Geschäftsführer