IMPLANT SOLUTIONS

DENTI® ROOT FORM+

TWO STAGE IMPLANT SYSTEM

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Prof. Dr. István Vajdovich

ABOUT US

DENTI SYSTEM LTD.

Denti System Kft. was founded in 1989 with the objective to develop and produce a dental implant system for the Hungarian dental practice that is widely available for dental patients in need and can be applied safely and economically by dentists. Our primary purpose was to fulfil the increasing demand of patients helping them to achieve a better quality of life.

SMILE FOR EVERYONE

The high quality of our products is guaranteed by Hager & Meisinger GmbH /Germany/ that has been the manufacturer of our implants, instruments and implant system elements since the Spring of 2005 as well as supports our company by its development background.

QUALITY AND TRADITION

The Denti® Implant System has 30 years of background. Its unbroken improvement is based on continuous research and development, theoretical knowledge collected since then and the experiences from the long term clinical practice. The always renewed, high quality Denti® Implant System is the result of many years of scientific experience and constant research and it is used both in Hungary and in other countries with great satisfaction. Denti® products can be easily, safely and highly successfully used in the everyday dental practice. The Denti® name is guaranteed by constant innovation and the strict quality assurance system.

30 YEARS OF INNOVATION

RESEARCH AND DEVELOPMENT

Research activities of our company are primarily focused on dental implants, related instruments and specific surgical methods for implantation. A series of experimental and clinical studies supports our successful research and development activity in implantology. In this research and development activity we consider cooperation with our users as a top priority.

EDUCATION

Education is important part of the activity of our company. We have delivered accredited course with recognized university professors since 1990. We organize courses both for beginner and expert implantologists-dentists and dental technicians. The purpose of our courses is to teach the participants to the practical methods of implantology by using Denti® Implant System and safely apply these methods in their everyday practice. We institutionalized our educational activities in Spring 2010 under the name of DentiDent Implant Clinic.

CONTENTS

DENTI® ROOT FORM+

HEX & TUBE CONNECTION

for the strong mechanical connection, and for the long-term stability

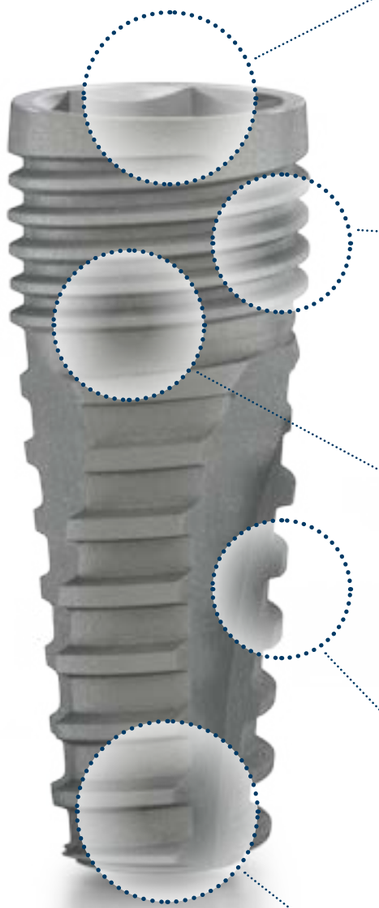
SPECIAL FINE-PITCH THREAD

with small lead angle on the neck of the implant to preserve the level of the peri-implant bone quality

TWO-STAGE

ROOT FORM GEOMETRY

for the optimal distribution of the occlusal force in the bone tissue



HELICAL

SELF-TAPPING DESIGN

up to the neck of the implant for the ideal screw-in torque

WIDE, DEEP THREAD

STRUCTURE

for an excellent primary stability even in case of lower bone quality

SUPERHYDROPHILIC SURFACE

DESIGN

for quick and reliable osseointegration

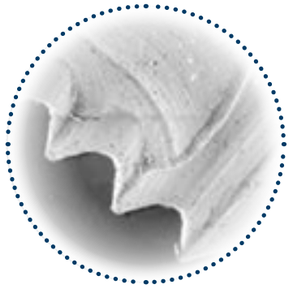
THE APPROVED SURFACE & MATERIAL

SURFACE

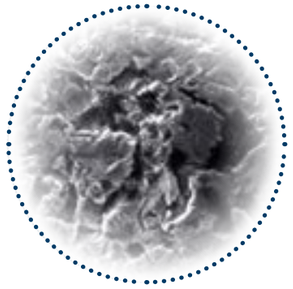
The Denti® implants' surface is the combination of a physical and mechanical surface treatments (sandblasting and chemical treatment). The surface of all implants connecting to the bone shows clear titanium-oxide (rutil) crystal structure which was first micro-roughened using the large particle "sand blasting" technique then thickened with passivation. This micro-roughened, passivated surface design with its high surface energies is the guarantee for the rapid and safe formation and permanent adherence of the bone tissue on the surface of the Denti® implants. Its highly micro-porous surface ensures a stable implant-bone connection thus creating an optimal condition for successful treatment. According to clinical control studies, the success rate of Denti® implants has been above 97% over the past 27 years.

MATERIAL

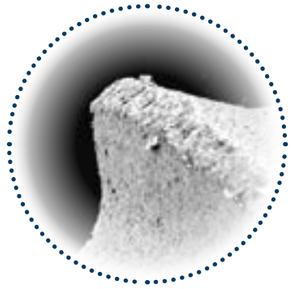
THE DENTI® IMPLANTS MADE OF UNALLOYED, PURE TITANIUM (Grade 4)



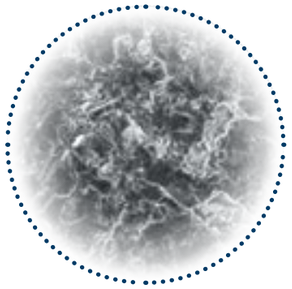
Raw processed surface



Sand-blasted surface



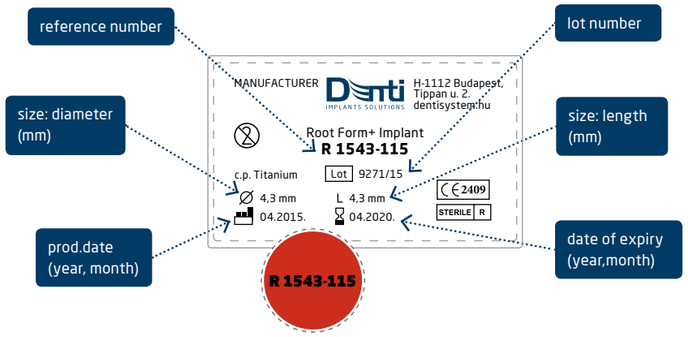
Passivated surface



Surface microstructure of Denti implants (MPS, SEM 500x)

Surface microstructure of Denti implants (MPS, SEM 10.000x)

PACKAGING AND STERILIZATION



ALL COMPONENTS

have double safety packaging one external and one internal (thermo sealed) part.

Titanium implants are sterilized with gamma irradiation.
The sterility is assured for 5 years.
Zirconium dioxide implants are sterilized in an autoclave.
The sterility is assured for 3 years.

THE PACKAGE CONTAINS:

- instructions for use
- guidance for insertion
- guidance for prosthetic procedures
- guidance for impression taking
- detachable labels (LOT number and bar code for unique identification)

All Denti® implants can be traced back according to product name and LOT number, which can be found on both the outer and the inner packings. Please keep the self-adhesive barcode labels.

The external packaging

USABILITY

	Date of expiry (year, month)
Ø	Size: diameter (mm)
LOT	Registry number - individual product identification
CE 1011	CE markings
	Date of manufacture (year, month)
	Do not use more than once
REF	Product code
STERILE R	This product was sterilized with gamma rays
	NON STERILE Not sterile
	Attention, see instruction for use
Rx only	Caution: Federal law (USA) restricts this product to sale by or on the order of a dentist or phisician
	This product was sterilized with heat or steam

label Legend

The internal packaging

REDESIGNED,
DOUBLE STERILE PACKAGING

INNOVATION – ENVIRONMENTAL AWARENESS

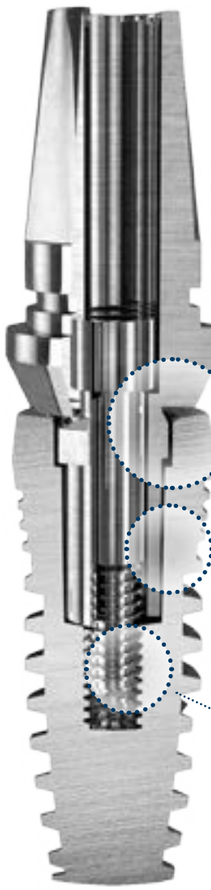
IMPLANT – ABUTMENT CONNECTION

The connection between the implant body and the abutment of Denti® implants is safe according to modern mechanical requirements, and it guarantees long-term stability for the abutments and the assembly on them.

HEX & TUBE connection

Triple safety in the connection between Denti® Root Form+ implants and abutments.

These characteristics significantly decrease the impact on abutment through-bolt, providing maximal stability and minimal possibility for loosening of the implant abutment.



1.

the precise inner hexagonal construction guarantees safe, rotation-free abutments

2.

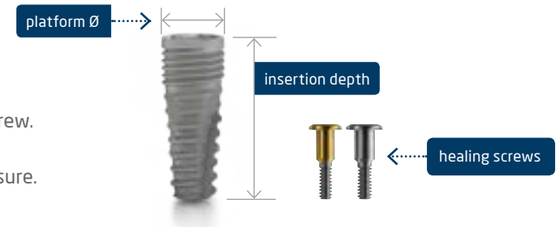
as a result of the precisely fitting long, parallel walls, the forces affecting the abutment are evenly led to the surrounding bone tissue

3.

reinforced, special TiAlV through-bolt to stabilize the implant-abutment unit

DR+ IMPLANT RANGE

The implants are delivered together with the healing screw.
Only single use of the healing screw is suggested.
When tightening the healing screw use only finger pressure.



Denti® DR+ implants Ø 3,3 mm

Reference No.	Insertion depth (mm)	Platform Ø (mm)
R 1033-115	11,5	4,0
R 1033-135	13,5	4,0
R 1033-155	15,5	4,0

Denti® DR+ implants Ø 3,8 mm

Reference No.	Insertion depth (mm)	Platform Ø (mm)
R 1538-095	9,5	4,3
R 1538-115	11,5	4,3
R 1538-135	13,5	4,3
R 1538-155	15,5	4,3

Denti® DR+ implants Ø 4,3 mm

Reference No.	Insertion depth (mm)	Platform Ø (mm)
R 1543-095	9,5	4,8
R 1543-115	11,5	4,8
R 1543-135	13,5	4,8
R 1543-155	15,5	4,8

Denti® DR+ implants Ø 4,8 mm

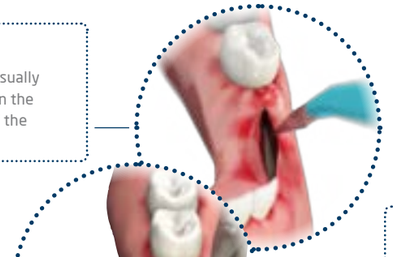
Reference No.	Insertion depth (mm)	Platform Ø (mm)
R 1548-095	9,5	5,3
R 1548-115	11,5	5,3
R 1548-135	13,5	5,3
R 1548-155	15,5	5,3

Denti® DR+ implants Ø 5,3 mm

Reference No.	Insertion depth (mm)	Platform Ø (mm)
R 1553-095	9,5	5,8
R 1553-115	11,5	5,8
R 1553-135	13,5	5,8
R 1553-155	15,5	5,8

1a EXPLORATION

The surface of the alveolar crest is usually explored via an incision performed on the alveolar ridge at the intended site of the implantation.



1b EXPLORATION

Preparation of the mucoperiosteum downwards to the sides should be performed so that the wound edges could be retracted tension-free. The bone surface that has been made visible is then examined.

2 MARKING

In case of a thin alveolar crest, we should mark the future site of the implant(s) on the bone surface with a initial drill.

Initial drill

Spherical drill

3 BALL ROUND BUR

A plate shaped indention of approximately 2 mm in depth is then performed in the bone surface /according to the previous marking/ with a ball round bur. The recommended rpm is 1500 to 2000 U/min for burring.
Any inequality of the alveolar ridge must be smoothed at the intended site of the implantation.

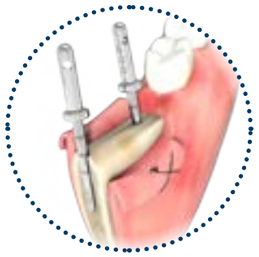
4 PRE-DRILLING

Special care should be paid to the size given on the package of the implant during the one stage surgical implantation of the DR+ implant, as this size indicates the whole length of the passivated body piece of the implant. If the DR+ implant is inserted with one stage surgical protocol, it should be considered on selecting the length of the body piece of the implant that the neck of the implant is 2 mm in length, and this 2 mm should be extracted from the length given on the package of the implant to get the length of the implant with the screw-thread. On preparing the guide channel, parallel holes should be prepared carefully.

Drilling a pilot is performed in the direction and depth determined previously by using a pilot drill with stop ring.

Pilot drill with stop

Parallel pin



5 CHECKING PARALLELISM

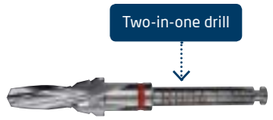
If we insert the parallel pin (8000-058) assuring parallel pre-drilling into the prepared hole, the direction of the previously prepared holes can be visualized. It can help us in preparing the next hole preferably being parallel with the previous one. The prepared hole is then rinsed with saline.

DR+ ONE STAGE SURGICAL PROTOCOL

6 PREPARING THE IMPLANT BED

Dilation of the pre-drilled hole to the core width and recession of the neck piece are performed in one step by using a two-in-one drill. A drill with an appropriate size should be used to prepare the implant nest so that it would fit the implant to be inserted.
The size of the parallel edges of the drill should be the same as the body pieces of the implant; therefore, they can also be used in one stage implantation. Thus, inserting the two-in-one drill into the bone only till the widening neck piece, a reamer hole being exactly of the same size as the body piece of the implant can be prepared.

Two-in-one drill



Thread cutter

7 THREAD CUTTING

In case the patient's mandible is D1 quality with thick and tough cortical layers, a thread should be cut into the hole with the appropriate size thread cutter before the insertion of the implant. Thread cutting is performed at low rpm (under 20 U/min) with a surgical drill.

8a MECHANICAL INSERTION OF THE IMPLANT

A plate shaped indention of approximately 2 mm in depth is then performed in the bone surface /according to the previous marking/ with a ball round bur. The recommended rpm is 1500 to 2000 U/min for burring.
Any inequality of the alveolar ridge must be smoothed at the intended site of the implantation.

Implant body driver with power drive

Manual implant body driver

8b MANUAL INSERTION OF THE IMPLANT

The implant is driven into the nest by the help of manual drivers (8034-059, 8455-078), and only the polished part of the head piece of the implant is raised above the bone surface. The mechanical drivers can be converted into manual ones by the help of the 8000-002 and 8000-102 adapters.

9 INSERTION OF THE HEALING SCREW AND WOUND CLOSURE

The implant body is covered with a healing screw, and then the mucoperiosteum is sealed around the neck of the implant with tension-free knotted sutures.

For more detail see the SURGICAL MANUALS available on dentisystem.com

DOWNLOAD
SurgicalOnePhaseManual(EN)-DR+.pdf

DR+ TWO STAGE SURGICAL PROTOCOL

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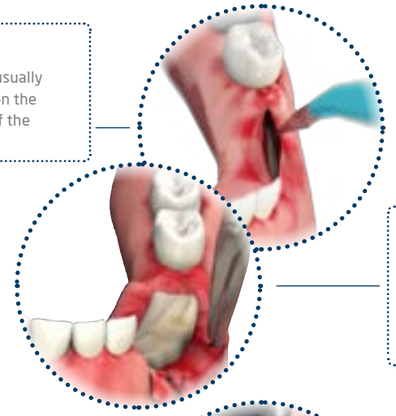

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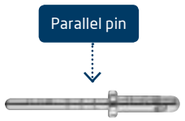

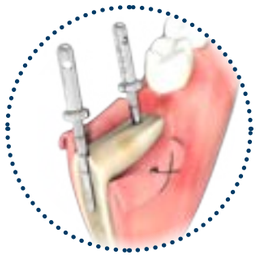

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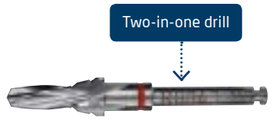

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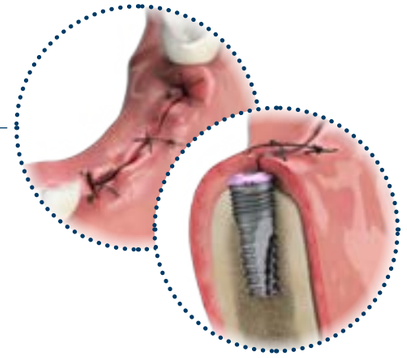


The implant body is covered with a healing abutment, and then the mucoperiosteum is sealed around the neck of the implant with tension-free knotted sutures.

DR+ TWO STAGE SURGICAL PROTOCOL

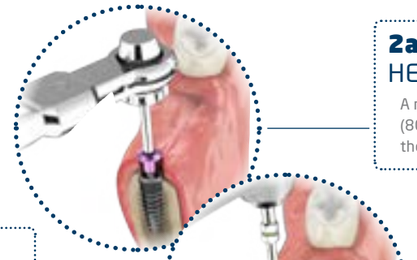
SECOND SURGICAL INTERVENTION

A drawback of the two stage surgical protocol is that a second surgical intervention should be performed to uncover the implants from the mucoperiosteum. A gingiva shaper is driven into the implants to promote the epithelial junction in the peri-implant sulci.

1 EXPLORATION

To uncover the neck piece of the implants, a gingiva cutter is used to excise the mucosa (8000-053, 8000-153).
The gingiva is incised in a circle with a gingiva cutter centrally to the abutment, and then the circularly incised part is removed.

gingiva cutter



2a REMOVAL OF THE HEALING SCREW (MANUAL)

A manual dental torque wrench (8000-076, 8000-077) is used to remove the healing screw.

2b REMOVAL OF THE HEALING SCREW (MECHANICAL)

Both the manual and the mechanical dental torque wrenches can be used (8000-076) to remove the healing screw.

3a FORMING OF THE GINGIVA

After removing the healing screw, a gingiva former is driven into the implant until the wound in the mucosa is healed. The dental torque wrench used in the previous step can be applied to insert them.
For further information on gingival shapers See the prosthetic manual / gingiva shapers.

3b FORMING OF THE GINGIVA

The gingiva former should stay in the jaw until the prosthetic work begins.

For more detail see the SURGICAL MANUALS available on dentisystem.com

DOWNLOAD
SurgicalTwoPhaseManual(EN)-DR+.pdf

PROSTHETICS

MILLING CENTER

With an experience of 20 years in dental techniques, our team introduced CAD-CAM technology in the DentMill Milling Center in June 2011. Our goal is to serve dental technicians and dentists at the highest level from design to production . Our Milling Center offers services on the whole scale of CAD-CAM technology. We design crowns, bridges, implants based prosthesis, individual structures, dental bars, inlays, inlay anchorage with high quality optical scanners.

dentmill MILLING CENTER

WHAT KIND OF SERVICES DO WE OFFER?

- scanning samples
- designing prosthesis with CAD software
- receiving variety of data from open systems, designed replacement
- 3M LAVA zircon planning, receiving data only here in Hungary
- production from titanium, zircon, CoCr
- all above from the highest quality materials, on professional machines and with skilled colleagues

MATERIALS WE WORK WITH

- ceramic fused and anatomic bridge structure from titanium, zircon and CoCr materials
- solo or zircon bridge piers inlay, onlay
- designing and producing individual implant abutments from zircon and titanium
- hybrid replacements, primer piers of titanium directly attached to the implant platform
- complete dental work starting from negatives
- ceramic frames of zircon, titanium, CoCr on sectional sample
- receiving data from CAD software and milling of zircon, titanium, CoCr
- professional, technical assistance to DWOS scanners

WWW.DENTMILL.HU

ABUTMENT DESIGN

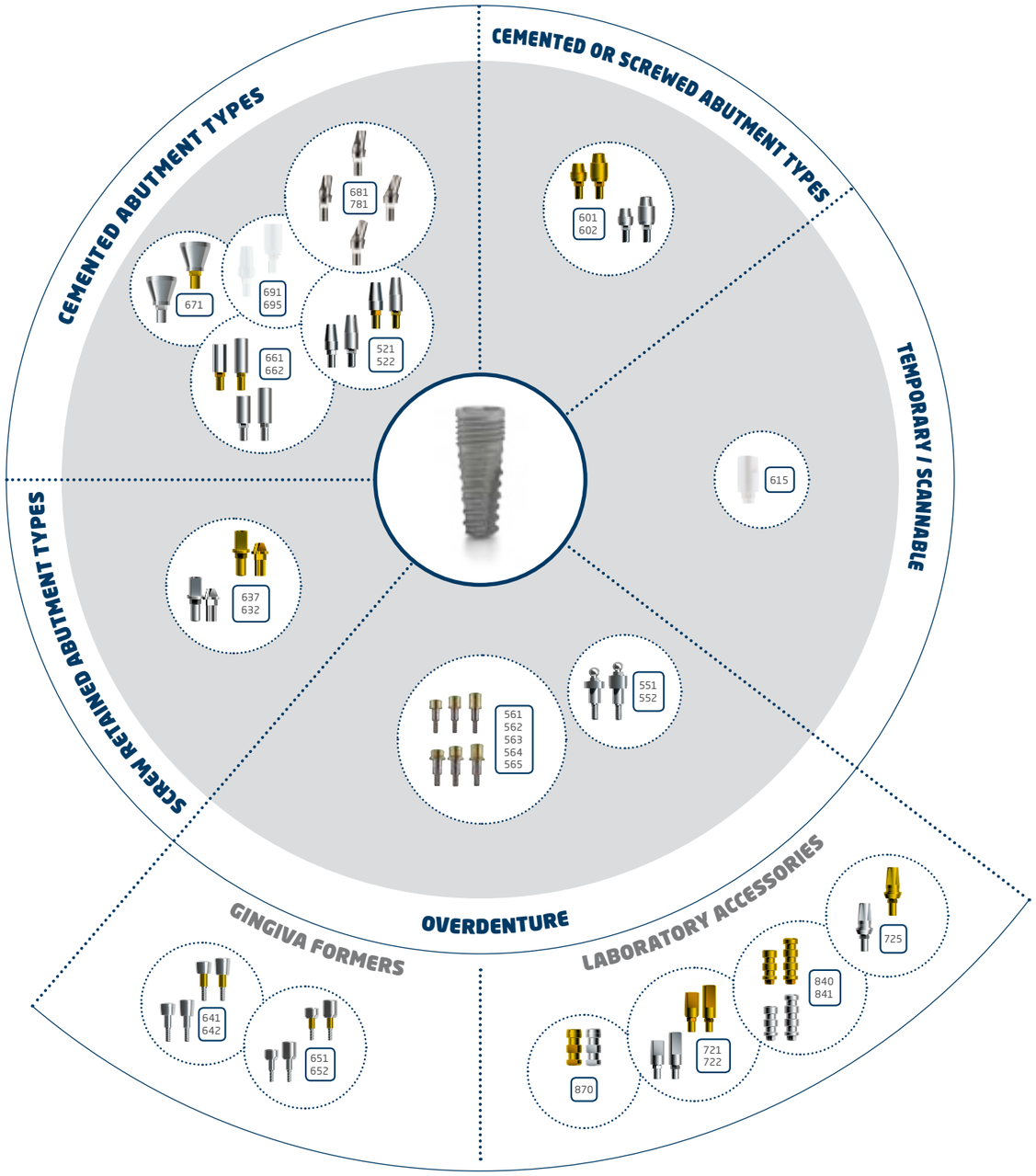
When the hexagonal positioning part of the head part fits into the hexagonal inside of the implant, there is a small click and the abutment becomes stable. After that, the fixing screw is screwed in and tightened with the torque ratchet through the abutment. The torque recommended for tightening of the abutment should be 25 Ncm and 30 Ncm for ball heads. Abutments are made of unalloyed titanium. The neck part of the abutments is smooth polished to ensure the development of the peri-implant permucosal seal. The low neck part encloses a threaded inner hole for fixing the screw that retains the prosthesis.

DENTI® ROOT FORM PLUS IMPLANT DESIGNS

COLOR CODE AND SIGNS

The purpose of color-coding is to make usability easier in surgical procedures as well as in prosthetic and laboratory stages.
The Denti® color codes are obtained by an electro-chemical reaction process without any addition of materials and thus without any change to the titanium properties.
The Denti® Root Form+ system are available in 4 diameters in order to meet different needs of the patients..

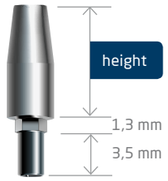
The chart indicates which color corresponds to each Denti® implant diameter	
Diameter of Root Form Implants (mm)	Color code
3.3; 3,8	yellow
4,3 - 4,8 - 5,3	natural



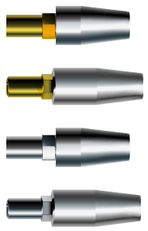
DENTI® CEMENTED ABUTMENT TYPES

UNI ABUTMENT

The conical, universal abutments can be used for the preparation of both individual crowns and bridges.



Universal abutment		
Uni abutment	Implant Ø (mm)	Height (mm)
2034-521	3,3; 3,8	7,5
2034-522	3,3; 3,8	9,5
2455-521	4,3; 4,8; 5,3	7,5
2455-522	4,3; 4,8; 5,3	9,5



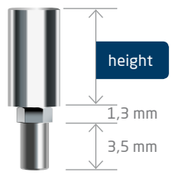
Instrument to use with:

- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual

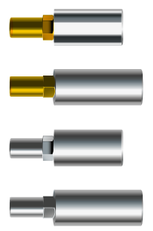
DENTI® CEMENTED ABUTMENT TYPES

CYLINDRICAL ABUTMENT FOR ANGLED CORRECTION

The cylindrical abutment is recommended whenever there is a need to correct axial deviations in case any implants are not adequately parallel with each other. It can be used for the preparation of both individual crowns and bridge and can be corrected by means of milling whenever necessary.



Cylindrical abutment		
Reference No.	Implant Ø (mm)	Height (mm)
2034-661	3,3; 3,8	7,5
2034-662	3,3; 3,8	9,5
2455-661	4,3; 4,8; 5,3	7,5
2455-662	4,3; 4,8; 5,3	9,5

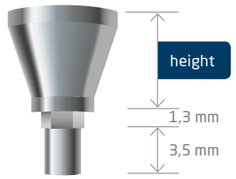


Instrument to use with:

- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual

REVERSE CONICAL ABUTMENT FOR ANGLED CORRECTION

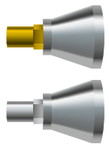
Denti® reverse conical abutments are recommended whenever there is a need for the correction of considerable axial deviations in case implants could not be prepared in an adequately parallel position. It can be used for the preparation of both individual crowns and bridges and can be reshaped by means of milling whenever necessary.



Universal abutment		
Reference No.	Implant Ø (mm)	Height (mm)
2034-671	3,3; 3,8	7,5
2455-671	4,3; 4,8; 5,3	7,5

Instrument to use with:

- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual



DENTI® CEMENTED ABUTMENT TYPES

ANATOMIC ANGULATED ABUTMENT

HEAD PIECES FOR 15 TO 25-DEGREE AXIS CORRECTION

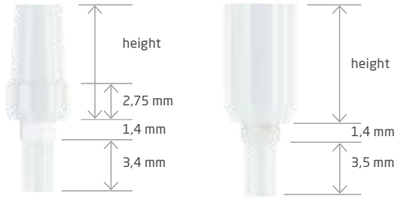
The head pieces available in the Denti Implant System for 15- to 25-degree axis correction have a circular shoulder and a curved shape, in accordance with the curve of the gingival margin. Head pieces with neck heights that are appropriate for the thickness of the mucoperiosteum can be selected during the second surgery. Correction head pieces are available with a neck height of 2.5 mm. Their correct use must be ensured even during the placement of the implant by positioning the head piece properly. To facilitate this, plastic sample head pieces are available.

Anatomic angulated abutment			
Reference No.	Implant Ø (mm)	Height (mm)	Degree
2034-681	3,3; 3,8	8,0	-15°
2034-781	3,3; 3,8	8,0	-25°
2455-681	4,3; 4,8 ; 5,3	8,0	-15°
2455-781	4,3; 4,8 ; 5,3	8,0	-25°

Plastic sample head pieces for correction head pieces	
Reference No.	Height (mm)
2455-777	8,0

CIRCON ABUTMENTS

Advantages of using Denti® circon abutments: the screw tightening the abutment is positioned deep into the head part. Thus, it does not turn the transparent crown gray. These abutments can be processed by means of traditional dental techniques. They are not fragile even despite their small size and can resist chewing tension in a flexible manner.
Circon abutments are available in two types: cylindrical and conical.



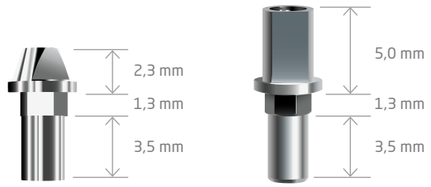
Circon abutment

Reference No.	Implant Ø (mm)	Height (mm)
2034-691	3,3; 3,8	8,0
2034-695	3,3; 3,8	8,0
2455-691	4,3; 4,8; 5,3	8,0
2455-695	4,3; 4,8; 5,3	8,0

Instrument to use with:
- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual

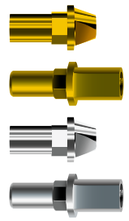
INTERMEDIARY ABUTMENTS

Denti® intermediary abutments are especially useful when treating tight edentulous gaps. Reshaping of the crowns and bridges can be carried out by prefabricated burnout plastic cylinder.



Intermediary abutments

Reference No.	Implant Ø (mm)	Height (mm)
2034-637	3,3; 3,8	2,3
2034-632	3,3; 3,8	5,0
2455-637,	4,3; 4,8; 5,3	2,3
2455-632	4,3; 4,8; 5,3	5,0



Instrument to use with:
- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual

BURNOUT PLASTIC CYLINDERS

The burnout cylinders provide full flexibility when selecting the method for single tooth restorations and multiple constructions.
Denti® burnout plastic cylinders are recommended for the precise copying of the intermediary abutments (Reference No.: 2455). Castable plastic cylinders are available in two types. Cylindrical types (-009) are recommended for several interposed abutments. Flat sided types are recommended for single or telescopic construction.

Burnout plastic cylinders

Reference No.	Implant Ø (mm)	
3034-009	3,3; 3,8	cylindrical
3034-010	3,3; 3,8	flat sided
3455-009	4,3; 4,8; 5,3	cylindrical
3455-010	4,3; 4,8; 5,3	flat sided

TEMPORARY/SCANABLE ABUTMENT

Due to the material of the temporary abutment (WHITE PEEK), it can be applied in dental scanners. For the construction of immediate, temporary prostheses, we recommend the use of temporary head pieces made of hard plastic (PEEK). The head piece is secured into the implant with a through bolt. After securing the head piece, it may be grinded in, as necessary, with a turbine in the mouth, and the impression for the temporary immediate prosthesis can be taken immediately. The completed crown/bridge is fixed to the head piece with adhesive cement.
Immediate head pieces are not recommended for long-term loading.

Scanable abutment

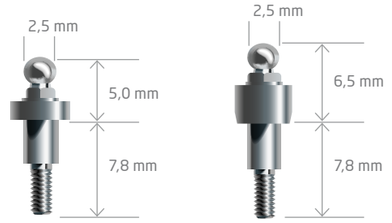
Reference No.	Implant Ø (mm)	Height (mm)
2034-615	3,3; 3,8	8
2455-615	4,3; 4,8; 5,3	8

Instrument to use with:
- 8000-002 hand adapter for ratchet
- 8000-143 torque ratchet

DENTI® REMOVABLE ABUTMENT TYPES

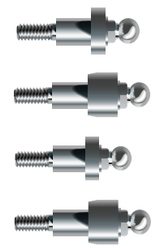
BALL ABUTMENT

The ball abutments are available in 2 heights. The ball abutment is made of titanium alloy TiAlV. The radius of the ball is 2 mm.
Ball abutments do not have internal hex connection.



Anatomic angulated abutment

Reference No.	Implant Ø (mm)	Height (mm)	Ball Ø (mm)
2034-551	3,3; 3,8	5,0	2,5
2034-552	3,3; 3,8	6,5	2,5
2455-551	4,3; 4,8; 5,3	5,0	2,5
2455-552	4,3; 4,8; 5,3	6.5	2,5



Instrument to use with:
- 8000-002 hand adapter for ratchet
- 8000-143 torque ratchet

Accessories

Reference No.	
3000-880	OT CAP retaining cap (standard)
3000-881	OT CAP retaining cap
3000-882	OT CAP retaining cap ((soft))
3000-883	OT CAP retaining cap worn out, undersized inside diameter sphere

DENTILOC ABUTMENTS

Dentiloc abutment provides spherical retention fixation. It can mainly be utilized in implants with larger axial deviation, where excellent fixation can be provided even at 40' axial deviation.
Another advantage is its low height resulting in smaller space requirement. Compared to ball-head assembly, it has a larger retention surface, better prevention of loosening, and it is more resistant to wear-and-tear due to the larger interface.

Anatomic angulated abutment

Reference No.	Implant Ø (mm)	Height (mm)
R 2034-561	3,3; 3,8	2,7
R 2034-562	3,3; 3,8	3,7
R 2034-563	3,3; 3,8	4,7
R 2034-564	3,3; 3,8	5.7
R 2034-565	3,3; 3,8	6.7
R 2455-561	4,3; 4,8; 5,3	2.7
R 2455-562	4,3; 4,8; 5,3	3.7
R 2455-563	4,3; 4,8; 5,3	4.7
R 2455-564	4,3; 4,8; 5,3	5.7
R 2455-565	4,3; 4,8; 5,3	6.7

Instrument to use with:
- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual

DENTI® REMOVABLE ABUTMENT TYPES

NOVALOC SWISS MADE

Anatomic angulated abutment

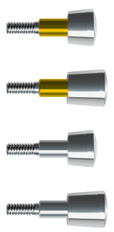
Reference No.		pcs.
2010.101	2010.101 Equipment Box with 3 tools 2010.731, 2010.741,2010.751 (excluding consumadles)	
2010.601	Processing package Titanium Titanium matrix housing incl. PEEK mounting insert black. Retention inserts white, yellow, green Mounting collar silisone	2
2010.611	Processing package PEEK PEEK matrix housing incl. PEEK mounting insert black Retention inserts white, yellow, green Mounting collar silicone	2
2010.710	Retention insert white PEEK, pull-off weight	4 extra-ligh
2010.712	Retention insert yellow PEEK, pull-off weight	4 medium
2010.713	Retention insert green PEEK, pull-off weight	4 strong
2010.714	Retention insert blue PEEK, pull-off weight	4 extra-strong
2010.715	Retention insert black. PEEK, pull-off weight	4 ultra strong
2010.731	Demounting tool for mounting insert+ model analogue reposition aid, blue	
2010.741	Mounting and demounting tool for retention inserts, brown	
2010.751	Matrix housing extractor grey	
2010.703	Titanium matrix housing with attachment option incl. PEEK mounting insert	4
2010.701	Titanium matrix housing incl. PEEK mounting insert	4
2010.702	PEEK matrix housing incl. PEEK mounting insert	4
2010.725	PEEK mounting insert black	4
2010.721	Model analog Aluminium	4
2010.722	Forming/fixing matrix, red PEEK	4
2010.723	Processing spacer, white POM C	4
2010.724	Mounting collar silicone	10

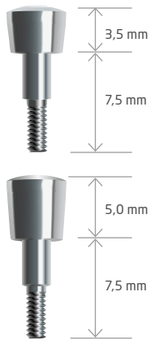
GINGIVA FORMERS

The gingiva formers are designed to use for the formation of a new periimplant gingival cuff. Denti® gingiva formers are designed with two heights. The appropriate one to choose is the one, which is approx. 1 mm longer than the mucoperiosteum. Thus when tightened, it will protrude into the mouth.
In case of implants assembled by a two-stage surgery method, we recommend the use of Denti® gingiva formers for the healing period after the second operation.
The so-called transgingival healing screws allow for the aesthetic shaping of the gingiva around the implant. These can be used when implants are assembled by a half-open, half-closed operation method.

Transgingival healing screw

Reference No.	Implant Ø (mm)	Height (mm)
2034-641	3,3; 3,8	3,5
2034-642	3,3; 3,8	5,0
2455-641	4,3; 4,8; 5,3	3,5
2455-642	4,3; 4,8; 5,3	5,0

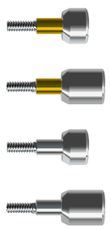


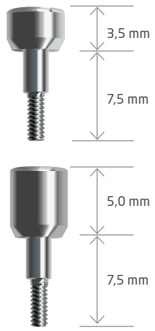


Gingiva former

Reference No.	Implant Ø (mm)	Height (mm)
2034-651	3,3; 3,8	3,5
2034-652	3,3; 3,8	5,0
2455-651	4,3; 4,8; 5,3	3,5
2455-652	4,3; 4,8; 5,3	5,0

Instrument to use with:
- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual

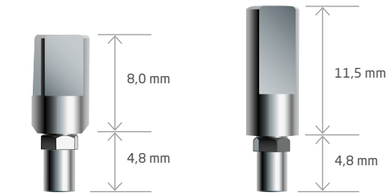




LABORATORY ACCESSORIES IMPRESSION TAKING

Transmitting the structure of implants to the specimen must be extremely precise. By using Denti® Implant Systems, taking a dental impression can be done with open or closed tray. The method of impression taking (open or closed tray) is dependent on the parallelism, number and angulation of the implants as well. There are appropriate impression copings for both impression methods. Each element is color coded according to the diameter of the implant.
A precise copy of the intraoral conditions can be made by the means of impression copings and labor analogues.

CLOSED TRAY IMPRESSION COPINGS



The closed tray impression copings are available in two heights.

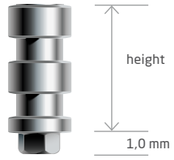
Closed tray

Reference No.	Implant Ø (mm)	Height (mm)
3034-721	3,3; 3,8	8,0
3034-722	3,3; 3,8	11,0
3455-721	4,3; 4,8; 5,3	8,0
3455-722	4,3; 4,8; 5,3	11,0

Instrument to use with:
- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual

OPEN TRAY IMPRESSION COPINGS

Denti® laboratory components allow an impression to be taken from the fixture. The impression copings and the laboratory analogues ensure correct transfer of the situation to the model.

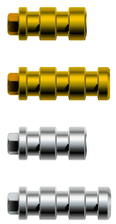


Open tray impression copings		
Reference No.	Implant Ø (mm)	Height (mm)
3034-840	3,3; 3,8	7,5
3034-841*	3,3; 3,8	9,5
3455-840	4,3; 4,8; 5,3	7,5
3455-841*	4,3; 4,8; 5,3	9,5

*special order

Instrument to use with:

- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual



CONICAL IMPRESSION COPING WITH CAP

The indirect, closed-tray impression technique is used when the placed implants show only a small degree of axis deviation (9 to 10° at most). This impression technique is primarily recommended for partial or subtotal tooth loss. An A-silicone or polyester impression material is recommended for obtaining the impression. Use hard plastic factory or custom trays for obtaining the impression. If possible, use a one-stage or a simultaneous two-stage impression technique.

Open tray impression copings		
Reference No.	Implant Ø (mm)	Height (mm)
3034-725	3,3; 3,8	8
3455-725	4,3; 4,8; 5,3	8

Instrument to use with:

- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual

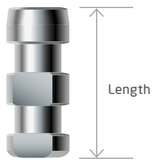
IMPRESSION CAP

Impression cap		
Reference No.	implants	Height (mm)
34724	3034-725 3455-725	8

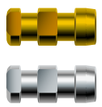
Three items per pack

LABOR ANALOGUES

Unalloyed titanium.
Analogous to the body of the fixture.



Labor analogues		
Reference No.	Implant Ø (mm)	Length (mm)
3034-870	3,3; 3,8	9,5
3455-870	4,3; 4,8; 5,3	9,5



SCREWS

Table of prosthetic screws

Fixing screws		
Reference No.	Size (mm)	Application
3034-160	10,0	2034-521, 2034-522
	10,0	2034-661, 2034-662
	10,0	2034-671
	10,0	2034-601-2034-602
3034-161	10,8	2034-691, 2034-695
3034-120	12,8	2034-637
3034-122	15,2	2034-632
3455-180	10,0	2455-501,-502
	10,0	2455-521,-522
	10,0	2455-661,-662
	10,0	2455-671
	10,0	2455-601,-602
3455-181	10,8	2455-691,-695
3455-121	12,8	2455-637
3455-123	15,2	2455-632
2455-571	3,0	2455-501,-502 2034-601,-602; 2455-601,-602
2455-572	4,0	2455-501,-502 2034-601,-602; 2455-601,-602
2455-573	1,8	2455-501,-502 2034-601,-602; 2455-601,-602
3034-111	15,7	3034-721
3034-112	18,7	3034-722
3455-114	15,7	3455-721
3455-115	18,7	3455-722
3034-140	21,5	3034-840
3455-141	21,5	3455-840
Healing screws		
1034-001	-	For DR+ Ø 3,3 - 3,8 mm implants
1455-002	-	DR+ Ø 4,3-4,8-5,3 mm implants

Instrument to use with:

- 8000-076 screwdriver with power drive
- 8000-077 screwdriver, manual

STEPS OF THE OPEN-TRAY (DIRECT) IMPRESSION TECHNIQUE

The open-tray impression technique is primarily advised when the placed implants show a considerable axis deviation (> 9 to 10°) but, because of its high precision, its use can be recommended in every case. This impression technique is recommended for partial or complete tooth losses.
An A-silicone or polyester impression material is recommended for obtaining the impression. Use a hard plastic factory or custom tray for obtaining the impression. If possible, use a one-stage or a simultaneous two-stage impression technique.

1. PREPARING THE MOUTH FOR TAKING THE IMPRESSION

For 1 week after the release of the implant, the patient will wear a healing abutment, which ensures the optimal shaping of the peri-implant gingiva. This method is primarily recommended in the case of implants placed with the two-stage surgical procedure. A healing abutment is not required if a one-stage surgical procedure was performed. After that, a situation impression is obtained for the custom tray.

GINGIVA FORMERS

2. OBTAINING THE IMPRESSION

After the regeneration of the gingiva (about 7 to 10 days), the healing abutment(s) is (are) removed, the impression post suitable for the open-tray impression technique is placed and secured with a long fixation screw, and then the custom tray is tried in its place. The custom tray should be laid over the impression abutment so that the fixation screws extend 1 to 2 mm beyond the openings on the occlusal side of the tray.

OPEN TRAY IMPRESSION COPINGS

3.

The tray filled with the impression material is placed in the mouth. Make sure to have the custom tray seated exactly in its place.

4.

The set impression can be removed from the mouth only by screwing off the long screws that secure the impression posts through the openings on the occlusion side of the tray, and then pulling them out, one by one, of the implants. After removing the impression tray from the mouth, we can see that the retentive impression abutments have remained secured in the impression. After that, the healing abutment(s) – or, in case of implants placed with the one-stage surgical procedure, the transgingival screw(s) – is (are) placed back in the mouth, and the implants are closed.

5.

Before casting the impression, the laboratory implants can be precisely and safely inserted in the impression abutments, and secured strongly with the long fixation screws going through them. This latter step can also be performed in the dental laboratory.

From that point on, the procedure is the same as the one used for the construction of prostheses in the dental practice.
The steps of obtaining the impression are independent of the type of the prosthesis. The open-tray impression technique may be suitable for the construction of any implant-based prosthesis (cemented, screwed-in or removable). There is no difference between the different prostheses in regard to the steps of obtaining the impression and the use of the prosthetic accessories.

STEPS OF THE CLOSED-TRAY (INDIRECT) IMPRESSION TECHNIQUE

The indirect, closed-tray impression technique is used when the placed implants show a relatively small degree of axis deviation (9 to 10° at most). This impression technique is primarily recommended for partial tooth loss. An A-silicone or polyester impression material is recommended for obtaining the impression. Use a hard plastic factory or custom tray for obtaining the impression. If possible, use a one-stage or a simultaneous two-stage impression technique.

1. PREPARING THE MOUTH FOR TAKING THE IMPRESSION

For 1 week after the release of the implant (second surgery), the patient will wear a healing abutment, which ensures the optimal shaping of the peri-implant gingiva. This method is primarily recommended in the case of implants placed with the two-stage surgical procedure. A healing abutment is not required if one-stage surgical procedure was performed.

GINGIVA FORMERS

2. OBTAINING THE IMPRESSION

After the regeneration of the gingiva (about 7 to 10 days), the healing abutment(s) is (are) removed, and the impression post(s) is (are) placed and secured in the implant(s) with a through fixation bolt.

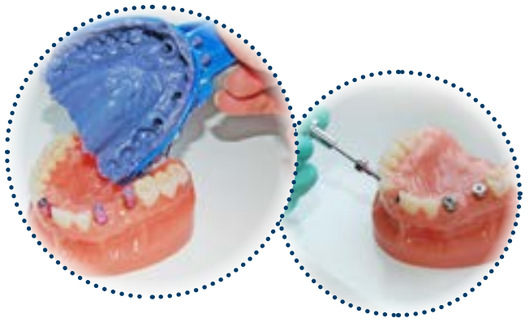
CLOSED TRAY IMPRESSION COPINGS

3.

The tray filled with the impression material is placed in the mouth.

4.

The set impression is removed from the mouth, and then the impression posts are screwed out of the implants one by one. After this, the healing abutments or transgingival healing screws are placed back into the implants in the mouth, closing the implants.



5.

Before casting the impression, the laboratory implants are secured onto the impression posts removed from the mouth with the through bolt. The obtained impression post-laboratory implant assembly must be placed back carefully into the impression. It may be performed in the office or in the dental laboratory.

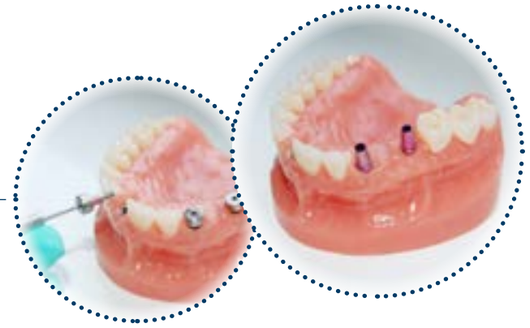
From that point on, the procedure is the same as the one used for the construction of prostheses in the dental practice.
The steps of obtaining the impression are independent of the type of the prosthesis. The closed-tray impression technique may be suitable for the construction of any implant-based prosthesis (cemented, screwed-in or removable). There is no difference between the different prostheses in regard to the steps of obtaining the impression and the use of the prosthetic accessories.

STEPS OF THE CLOSED-TRAY (INDIRECT) IMPRESSION TECHNIQUE WITH THE USE OF AN IMPRESSION CAP

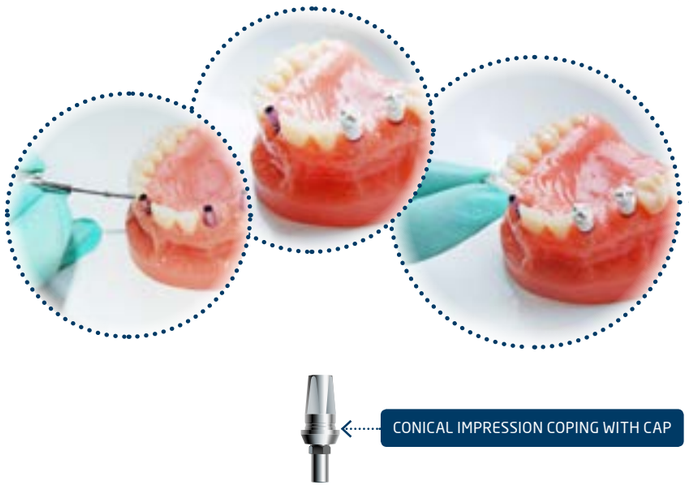
The indirect, closed-tray impression technique is used when the placed implants show only a small degree of axis deviation (9 to 10° at most). This impression technique is primarily recommended for partial or subtotal tooth loss. An A-silicone or polyester impression material is recommended for obtaining the impression. Use hard plastic factory or custom trays for obtaining the impression. If possible, use a one-stage or a simultaneous two-stage impression technique.

1. PREPARING THE MOUTH FOR TAKING THE IMPRESSION

For 1 week after the release of the implant (second surgery), the patient will wear a healing abutment, which ensures the optimal shaping of the peri-implant gingiva. This ethod is primarily recommended in the case of implants placed with the two-stage surgical procedure. A healing abutment is not required if a one-stage surgical procedure was performed.



GINGIVA FORMERS



2. OBTAINING THE IMPRESSION

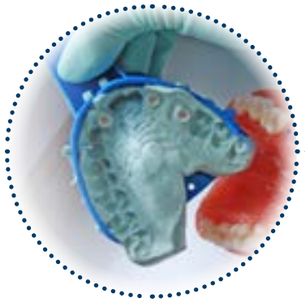
After the regeneration of the peri-implant gingiva (about 7 to 10 days), the healing abutment(s) is (are) removed, and the No. 725 abutment(s) suitable for obtaining the impression with a cap is (are) placed and secured in the implant with a fixation screw.

CONICAL IMPRESSION COPING WITH CAP

3.

After that, the impression caps (Catalog No.: 34724) are secured in the proper position by pressing them onto the abutments.

34724
H: 8mm



4.

The tray filled with the impression material is placed in the mouth.

5.

After setting, the impression is removed from the mouth. The impression caps with the retention screw stay secured in the impression material.

6.

Then, the impression components (abutments) are screwed out of the implants one by one, and are placed back on the caps in the impression material by applying light pressure. After that, the healing abutments or the transgingival healing screws are placed back into the implants in the mouth, closing the implants this way.

From that point on, the procedure is the same as the one used for the construction of prostheses in the dental practice. The steps of obtaining the impression are independent of the type of the prosthesis. The impression technique with the use of an impression cap may be suitable for the construction of any implant-based prosthesis (cemented, screwed-in or removable). There is no difference between the different prostheses in regard to the steps of obtaining the impression and the use of the prosthetic accessories.

SURGICAL TRAY - PROFESSIONAL KIT

PROFESSIONAL SURGICAL SET

The surgical instrument trays can be used for the storage and sterilization of the instruments that belong to the implants of each system.
The surgical trays are made up of the following components: basic tray, holder, cover, colored rings, which may be exchanged when damaged.

SIZE OF THE Professional SURGICAL TRAY:
199 mm x 174 mm x 60 mm.

For DR+ implants	
Reference No.	
R 8265	Denti Root Form Plus surgical tray
R 8365	Denti Root Form Plus surgical set

SURGICAL TRAY - ECO KIT

ECO SURGICAL SET

SIZE OF THE ECO SURGICAL TRAY:
176 mm x 143 mm x 63,5 mm.

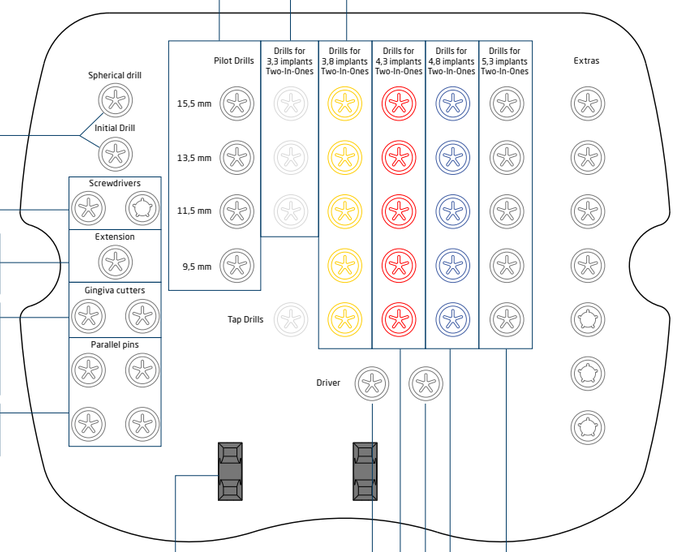
For DR+ implants	
Reference No.	
R 8264	Denti Root Form Plus ECO surgical tray
R 8364	Denti Root Form Plus ECO surgical set

PROFESSIONAL SURGICAL KIT

DR+ 3,3 | DR+ 3,8 | DR+ 4,3 | DR+ 4,8 | DR+ 5,3

DR ROOT FORM

TRANSPARENT SCALE
DR 8000-751



8010-095
8010-115
8010-135
8010-155

8000-067
8000-066

R 8033-115
R 8033-135
R 8033-155
R 8033-030

RN 8038-095
RN 8038-115
RN 8038-135
RN 8038-155
R 8038-040

RN 8043-095
RN 8043-115
RN 8043-135
RN 8043-155
R 8043-040

8000-076
8000-077

RN 8048-095
RN 8048-115
RN 8048-135
RN 8048-155
R 8048-040

RN 8053-095
RN 8053-115
RN 8053-135
RN 8053-155
R 8053-040

8000-057
8000-153
8000-053
8000-058

8000-343
8000-002

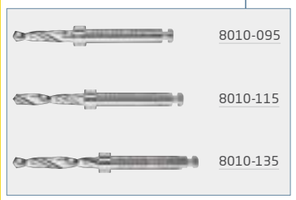
PROFESSIONAL SURGICAL KIT

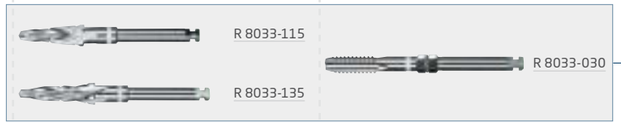
ECO SURGICAL KIT

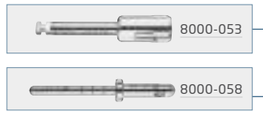
DR+ 3,3

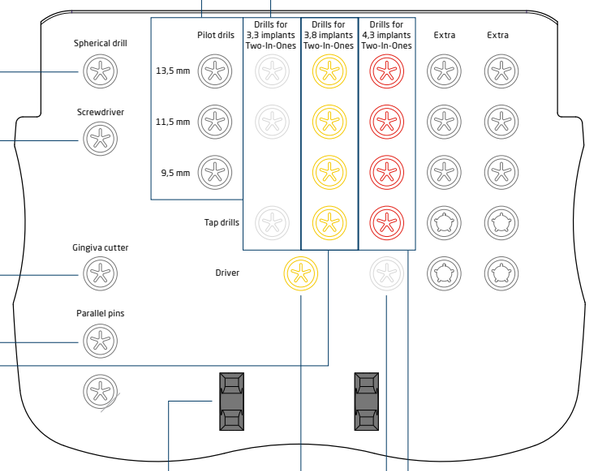
DR+ 3,8

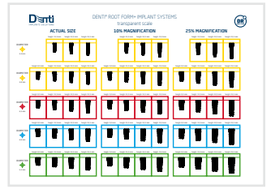
DR+ 4,3











TRANSPARENT SCALE
DR 8000-751

ECO SURGICAL KIT

SURGICAL INSTRUMENTS

Special surgical kit was developed for the insertion of Denti® Root Form implants. Drills and bone cutters with a cutting edge used for cutting the living bone are made of stainless steel meeting international standards. The instruments are designed to ensure atraumatic preparation. High primary stability is guaranteed for every quality bone.

THE DRILLS ARE:

- made from surgical stainless steel
- color coded on the shaft for quick identification
- numerically coded on the shaft
- laser marked for marking depths and diameters
- autoclavable
- available in a complete kit or as individual items
- for external irrigation
- double possibility: using with contra angle or manually

INITIAL DRILLS

For DR+ implants			
Reference No.		Ø mm	Cutting depth (mm)
8000-067	Initial drill	Ø 2,00	16,0
8000-066	Spherical drill		
8010-095	Pilot drill with stop	Ø 2,00	9,5
8010-115	Pilot drill with stop	Ø 2,00	11,5
8010-135	Pilot drill with stop	Ø 2,00	13,5
8010-155	Pilot drill with stop	Ø 2,00	15,5

TWO-IN-ONES

The Denti® two-in-one drills are designed to prepare the implant bed precisely. The size of the drills is adjusted to the length and diameter of the corresponding fixture. The profile of the two-in-one drills is designed to collect and conserve the vital bone.

For Dr+ implants 3,3 mm	
Reference No.	Cutting depth (mm)
R 8033-115	11,5 mm
R 8033-135	13,5 mm
R 8033-155	15,5 mm

For DR+ implants 3,8 mm	
Reference No.	Cutting depth (mm)
R 8038-095	9,5 mm
R 8038-115	11,5 mm
R 8038-135	13,5 mm
R 8038-155	15,5 mm

For DR+ implants 4,3 mm	
Reference No.	Cutting depth (mm)
R 8043-095	9,5 mm
R 8043-115	11,5 mm
R 8043-135	13,5 mm
R 8043-155	15,5 mm

For DR+ implants 4,8 mm	
Reference No.	Cutting depth (mm)
R 8048-095	9,5 mm
R 8048-115	11,5 mm
R 8048-135	13,5 mm
R 8048-155	15,5 mm

For DR+ implants 5,3 mm	
Reference No.	Cutting depth (mm)
R 8053-095	9,5 mm
R 8053-115	11,5 mm
R 8053-135	13,5 mm
R 8053-155	15,5 mm

TAPS

The tap drills are to be used for tapping hard, dense bone. The tap drills are designed with a thread having the same configuration as the implant. There is no tap drill for screw implants with thread height 1,25 mm as they are designed for soft bone where there is no tap drilling is needed.

DR+ implants	
Reference No.	Application
R 8033-030	for Ø3,3 mm implants
R 8038-040	for Ø3,8 mm implants
R 8043-040	for Ø4,3 mm implants
R 8048-040	for Ø4,8 mm implants
R 8053-040	for Ø5,3 mm implants

Both Denti® Root Form self threading implants may require the use of tap drills (different for each diameter) only in case of hard bone.

IMPLANT BODY DRIVERS

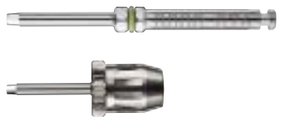
The inner shaping of the Ø 3,3-3,8 mm DR+ implants are identical, so the drivers can be equally used for these fixtures.
The inner shaping of the Ø 4,3-4,8-5,3 mm DR+ implants are identical as well, so the drivers can be equally used for these fixtures. The implant body drivers are available in two lengths:
short and long.

With short shaft		
Reference No.	Total length (mm)	Application
8034-070	24,5	For Ø 3,3-3,8 DR+ implants
8455-079	24,5	For Ø 4,3-4,8-5,3 DR+ implants

With long shaft		
Reference No.	Total length (mm)	Application
8034-170	29,5	For Ø 3,3-3,8 DR+ implants
8455-179	29,5	For Ø 4,3-4,8-5,3 DR+ implants

Optional instruments for DBL and/or for DCR systems		
Reference No.	Total length (mm)	Application
8034-059	29,5	For Ø 3,3 - 3,8 DR+ implants
8455-078	29,5	For Ø 4,3-4,8-5,3 DR+ implants

Screwdrivers			
Reference No.		Hex (mm)	Length (mm)
8000-076	Screwdriver with power drive	1,2	30,0
8000-077	Screwdriver, manual	1,2	22,0



The Denti® screwdrivers are applicable for driving in all kinds of Denti® screws

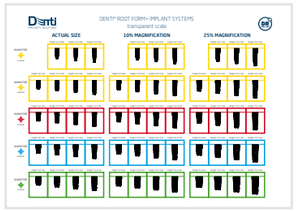
- healing screws
- fixing screws
- bridge screws
- gingiva formers
- impression copings

The surgical tray contains the screwdrivers with power drive. The manual screwdriver can be purchased separately.

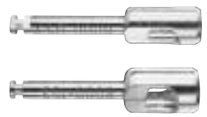
TRANSPARENT SCALE

The transparent scale is used to select the proper implant length and diameter. It can be placed on the panoramic radiograph. Available magnification rates are 1:1; 1:1,1; 1:1,25

DS	DR		
Reference No.	Reference No.		Scale
8000-751	DR 8000-751	Tranparent scale	1:1, 1:1,1 1:1,25



Standard instruments		
Reference No.		Diameter (mm)
8000-053	Gingiva cutter	4,0
8000-153	Gingiva cutter	5,0

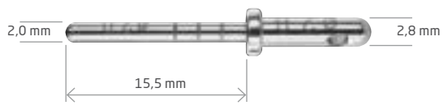


Reference No.	
8000-057	Drill extension

Reference No.	
8000-056	Titanium forceps

Reference No.	
8000-058	Parallel pin

The parallel pin is to check inclination of surgical site, parallelism as well as to check the insertion depth.



Torque ratchet	with adjustable torque
Reference No.	
8000-143	Torque ratchet
8000-002	Hand adapter for ratchet - standard
8000-102	Hand adapter for ratchet - long

The torque ratchet with hand adapter can be used:

- for implant insertion
- with tap drills
- with implant body drivers
- for fixing the ball abutment

SUGGESTED TORQUE WITH DENTI® SCREWS

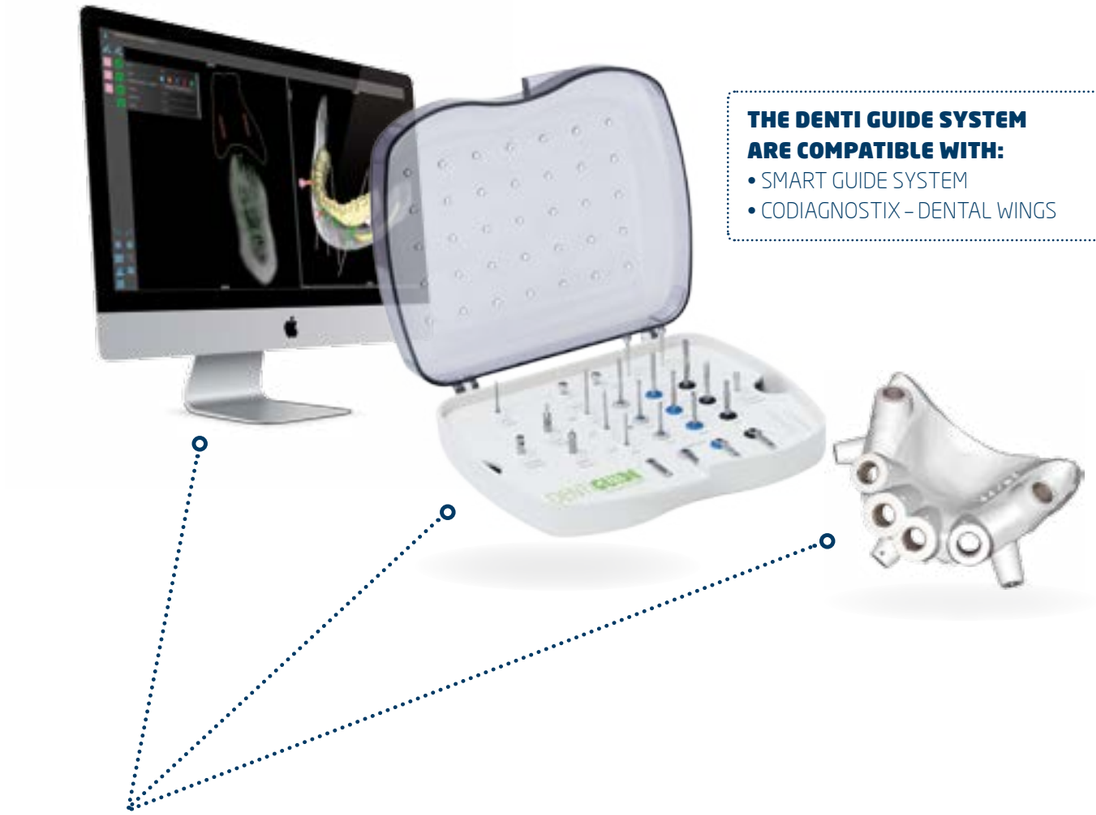
Healing screw	manual
Impression coping	manual
Gingiva formers	manual
Fixing screws	25 Ncm
Ball abutment	30 Ncm
Bar retention	30 Ncm

SURGERY – MAXIMUM TORQUE

Suggested torque with surgery		
	IN	OUT
Implant placement	50Ncm	
Checking primary stability		20 Ncm
Tap retrieving	50 Ncm	-

DENTIGUIDE

A GUIDED TECHNOLOGY
DESIGNED FOR
DENTI® IMPLANT SYSTEMS



THE DENTI GUIDE SYSTEM ARE COMPATIBLE WITH:

- SMART GUIDE SYSTEM
- CODIAGNOSTIX – DENTAL WINGS

ONE SYSTEM CONTAINING THE ENTIRE WORKFLOW

- straightforward and quick procedure for all edentulous cases
- simple planning on a clear software platform
- denti implants integrated into the software
- high quality surgical guide with a patient specific surgical protocol arriving on the following day of the first patient visit

SURGICAL PLANNING
GUIDED IMPLANT SITE PREPARATION
GUIDED IMPLANT PLACEMENT

The most advanced guided system totally fitting into your daily practice

Surgical tools completly in line with the entire technology

IMPLANT BODY SURFACE AREAS

Tooth type	Maxillary	Mandibular
	Surface area (mm2)	Surface area (mm2)
Central incisor	204	154
Lateral incisor	179	138
Canine	273	268
First premolar	234	180
Second premolar	220	207
First molar	433	431
Second molar	431	421

IMPLANT TYPE	Reference No.	Implant diameter (mm)	Implant length (mm)	Implant surface area (mm2)
DR+	R 1033-115	3,3	11,5	270
	R 1033-135	3,3	13,5	315
	R 1033-155	3,3	15,5	360
	R 1538-095	3,8	9,5	255
	R 1538-115	3,8	11,5	305
	R 1538-135	3,8	13,5	360
	R 1538-155	3,8	15,5	415
	R 1543-095	4,3	9,5	310
	R 1543-115	4,3	11,5	370
	R 1543-135	4,3	13,5	430
	R 1543-155	4,3	15,5	490
	R 1548-095	4,8	9,5	340
	R 1548-115	4,8	11,5	415
	R 1548-135	4,8	13,5	490
	R 1548-155	4,8	15,5	560
	R 1553-095	5,3	9,5	380
	R 1553-115	5,3	11,5	465
	R 1553-135	5,3	13,5	550
	R 1553-155	5,3	15,5	625

WARRANTY

GENERAL WARRANTY CONDITIONS

Since 1989, the Denti Systems Kft. produces and develops dental implants, thus provides dental experts/technicians (herein after referred to as "User") an extensive product portfolio.

Our product line, which satisfies all needs, includes almost all type of implants according to the current market requirements, starting with the two-phase system down to the single-phase titanium implants.

As a matter of course, all of our systems can be used versatilely. In interest of short and long term clinical results and success of implantation, implantologists who use our system, may find the most adequate implant family from our product portfolio.

Our products are scientifically documented and tested in everyday clinical practice for decades, and used day by day.

Below you can find the prospectus with full information of warranty conditions regarding implant systems of Denti System:

WARRANTY CONDITIONS

- The warranty applies to all implants and its components, which has been marketed by Denti System Kft. This includes implants, locking screw, healing cap and all kinds of prosthetic structures, heads and other clinical instruments.
- The warranty claims, the rights and advantages specified below may exclusively be enforced by the user of our company. The warranty is non-transferable, third-party (with the exception of product liability claim of Civil Code § 6:168), laboratories or suppliers cannot exercise it.

PERIOD OF WARRANTY

The warranty period for the products of Denti System Kft. are as follows:

- 3 years on clinical instruments and other surgical accessories;
- 5 years on components of Denti Systems Kft. made of zirconium;
- unlimited warranty on clinical components of Denti System Kft. (except components made of zirconium).

VALIDITY OF WARRANTY

The following conditions define the merit of the warranty claim regarding the products of Denti System Kft.:

- Material quality of the product:
  - » In case the material of the product is defective or it has been damaged during production and so it has come to the market - that means the product does not meet the quality requirements of the Denti System Kft. - as a matter of course our company will replace it at no additional costs.
- Osseointegration (bone-healing):
  - » The Denti System Kft. provides replacement implant for its doctors (users) at no additional costs, in case the osseointegration is not proper after the implantation. The entitlement above shall be exercisable after the adequate clinical investigation, in case the investigation exludes the possible responsibility of the doctor performing implantation and the patient.

ENFORCEMENT OF WARRANTY CLAIM

The warranty claim is enforceable, if the User fulfills the following conditions:

- A warranty claim shall be accepted, if the User notifies it to our company without any delay, but latest in 30 days after the implantation. The User is obligated to contact the regional representative of Denti System Kft. or the company's customer service centre and shall fill out a complaint sheet.
- All the relevant information regarding the complaint shall be represented on the complaint sheet and the product shall be attached as well. The claimed product shall be disinfected before returning.
- The whole documentation of the case in question shall be enclosed to the complaint sheet and the User shall prove that there was no contraindication in the given case for the patient.
- The User, who vindicates its warranty claim, shall prove that the operation has been conducted in compliance with the surgical protocol specified by Denti System Kft.
- In case of enforcement of the warranty, the costs regarding the shipping of the defective product shall be advanced by the User, while the costs concerning the shipping of the replacement product shall be advanced by the Denti System Kft. In case of substantiated warranty claim the costs above shall be borne by Denti System Kft, in case of unsubstantiated warranty claim the costs above shall be borne by the User.

LIMITATION OF WARRANTY

The Denti System Kft. is liable for warranty claims and lack of conformity exclusively in accordance with the present prospectus and § 6:157-173 of the Civil Code.

The Denti System is liable exclusively regarding the conformance of the products produced and distributed by itself, whether these products meet the legal provisions, respectively the technical requirements specified in the product specification and user guide.

Making legal statements by unauthorized third parties on behalf of the Denti System Kft either orally or in writing, which are different from above or supplement its statements, do not constitute a commitment by the Denti System Kft., thus Denti System Kft. has no responsibility regarding these statements.

The liability for damages of Denti System Kft. is excluded for any other damages in the asset of the User and for the loss of pecuniary advantage, except the damages caused intentionally and which injure human life, limb or health.

EXCLUSIONS FROM WARRANTY CLAIMS AND ITS ENFORCEMENT

No warranty claim is enforceable against the Denti System Kft. in the following cases:

- Failure concerning the implant or any other component may be originated from an accident, trauma or any other injury, which has been caused by the patient or a third party.
- The cause of the failure and the damage is a component distributed by Denti System Kft., which has been implanted in such patient, by whom the success of the operation was not guaranteed. These contraindications are (but shall not be limited to) the following:
  - » o not appropriate bone supply
  - » o circulatory, blood coagulation problems
  - » o diabetes
  - » o cancer
  - » o bite disorder
  - » o not appropriate oral hygiene
  - » o heavy smoking
  - » o daily alcohol consumption
  - » o drug addiction
  - » o in the growth phase of the the jaw and the skull bone
  - » o during pregnancy
- The cause of the failure is natural wear and tear.
- The product, in respect of which the warranty may be enforced, was directly or indirectly used together with another instrument made by another producer.

Modification of the warranty period or the possible withdrawal of warranty

Denti System Kft. reserves all rights to change the warranty conditions without any prior notice, or to revoke them. As a matter of course these modifications and revocations do not apply to those components of Denti, which have been already implanted or used, respectively to products purchased by the User before the modification.

The above conditions come into effect on: 2015

DOWNLOADS

DENTYSISTEM.COM/DOWNLOADS

IMPLANT SYSTEM MAPS
• Map - DBL (PDF)
• Map - DBL Eco (PDF)
• Map - DN (PDF)
• Map - DOP (PDF)
• Map - DR (PDF)
• Map - DR Eco (PDF)
• Map - DR+ (PDF)
• Map - DR+ Eco (PDF)

For more detail see the DOWNLOAD section available on dentisystem.com

dentisystem.com/downloads

MANUALS
• MANUAL - Surgical 1 phase DBL (PDF)
• MANUAL - Surgical 1 phase DR+ (PDF)
• MANUAL - Surgical 2 phase DBL (PDF)
• MANUAL - Surgical 2 phase DR+ (PDF)

GUIDES
• CATALOGUE - One stage implants (PDF)
• CATALOGUE - Two stage implants (PDF)
• Patient Guide (PDF)
• User guide (PDF)
• User guide - prosthetics (PDF)

MISCELLANEOUS
• EC Designe Examinatiom Certificate (PDF)
• EC Designe Examinatiom Certificate - Annex (PDF)
• EK Certificate (PDF)

DENTIGEN

WITH USING DENTIGEN

in the dental implant diagnosis a quality management can be achieved within the implantology field and economically sound success rates could be maintained.

Principlesofthe genetic testingoftheDentiGentest
The DentiGen test enables to predispose the Interleukin-1

INDICATIONS OF DENTIGEN

REFRACTORY, THERAPY RESISTANT PERIODONTITIS
The positive test result can explain why previous therapies have been ineffective and may be an indication of the initiation of an alternative therapy.

PROGRESSIVE PERIODONTITIS
The positive test result is an indication of more aggressive therapy and more frequent follow-ups.

EARLY SIGNS OF PERIODONTITIS
The test result achieved prior to the initiation of the treatment can be helpful in planning the therapy tailored to the patient's needs, in stopping the progression of the disease and in preventing 'overmedicalization'.

RELATIVES OF GENOTYPE-POSTIVE PATIENTS
It may call the attention to the importance of more intense preventive measures in genotype-positive patients even if no symptoms are present.

BEFORE A COSTLY RESTORATIVE TREATMENT
Identifying the patient's genotype may be helpful in assessing the possible risks of complications that may develop after implantation and tissue regeneration.

NONCOMPLIANT PATIENT
Patient compliance can significantly be improved if the patient is aware of the genetic risk factors.

CERTIFICATES

ISO, CE, WARRANTY

EN ISO 9001 and EN ISO 13485.
The Denti® products are CE marked.
All Denti® products undergo strict testing
by the Quality Department.

THE DENTI® SYSTEM LTD.

commits itself to a 10-year period warranty
after purchase to substitute its implants.
The procedure for substitition is detailed
in the General Terms and Conditions.

Denti
IMPLANT SOLUTIONS

WWW.DENTISYSTEM.COM