



EC Certificate

Full Quality Assurance System

Certificate No.:
245818-2017-CE-NOR-NA-PS Rev. 4.0

Project No.:
PRJC-513705-2014-MSL-NOR

Valid Until:
01 April 2023

This is to certify that the quality system of:

Ortho Organizers, Inc.

1822 Aston Ave.
Carlsbad, CA 92008-7306
USA

For design, production and final product inspection/testing of:

Non-active medical devices with measuring function and dental materials

Has been assessed with respect to:

The conformity assessment procedure described in Annex II excluding section 4 of Council Directive 93/42/EEC on Medical Devices, as amended

and found to comply.

Further details of the product(s) and conditions for certification are given overleaf.

Place and Date:
Høvik, 31 August 2018



For:
DNV GL PRESAFE AS

Tone Kolpus

Tone Kolpus

The Certificate has been digitally signed.
See www.presafe.com/digital_signatures for more info

Notice: The Certificate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid.



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Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as "Forskrift om Medisinsk Utstyr" by the Norwegian Ministry of Health and Care Services.

Certificate history:

Revision	Description	Issue Date
3.0	Replaces Nemko AS (NB 0470) certificate no. EU1209401 and certificate no. EU1208413 following transfer of notified body functions to DNV GL Nemko Presafe AS (NB 2460)	2017-10-18
1.0	Addition of Metal Brackets and Buccal Tubes already certified in former Nemko AS (NB 0470) certificate no. EU1209401 and certificate no. EU1208413	2017-12-04
2.0	Correction of validity date	2017-12-22
3.0	Recertification	2018-08-17
4.0	Devices marked with bold added to the certificate	2018-08-31



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Products covered by this Certificate:

Product Description	Product Name	Class
Non-Active Medical Devices with Measuring Function	Measuring Device (Ruler)	Im
Dental Materials	Ceramic Brackets	Ila
	Composite and Plastic Brackets	Ila
	Metal Self-Ligating Brackets	Ila
	Metal Brackets and Buccal Tubes	Ila
	Ceramic Self-Ligating brackets	Ila
	Orthodontic Bands and Assemblies	Ila
	Auxiliaries	Ila
	Orthodontic Wires and Accessories	Ila
	Elastomerics	Ila
	Intraoral Appliances	Ila
	Fixed Attachments	Ila
	Seperators	Ila

The complete list of devices is filed with the Notified Body

Sites covered by this certificate

Site Name	Address
Ortho Organizers, Inc.	1822 Aston Avenue, Carlsbad, CA 92008-7306, USA

EU Representative

MDI Europa GmbH, Langenhagener, Straße 71, D-30855 Langenhagen, Germany



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Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform Presafe of any intended updating of the quality system and Presafe will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system. Presafe reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number of Presafe.

End of Certificate



Notified Body Confirmation Letter Reference: C569778

To whom it may concern,

Confirmation of the status of a formal application, written agreement, and appropriate surveillance in the framework of Regulation EU 2023/607 amending Regulations (EU) 2017/745 and (EU) 2017/746 as regards the transitional provisions for certain medical devices and in vitro diagnostic medical devices

This letter confirms that, DNV Product Assurance AS, a Notified Body (NB) designated against Regulation (EU) 2017/745 (MDR) and identified by the number NB 2460 on NANDO, has received a formal application in accordance with Section 4.3, first subparagraph of Annex VII of MDR and has signed a written agreement in accordance with Section 4.3, second subparagraph of Annex VII of MDR with the following manufacturer:

Ortho Organizers, Inc.

1822 Aston Ave., 92008-7306 Carlsbad, CA USA

SRN Number: US-MF-000010061

The devices covered by the formal application and the written agreement mentioned above are identified in the Tables below. Table 1 identifies the devices for which an MDR application has been received, written agreement concluded and for which the NB is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive. Table 2 identifies the devices for which an MDR application has been received and a written agreement concluded, but the NB has not yet taken the responsibility for appropriate surveillance of the corresponding devices under the applicable Directive.

In the case of devices covered by certificates issued under Directive 93/42/EEC (MDD) that expired after 26 May 2021 and before 20 March 2023, without having been withdrawn, this letter also confirms that the manufacturer signed the written agreement under MDR by the date of MDD certificate expiry; or provided evidence that a competent authority of a Member State had granted a derogation or exemption from the applicable conformity assessment procedure in accordance with Article 59(1) of MDR or Article 97(1) of the MDR respectively, by the 20 Mar 2023 for the relevant devices.

The transition timelines that apply to the devices covered by this letter, subject to the manufacturer's continued compliance to the other conditions specified in Article 120.3c of MDR (as amended by (EU) 2023/607), are shown below:

- 26 May 2026 for Class III custom-made implantable devices
- 31 December 2027 for Class III devices and Class IIb implantable devices excluding Well-established technologies (WET - sutures, staples, dental fillings, dental braces, tooth crowns, screws, wedges, plates, wires, pins, clips and connectors)
- 31 December 2028 for other Class IIb devices, Class IIa, Class I devices placed on the market in sterile condition or have a measuring function

Place and date:
Høvik, 2024/07/17



For the issuing office:
DNV Product Assurance AS – Notified Body 2460
Veritasveien 1, 1363 Høvik, Norway

C. Rajesh Kumar

Rajesh Kumar Chellappan
Management Representative

Lack of fulfilment of conditions as set out in the Certification Agreement may render this letter invalid.

NOTIFIED BODY 2460: DNV Product Assurance AS, Veritasveien 1, 1363 Høvik, Norway, Tel +47 67 57 88 00, www.dnv.com



- 31 December 2028 for devices not requiring the involvement of a notified body under MDD but requiring it under MDR (e.g., class I devices that qualify as re-usable surgical instruments)

Table 1: Devices covered by this letter and for which the NB is also responsible for appropriate surveillance of the corresponding devices under the applicable Directive:

Device name and Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the quotation request review stage)	If the MDR device is a substitute device, identification of the corresponding MDD device	MDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
Orthodontic Bands and Assemblies 019070702MolarBands6B	Class IIa	NA	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Latex Elastomerics 019070702L-ElastomericsKX	Class IIa	Elastomerics (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Latex- Free Elastomerics 019070702LF-ElastomericPU	Class IIa	Elastomerics (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Metal Brackets 019070702MetalBracketsEZ	Class IIa	Metal Brackets and Buccal Tubes (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Metal Brackets- (SLX) 019070702MetalBracketsEZ	Class IIa	Metal Self-Ligating Brackets (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Motion 3D Metal 019070702Motion3DApplncMV	Class IIa	Intraoral Appliances (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Motion 3D Clear 019070702Motion3DApplncMV	Class IIa	Intraoral Appliances (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Ceramic/ Composite/ Plastic Brackets 19070702PlasticBracketCM	Class IIa	Ceramic Self-Ligating brackets Composite and Plastic Brackets Ceramic Brackets (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Metal Bracket Auxiliaries (Buccal Tubes, Hooks, and Button) 019070702MetIBracketAuxQB	Class IIa	Auxiliaries Fixed Attachments (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460

Device name and Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the quotation request review stage)	If the MDR device is a substitute device, identification of the corresponding MDD device	MDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
Archwire Accessories 019070702ArchwireAccess2H	Class IIa	Orthodontic Wires and Accessories Auxiliaries (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Fixed and Functional Appliances 019070702F&FAppliancesV4	Class IIa	Intraoral Appliances (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Measuring Devices 019070702MeasureDeviceVH	Class I devices with a measuring function	Measuring Device (Ruler) (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460
Archwires 019070702ArchwiresW9	Class IIa	Orthodontic Wires and Accessories Auxiliaries (Only Name Change)	Certificate number: 245818-2017-CE-NOR-NA-PS Rev. 4.0 NB number NB: 2460

Table 2: Devices covered by this letter and for which the NB is NOT responsible for appropriate surveillance of the corresponding devices under the applicable Directive:

Device name and Basic UDI-DI (under MDR application)	MDR Device classification (as proposed by the manufacturer and verified at the pre-application stage)	If the MDR device is a substitute device, identification of the corresponding MDD device	MDD Certificate Reference(s) of the devices under MDR application, and the NB Identification
NA	NA	NA	NA

Confirmation Letter Revision History

Date	NB internal reference traceable to each version of the letter	Action
2023/10/18	C569778	Initial issue
2024/07/17	C569778	Addition of devices to the list: Metal Brackets, Metal Bracket Auxiliaries, Archwire Accessories, Fixed and Functional Appliances, Archwires

Lack of fulfilment of conditions

The following may render this letter of confirmation invalid:

- Lack of compliance to the requirements of Regulation (EU) 2023/607
- Significant changes to design or intended purpose of the devices
- Changes in the quality system affecting production
- Periodical audits not held within the timeframe

EU DECLARATION OF CONFORMITY

Product Name/Trade Name: Reusable Instruments REF	
Basic UDI-DI Information: See Attachment 1 Device Product Codes: See Attachment 3	
 <p>Ortho Organizers, Inc. 1822 Aston Avenue Carlsbad, CA 92008 USA Phone: +760-448-8600 SRN: US-MF-000010061</p>	
<div style="border: 1px solid black; display: inline-block; padding: 2px;"> <div style="border: 1px solid black; display: inline-block; padding: 2px;">EC</div> <div style="border: 1px solid black; display: inline-block; padding: 2px;">REP</div> </div> <p>mdi Europa GmbH Langenhagener Str. 71 30855 Langenhagen / Germany Europe Phone: +49 511 39089530 Website: https://mdi-europa.com/ SRN: DE-AR-000006218</p>	
Quality Certificates Issued: Certificate No: 908347	

The object of the declaration described above is in conformity with the following regulations:

EU Regulation	Regulation (EU) 2017/745 of the European Parliament and of the Council of 5 April 2017 on medical devices (EU MDR)
Device Risk Classification	Class I based on Annex VIII and Rule 5.
Conformity Assessment Path	N/A
Notified Body Name, Address, and ID	Not applicable for this class I medical device that does not have a measuring function nor is it a reusable surgical instrument.
Standards	The products listed on this Declaration of Conformity have been assessed and/or tested in a typical configuration as described in the Manufacturer's accompanying documentation in accordance with the product standards listed below. Refer to Attachment 2.



This declaration of conformity is issued under the sole responsibility of the manufacturer:
Ortho Organizers, Inc.

Signed by:
Nicole Rajek
Signer Name: Nicole Rajek
Signing Reason: I approve this document
Signing Time: 9/5/2024 | 1:26:32 PM PDT
A31D4E89BEFD406B92A150785D079782

Carlsbad, CA

9/5/2024

Signature	Issued in	Date
Nicole Rajek		
Printed Name	Sr. Director, QA/RA	Title

Attachment 1

Product Name	Reusable Instruments
Product Part Number(s) and Descriptions	Attachment 3.
Intended Use	Orthodontic Instruments are to be used only by licensed dental professionals and/or orthodontic professionals during orthodontic treatment. Orthodontic instruments are provided non-sterile and require sterilization before each use. Orthodontic Hand Instruments are to be used during dental and/or orthodontic treatment to adjust and/or manipulate orthodontic wires and/or appliances.
Product Options/Accessories Part Number(s) and Descriptions	None
Unique Device Identifier	0190707O2ReusInstrument94
Category of Devices	Orthodontic Reusable Instruments
Global Medical Device Nomenclature Codes (GMDN) / (CND) and Description	GMDN Code: 33209- Orthodontic pliers 62072-Orthodontic elastomeric appliance pusher 35320-Manual dental scaler, reusable 37413-Orthodontic ligature director 62086-Self-ligating orthodontic bracket open/close tool CND Code: Q010499 – ORTHODONTIC DEVICES - OTHERS
Common Specifications	None

Attachment 2

Standard No.	Title
ISO 13485:2016+AC:2018	Medical devices – Quality management systems – Requirements for regulatory purposes
ISO 14971:2019	Medical devices – Application of risk management to medical devices
ISO 10993-1:2018	Biological evaluation of medical devices – Part 1: Evaluation and testing within a risk management process
ISO 10993-2:2006	Biological evaluation of medical devices – Part 2: Animal welfare requirements
ISO 10993-3:2014	Biological evaluation of medical devices – Part 3: Tests for genotoxicity, carcinogenicity and reproductive toxicity
ISO 10993-5:2009	Biological evaluation of medical devices – Part 5: Tests for in vitro cytotoxicity
ISO 10993-10:2010	Biological evaluation of medical devices – Part 10: Tests for irritation and skin sensitization
EN ISO 10993-11:2017	Biological Evaluation of Medical Devices – Part 11: Tests for system toxicity
EN ISO 10993-12:2013	Biological evaluation of medical devices – Part 12: Sample preparation and reference materials
EN 1641:2009	Dentistry – Medical devices for dentistry – Materials
IEC 62366-1:2015 +COR1:2016	Medical Devices – Application of usability engineering to medical devices
ISO 7405:2018	Dentistry – Evaluation of biocompatibility of medical devices used in dentistry
ISO 15223-1:2021	Medical Devices – Symbols to be used with medical device labels, labeling and information to be supplied – Part 1: General requirements
ISO 20417:2021	Information to be supplied by the manufacturer of medical devices
MEDDEV 2.7/1 Rev 4	Clinical evaluation: A guide for manufacturers and notified bodies under directives 93/42/EEC and 90/385/EEC
ISTA 2A: 2011	The Association for Transport Packaging
AAMI TIR12:2010	Designing, testing and labeling reusable medical devices for reprocessing in health care facilities: A guide for medical device manufacturers

Standard No.	Title
AAMI TIR30:2011	A compendium of processes, materials, test methods, and acceptance criteria for cleaning reusable medical devices
AAMI TIR 34:2014	Water for the Reprocessing of Medical Devices
ANSI/AAMI T77:2013/ (R) 2018	Containment Devices for Reusable Medical Device Sterilization
ANSI/AAMI T79:2017	Comprehensive Guide to Steam Sterilization and Sterility Assurance in Health Care Facilities
DIN EN 556-1:2001/AC:2006	Sterilization of Medical Devices – Requirements for Medical Devices to be Designated “STERILE” – Part 1: Requirements for Terminally Sterilized Medical Devices
ISO 17664-1:2004	Processing of health care products — Information to be provided by the medical device manufacturer for the processing of medical devices — Part 1: Critical and semi-critical medical devices
ISO 17665-1:2006	Sterilization of Health Care Products – Moist Heat – Requirements for Development, Validation and Routine Control of a Sterilization Process for Medical Devices
ISO/TS 17665-2:2009	Sterilization of Health Care Products – Moist Heat – Part 2: Guidance on the Application of ISO 17665-1
ISO 11138-3:2017	Sterilization of Health Care Products – Biological Indicators – Part 3: Biological Indicators for Moist Heat Sterilization Process
ISO 11607-1:2019	Packaging for Terminally Sterilized Medical Devices – Part 1: Requirements for Materials, Sterile Barrier Systems and Packaging

Attachment 3

Part Number	Descriptions
4150814	Plier Heavy Duty Cutter
4150893	Mathieus Hollo-Form Tip Instrument
4150895	Mathieus Hook Tip Instrument
4150898	Mathieus Regular Medium Tip Instrument
4150899	Mathieus Regular Narrow Tip Instrument
4150953	Elastic Separating Plier
4151604	Choice Micro Pin & Ligature Cutter
4151605	Choice Pin & Ligature Cutter
4151609	Choice Flush Cut Distal End Cutter w/ Safety Hold
4151610	Choice Hard Wire Cutter
4151611	Choice Angled Slim Nose Pin & Ligature Cutter
4151612	Choice Three Prong Bending Pliers
4151616	Choice Headgear Pliers
4151617	Choice Distal End Cutter w/ Safety Hold
4151618	Choice Mini Distal End Cutter w/ Safety Hold
4151619	Choice Distal End Cutter w/ Safety Hold - Long Handle
4151621	Choice Spring Flush Cut DistalEnd Cutter w/ Safety Hold
4151623	Choice Torquing Pliers - Male
4151624	Choice Torquing Pliers - Female
4151626	Choice Jaraback w/ Cutter
4151633	Choice Light Wire Bird Beak w/ Cutter
4151634	Choice Secure Hold Bird Beak Pliers
4151639	Choice Three Prong Pliers w/ Groove
4151640	Choice Tweed Arch Forming Pliers
4151641	Choice Three Prong Lab Pliers w/ Groove
4151643	Choice Intraoral Detailing Pliers - Left
4151644	Choice Intraoral Detailing Pliers - Right
4151645	Choice Double Sided Detailing Pliers - 1mm
4151649	Choice V Notch Stop Pliers
4151650	Choice Adams Pliers
4151652	Choice Straight Bracket Removing Pliers
4151653	Choice Angled Bracket RemovingPliers
4151654	Choice Posterior Band RemovingPliers- Long Handle
4151655	Choice Adhesive Removing Pliers
4151656	Choice High Force Band Removing Pliers
4151662	Choice Crown and Band Crimping Pliers



Part Number	Descriptions
4151663	Choice Hook Crimping Pliers
4151670	Choice Lab Wire Bird Beak Pliers
4151681	Choice Tweed Loop Forming Pliers
4151687	Choice Weingart Pliers
4151695	Choice Distal Cinch Back Pliers
4151937	Replacement Tungsten Carbide Tip
4151942	Replacement Bushings (6)
106-215	Abrasive Strip Placer
200-109	Elastic Remover
200-311	Posterior Direct Bond Bracket Holder
200-329	Bracketizer Instrument
200-407	Elastic Separating Pliers
200-407D	Deluxe Elastic Separating Pliers
200-412	Extra-Heavy Wire Cutter
200-450	Nitanium Expander Adjusting Pliers
201-101	Endura Plus Mathieu Wide Tip
201-101TC	Endura Plus Mathieu Wide Tip — Tungsten Carbide Deluxe
201-110	Endura Plus Mathieu Hole Tip
201-118	Endura Plus Mathieu Hook Tip
201-121	Endura Plus Mathieu Narrow Tip
201-121TC	Endura Plus Mathieu Narrow Tip — Tungsten Carbide Deluxe
201-123	Endura Plus Mathieu Half-Fine Tip
201-125	Endura Plus Mathieu Wide Tip with Safe-T-Lock — Tungsten Carbide Deluxe
201-126	Endura Plus Mathieu Narrow Tip with Safe-T-Lock — Tungsten Carbide Deluxe
201-199ATE	Eraser Plier-Long Handle
201-199ATPP	Pressure Point Plier-Long Handle
201-500	Band Pusher Scaler - 1/4" Handle
201-500W	Band Pusher Scaler - 3/8" Handle
201-501	Double-Ended Ligature Director - 1/4" Handle
201-501W	Double-Ended Ligature Director - 3/8" Handle
201-502	Ligature Director Scaler - 1/4" Handle
201-502W	Ligature Director Scaler - 3/8" Handle
201-503	Ligature Remover Director - 1/4" Handle
201-503W	Ligature Remover Director - 3/8" Handle
205-402	Narrow Tip
205-403	Smaha Ultra Fine Tip



Part Number	Descriptions
201-510	SLX 3D EZ Twist Opener
201-505	Carriere Double Ended Opener Tool
201-507	Carriere Motion 3D Placement Instrument
204-101	Endura Plus Distal End Cutter (Safety Hold)
204-101XL	Endura Plus Distal End Cutter (Safety Hold) - Long Handle
204-103	Endura Plus Hard Wire Cutter, 15° Angled
204-105	Endura Plus Multi-Use Cutter
204-107	Endura Plus Micro Mini Pin & Ligature Cutter
204-107XL	Endura Plus Micro Mini Pin & Ligature Cutter - Long Handle
204-109	Endura Plus Micro Mini Pin & Ligature Cutter, 15° Angled
204-110	Endura Plus Micro Cutter
204-111	Endura Plus Distal End Cutter Safety Hold & Flush Cut
204-202	Endura Plus Slim Weingart Pliers
204-203	Endura Plus How Pliers
204-206	Endura Plus Adhesive Removing Pliers
204-207	Endura Plus Posterior Band Removing Pliers - Long
204-213	Endura Plus Direct Bond Bracket Removing Pliers
204-213B	Endura Plus Direct Bond Bracket Removing Pliers, Blade
204-219	Endura Plus Debonding Pliers
204-220XL	Endura Plus Angulated Debonding Pliers
204-221	Endura Plus Crown & Band Contouring Pliers
204-301	Endura Plus Jarabak Pliers
204-302	Endura Plus Three Jaw Pliers
204-304	Endura Plus Bird Beak Pliers
204-305	Endura Plus Light Wire Bird Beak Pliers w/Groove
204-306	Endura Plus Tweed Loop Forming Pliers
204-307	Endura Plus Arch Bending Pliers
204-309	Endura Plus Lingual Arch Forming Pliers
204-310	Endura Plus Hollow Chop Contouring Pliers
204-312	Endura Plus Mini Three Jaw Pliers
204-318	Endura Plus Light Wire Bird Beak Pliers
204-319	Endura Plus Nance Loop Forming Pliers
204-320	Endura Plus Adams Pliers
204-321	Endura Plus Stop (V-Bend) Pliers
204-323	Endura Plus Optical Pliers
204-325	Endura Plus Bird Beak Pliers w/Cutter
204-328	Endura Plus Loop Tie Back Pliers
204-329	Endura Plus Detailing Step Pliers 0.50 mm
204-330	Endura Plus Detailing Step Pliers 0.75 mm
204-331	Endura Plus Detailing Step Pliers 1mm
204-332	Endura Plus Hook Crimping Pliers (Straight)
204-333	Endura Plus Hook Crimping Pliers (Angled)



Part Number	Descriptions
204-334	Endura Plus Utility Arch Pliers
204-335	Endura Plus Angled Utility Arch Pliers
204-336	Endura Plus Utility Step Pliers
204-337	Endura Plus Torquing Pliers Set w/Key
204-338	Endura Plus Small Distal End Cutter(SafetyHold)w/LongHandle
205-011	SAFETY HOLD REPLACEMENT SILICONE INSERT (3)
205-101	TRIUMPH DISTAL END CUTTER SAFETY HOLD
205-101XL	TRIUMPH DISTAL END CUTTER LONG HANDLE SAFETY HOLD
205-104	TRIUMPH HARD WIRE CUTTER
205-105	TRIUMPH MULTI-USE CUTTER
205-106	TRIUMPH MINI PIN & LIG CUTTER
205-107	TRIUMPH MICRO MINI PIN & LIG CUTTER
205-109	TRIUMPH MICRO MINI PIN & LIG CUTTER, 15 DEGREE
205-110	TRIUMPH MICRO CUTTER
205-111	TRIUMPH DISTAL END CUTTER SAFETY HOLD FLUSH CUT
205-202	TRIUMPH WEINGART PLIER
205-203	TRIUMPH HOW PLIERS
205-206	TRIUMPH ADHESIVE REMOVING PLIERS
205-206B	ADHESIVE RMR REPLACEMENT BLADE TRIUMPH
205-206T	ADHESIVE RMR 1/4 REP PADS (6) TRIUMPH
205-207	TRIUMPH BAND REMOVER, LONG POSTERIOR
205-207T	BAND RMR 3/16 REPLAC. PADS (6) TRIUMPH
205-207TT	BAND RMR 1/4 REPLAC. PADS (6) TRIUMPH
205-219	TRIUMPH DEBONDING PLIERS
205-220XL	TRIUMPH ANGULATED DEBONDING PLIERS
205-221	TRIUMPH CROWN BAND CONTOURING PLIERS
205-301	TRIUMPH JARABAK PLIERS
205-302	TRIUMPH THREE JAW PLIERS
205-304	TRIUMPH BIRD BEAK PLIERS
205-305	TRIUMPH LIGHT WIRE WITH GROOVE PLIERS
205-306	TRIUMPH TWEED LOOP FORMING PLIERS
205-308	TRIUMPH RECT. ARCH FORMING PLIERS
205-309	TRIUMPH LINGUAL ARCH FORMING PLIERS
205-310	TRIUMPH HOLLOW CHOP CONTOURING PLIERS
205-318	TRIUMPH LIGHT WIRE PLIERS
205-321	TRIUMPH STOP (V-BEND) PLIERS
205-322	TRIUMPH NITI THREE JAW PLIERS
205-325	TRIUMPH BIRD BEAK PLIERS WITH CUTTER
205-326	TRIUMPH LONG TAPERED BIRD BEAK PLIERS
205-327	TRIUMPH NITI DISTAL CINCH BACK PLIERS