



Endo-Model® SL®

Rotational and Hinge Knee Prosthesis System

Surgical Technique

Presented by:



CE 0482

Waldemar Link GmbH & Co. KG

Barkhausenweg 10 • 22339 Hamburg, Germany

Phone +49 40 53995-0 • info@linkhh.de

www.linkorthopaedics.com

02	System Overview: Modular Components
03	Variety of Assembly
04	Indications/Contraindications

Surgical Technique

05	Preoperative Planning
06	Standard Preparation Tibia
14	Standard Preparation Femur
22	Assembling the Tibial Components
27	Optional: Patella Preparation (Patella Resurfacing)

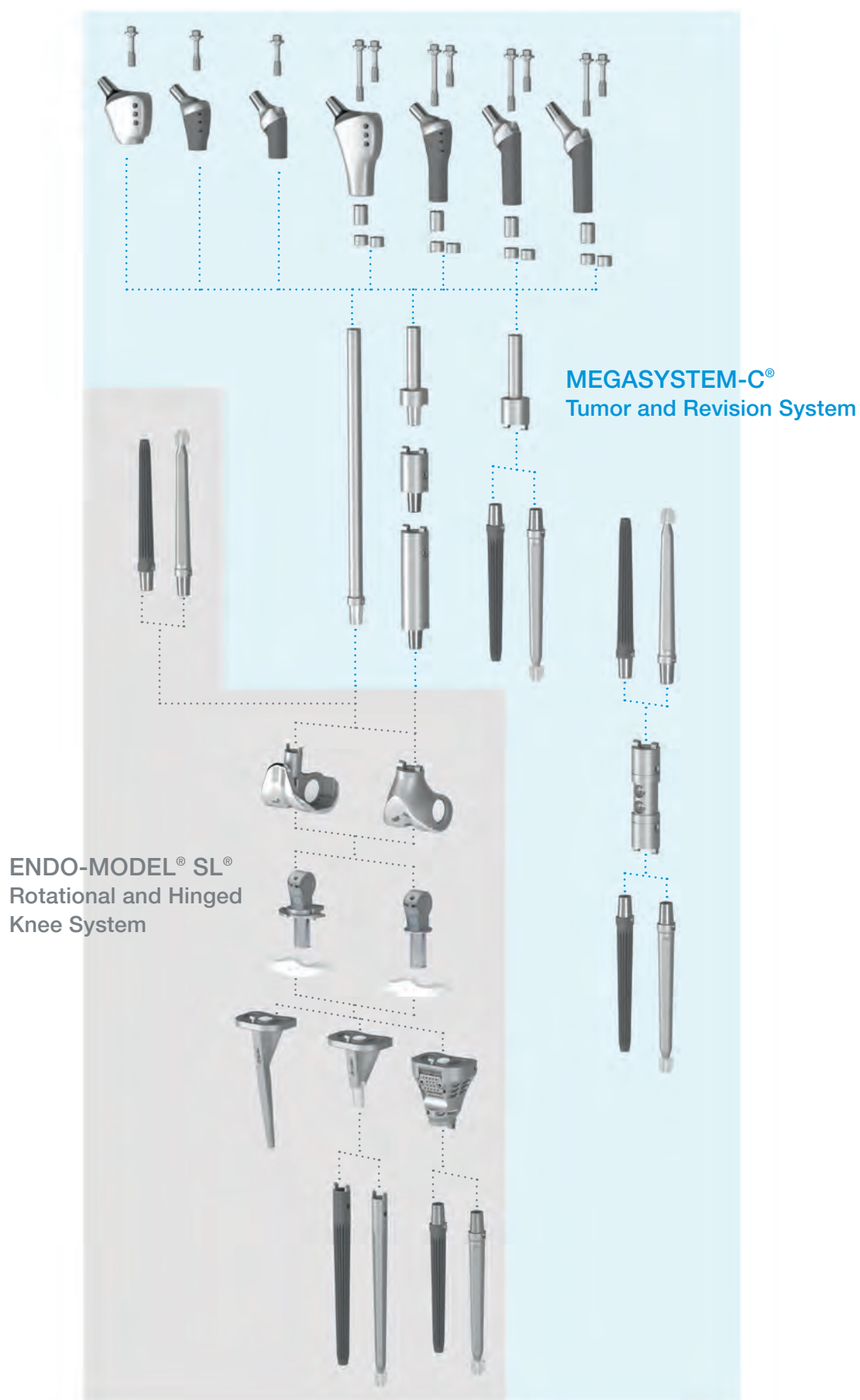
Implants

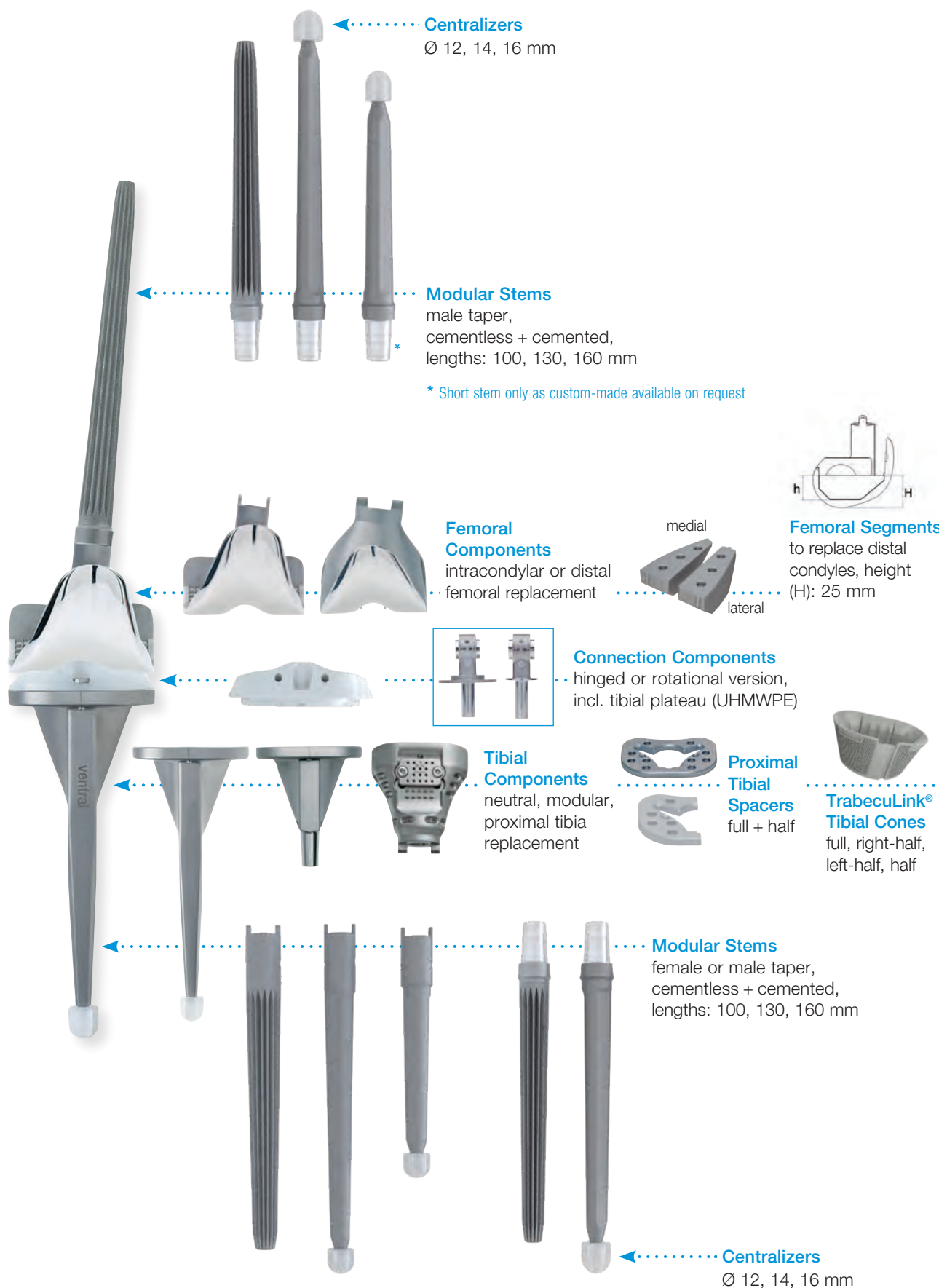
32	Joint Components Endo-Model® SL® for Rotating and Non-Rotating Hinge Knee
32	Femoral Components, intracondylar
33	Distal Femoral Replacement Components, slim
34	Femoral Segments, cemented
35	Femoral Segments, cementless
36	Tibial Components
37	Connection Components
38	Proximal Tibial Spacers, Tilastan® and UHMWPE
39	Replacement Set: Sparepart Kit Boxes for Endo-Model® SL®
40	Modular Stems with male taper, cementless + cemented
41	Modular Stems with female taper, cementless
42	Modular Stems with female taper, cemented, Modular Stems short
43	Centering Stars, LINK® Patella Components

Instruments

44	Instrument Set – Overview Containers for Endo-Model® SL® Rotating and Non-Rotating Hinge Knee
48	Container A – Tapered Reamers 100 mm
49	Container B – Tapered Reamers 130 mm
50	Container C – Tapered Reamers 160 mm
51	Container D – General Instruments
52	Container F – Femur I
53	Container G – Femur II
54	Container H – Tibia
55	Instrument Set – Femur Trial Stems 100 mm, 130 mm
56	Instrument Set – Femur Trial Stems 160 mm
57	Instrument Set – Tibial Trial Stems 100 mm, 130mm, 160 mm
58	Instrument Set – Trial Instruments for Distal Femur and Proximal Tibia Replacement
59	Instrument Set – Femur and Tibial Trial Prostheses, intracondylar
60	Container Assembling Instrument Set
61	Instruction: Assembling Instruments for Endo-Model® SL® Rotating and Non-Rotating Hinge Knee
68	Instrument Set for Patella Components
69	Additional Instruments: Adapter, Sawblades
70	Additional Instruments: Revision Drill Guide, Extractor
70	Accessories: X-ray Templates
71	Surface Modifications
72	Additional Information

Important Information





Product	Specified indications and contraindications: Endo-Model® SL® Rotational and Hinge Knee System	
	Rotational version	Hinged version
General Indications		
• Severe joint diseases with limitation of mobility due to degenerative, rheumatoid or post-traumatic arthrosis or arthritis. Joint fractures which disallow an osteosynthetic reconstruction	X	X
Indications		
• Bone necroses	X	X
• Bicondylar arthrosis by partly damaged collateral ligaments	X	-
• Bicondylar arthrosis by completely damaged ligaments and muscular instability	-	X
• Revision after primary total knee replacement	X	X
• Revision surgery after hinge knee or rotational knee joint	X	X
• Revision surgery by insufficient / inadequate bone mass	X	X
Differential Indications		
• Arthrosis of patella flange	X	X
• Valgus/Varus deformities <10°	X	X
• Valgus/Varus deformities 10–15°	X	X
• Valgus/Varus deformities 15–20°	X	X
• Valgus/Varus deformities 20–30°	-	X
• Sensitization against one or more components of used CoCrMo implant materials	-	-
Contraindications		
• Acute or chronic infections, local and systemic	X	X
• Allergies to (implant) materials	X	X
• Distinctive muscular, nerve, vascular or other diseases which put the affected limb at risk	X	X
• Insufficient / inadequate bone mass- or quality which prevents a stable anchor of the prosthesis	X	X
Relative Contraindications		
• Adiposity	X	X
• Insufficient musculature	X	-
• Lacking or foreseeable not assured compliance	X	X
• Foreseeable overload of joint prosthesis	X	X

Please note:

These indications/contraindications refer to standard cases. The ultimate decision on whether or not an implant is suitable for a patient must be made by the surgeon based on his/her individual analysis and his/her experience.

Indications/Contraindications Patella Resurfacing

Patella Resurfacing is not a necessary part of the method for implantation of a total knee replacement. It is an optional procedure carried out if femoropatellar syndrome is present.

Contraindication

The specific contraindications for the knee system being used must always be observed.

Account must also be taken of the following:

- Degree of retropatellar arthrosis
- Patient's height and weight
- Pre-existing anterior knee pain

Preoperative Planning for Endo-Model® SL® Rotational and Hinge Knee Prosthesis

Measurement tables and X-ray templates are available for the preoperative planning of revision and tumor surgery with the Endo-Model® SL® Rotational and Hinge Knee Prostheses which enable the surgeon to plan precisely the implants that will be used.

True-to-scale radiographs or precise knowledge of the actual magnification factor are the foundation for exact preoperative planning. LINK® X-ray templates show the implant illustrations in 110% magnification as standard. If different scales are desired, we will meet these wishes as far as technically possible. We provide data for digital planning on request to providers of digital planning software in the current formats.

Despite good preoperative planning, unforeseeable extensive bone loss in tumor and revision cases often presents a challenge for the surgeon. Especially in these cases the Endo-Model® SL® Rotating and Hinge Knee Prosthesis also in conjunction with the MEGASYSTEM-C® – proves its user friendliness. There is a high degree of modularity and flexibility in the adaptation of the implants to the respective bone situation.

In contrast to the use of normal hip and knee joint prostheses, management of extensive bone loss depends on the conditions in each individual situation. Structural changes in the muscles and ligaments, fixation conditions etc. increase the operative demands of tumor prostheses. Accordingly, management of extensive bone loss presents particular problems and is therefore subject to greater risk compared with the use of normal joint prostheses.

Standard Preparation Tibia



Intramedullary alignment

01

Mark the entry site with the awl (317-658/01) and open the tibial canal with the conical drill (16-3202/00).



02

Mount the awl of the previously planned length (100 mm, 130 mm or 160 mm) at the handle (15-6053/00). The impaction plate (16-3203/00) latches into the slot on the shaft of the awl.

When uncemented modular stems are used, ream with an increasing diameter until the awl makes cortical contact over a continuous distance of approx. 50 mm. The uncemented implant that will be used must correspond in length and diameter to the last awl used.

For cemented modular stems, the awl should be at least 2 mm larger than the planned stem diameter.

Important notes:

The position of the impaction plate represents the level of the joint line. Using the awls with a power tool is not permitted.



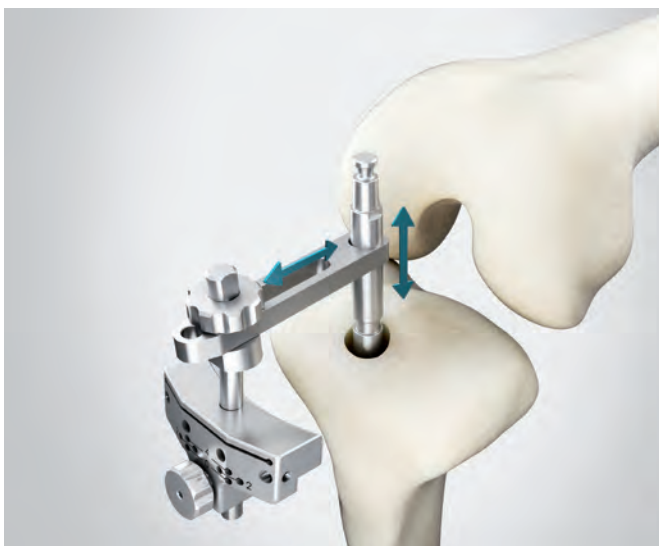
03

After the desired stability is achieved, the handle (15-6053/00) and the impaction plate (16-3203/00) are removed.



04

Attach the connector (16-3212/08) to the shaft of the awl.



05

Attach the tibial saw guide (16-3241/00) to the anterior shaft of the connector and fix it provisionally by tightening the knurled screw.



06

Attach the stylus for the tibial saw guide (317-802/52), preferably medially. The stylus tip marked 10 marks the resection level in the primary procedure (10 mm resection level). The stylus tip marked 2 can be used in revision surgery and marks a resection level of 2 mm. Alternatively, the stylus can be omitted and the resection level can be set using the cutting template (317-607/50).



07

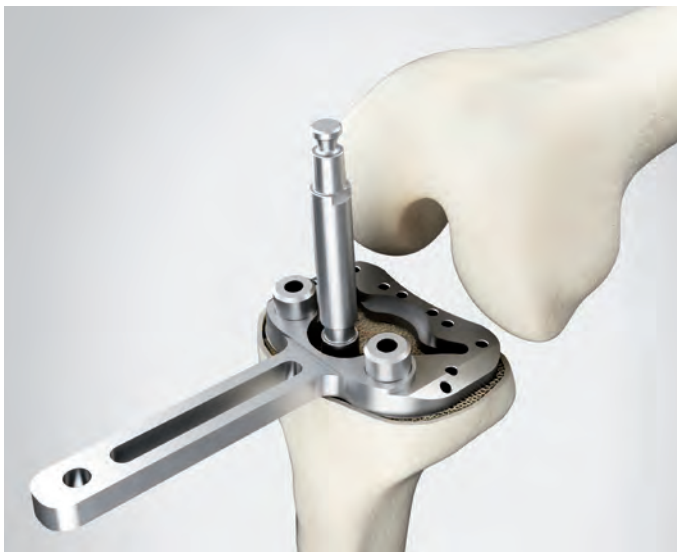
The tibial saw guide (16-3241/00) is fixed to the proximal tibia by means of two wire pins (317-585/65 or /95) through the lower row of parallel holes.



08

The bone is resected following removal of the stylus, connector and awl. The resection can be extended distally by 2 mm or 4 mm by shifting the tibial saw guide.

To achieve the correct resection geometry, sawblades with a thickness between 1.24 mm and 1.27 mm must be used.



09

The last-used awl is inserted into the medullary cavity again. By placing the drill template (16-3198/12, /13, /14) that corresponds exactly to the implant size, the definitive implant size is determined. It is important that the implant covers the resection surface as far as possible. Projection over the cortical margin of the tibia must be avoided.



10

The alignment gauge (16-3266/00) is placed over the shaft of the awl and connected to the pegs of the drill template. After rotational alignment of the drill template, it is fixed to the resection surface with at least two wire pins.

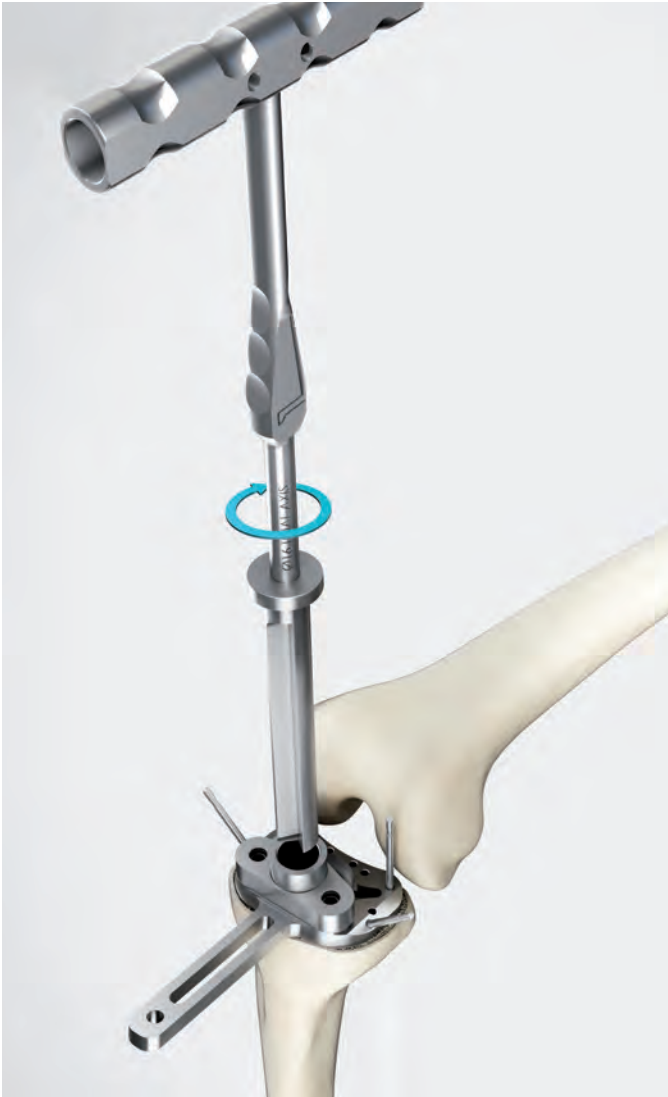


For rotational alignment, the alignment rod (16-3242/00) can be pushed from ventral onto the drill template. The ventral alignment rod must be aligned in the area between the middle of the tibial tuberosity and its medial margin.



11

Remove all of the alignment gauge. The awl Ø 24 mm must also be removed temporarily and then reinstalled.



12

Attached Ø 16 mm drill guide (16-3267/00) and drill the proximal tibia (manually or machine-operated) with the Ø 16 mm drill (16-3207/16) until stop.



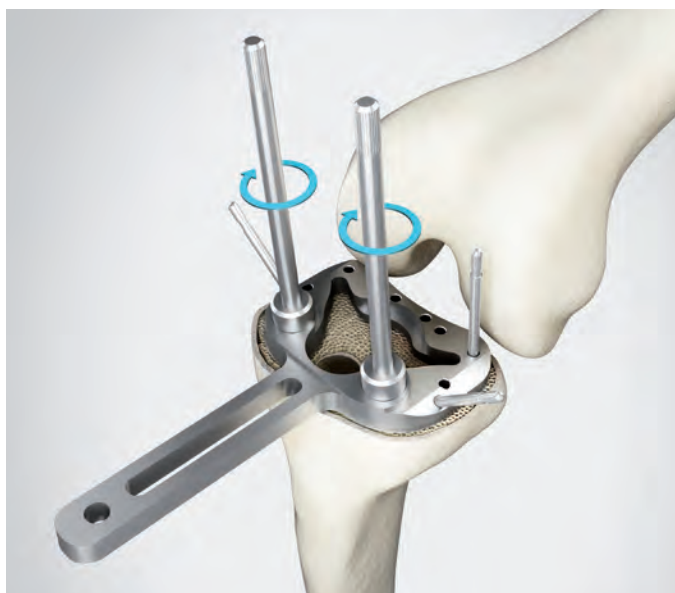
13

After removing the Ø 16 mm drill guide, the drill guide (16-3270/18, /20, /22) is attached to drill the central tibial opening. The drill guide must correspond to the size of the drill template.



14

Drill the central tibial opening manually with the drill corresponding in diameter to the drill guide (16-3208/18, /20, /22) until stop.



15

Screw the guide rods (16-3211/00) into the anterior threaded holes of the drill template.



16

Screw the stem compressor (16-3201/02, /03, /04) to the corresponding compressor (16-3199/12, /13, /14) for the proximal contour. Attach the handle (16-3197/00).



17

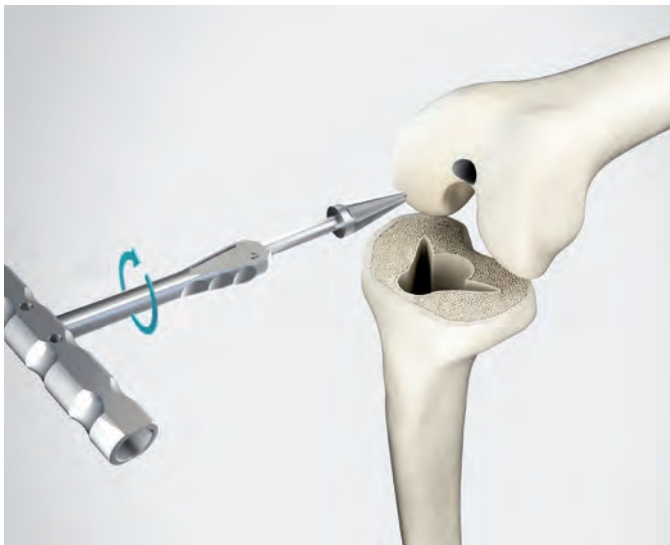
Drive in the compressor over the guide rods until the compressor touches the drill template.



18

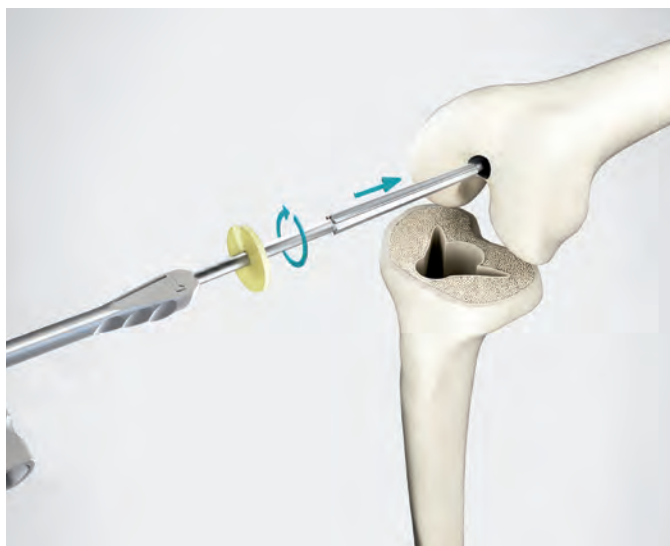
Preparation of the tibia is now complete.

Standard Preparation Femur



19

Mark the entry site with the awl (317-658/01) and open the femoral canal with the conical drill (16-3202/00).

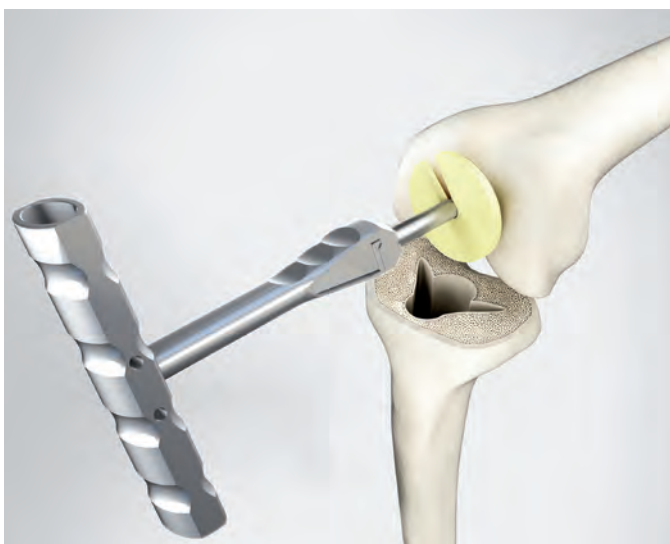


20 + 21

Mount the awl in the planned length (100 mm, 130 mm or 160 mm) in the handle (15-6053/00). The impactation plate (16-3203/00) latches into the slot on the shaft of the awl.

When uncemented modular stems are used, ream with an increasing awl diameter until the awl makes cortical contact over a continuous distance of approx. 50 mm. The uncemented implant that will be used must correspond in length and diameter to the last awl used.

For cemented modular stems, the awl should be at least 2 mm larger than the planned stem diameter.



Important notes:

The position of the impactation plate represents the level of the joint line. Using the awls with a power tool is not permitted.



22

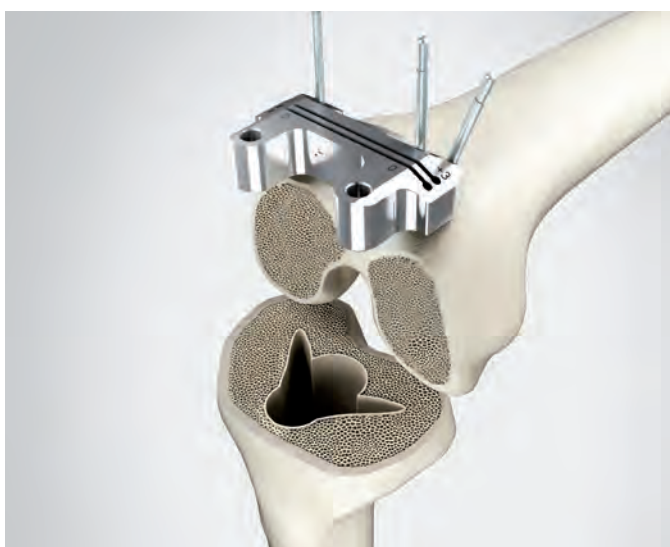
Remove the impaction plates and attach the alignment instrument for valgus angulation (16-3275/00). Ensure that the correct instrument for the right or left side is attached. The word “Left” or “Right” must face upward.



23

The appropriate saw block (16-3228/02, /03, /04) for the distal saw cut – according to the previously determined size – is fixed to the valgus alignment instrument using the clamp. The cut can be simulated with the cutting template (317-607/50).

There is a +3 mm slot for proximal offset of the cut or the instrument can be moved by +2 mm after it is fixed by wire pins.



24

After fixing the saw guide by means of two parallel and one oblique wire pins, the valgus alignment instrument and the awl are removed and the distal cut is made.

To achieve the correct resection geometry, sawblades with a thickness between 1.24 mm and 1.27 mm must be used.



25

With the alignment instrument for determination of external rotation (16-3276/00), the selected femoral size is first set and fixed with a pin.

The alignment instrument allows external rotation to be set to 0°, 3° and 5° with reference to the posterior condylar tangent. Alternatively, external rotation can also be aligned using the Whiteside line with the small dipstick in the center of the instrument. Small alignment rods can be attached medially and laterally for orientation to the epicondylar line (Insall line). Deficits in flexion and extension gap can be balanced by using femoral segments or tibial spacers.



26

Once the correct position is found, the instrument is fixed with two wire pins through the medial and lateral holes.



27

After the wire pins and alignment instrument have been removed, the dovetail adapter (317-802/36) is inserted in the depressions created by the wire pins.



28

The cutting block for chamfer cuts (16-3250/02, /03, 04) is pushed onto the side of the dovetail adapter and the central hex screw is fixed in the selected position with the hex screwdriver, wrench size 2.5 mm (10-5373/01). 2 wire pins can then be inserted for additional fixation. The anterior cut is made first, then the dorsal and finally the anterior and posterior oblique cut.

To achieve the correct resection geometry, sawblades with a thickness between 1.24 mm and 1.27 mm are to be used.



29

Before the trochlea is prepared with the chisel (317-802/32) for the patellar gliding groove, the cutting block for chamfer cuts is aligned somewhat lateral to the center. Then the chisel is connected to the handle (15-8516/45), and the trochlea is then prepared with it.



30

Following preparation of the distal femur, the last-used awl is inserted into the medullary canal again.



31

The condyle cap (16-3240/02, /03, /04) is placed on the prepared bone surfaces. The shaft of the awl forms the center.



32

A drill cap (16-3213/02, /03, /04) of the same size as the condyle cap is placed on the pegs of the condyle cap. The word "Left" or "Right" must be situated horizontally.



33

Using the centre sleeve (16-3281/00), the instruments are aligned and centered on the shaft of the awl.



34

Following alignment, the drill cap is fixed to the condyle cap with the holding clamp (16-3279/00). The condyle cap is fixed to the bone with 2 wire pins. The center sleeve and awl are removed. If necessary, the drill cap must also be removed temporarily and then re-attached again.



35

The drill for femur Ø 20 mm (16-3206/20) is inserted as far as stop.



36

After removing the holding clamp, the drill cap is removed, the saw attachment matching the selected prosthesis size (e.g. 16-3223/02) is attached and secured again with the holding clamp. The femur box is then prepared with an oscillating saw.



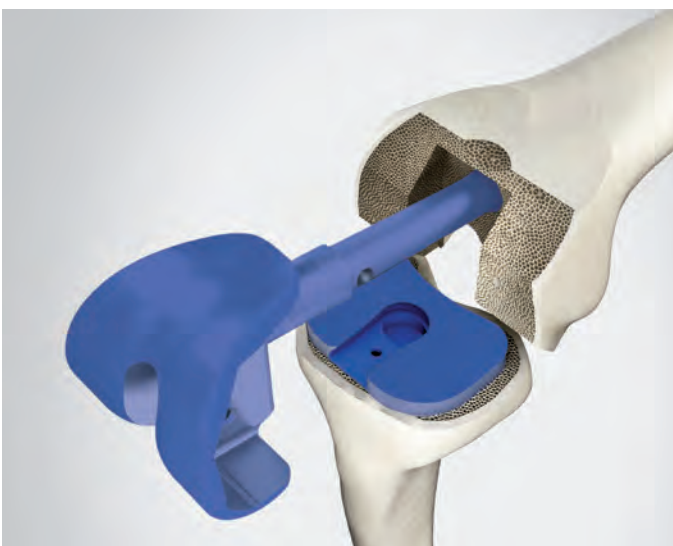
37

Preparation of distal femur is complete.



38

The tibial trial stem and trial prosthesis are joined by screwing them together and inserted into the prepared tibia.



39

The femoral trial stem and trial prosthesis are joined by screwing them together and inserted into the prepared femur.



40

Connect the two joint pieces by inserting the dorsal recess of the trial axis into the axis of the femoral component and then pushing the tibia up. Fix the screw with the hex screwdriver (64-8008/02). Test the prosthesis. Deficits in the flexion and extension gap are compensated for with femur trial segments and/or tibial trial spacers. The components are separated by proceeding in reverse order.

Note: A combination of several femoral segments or proximal tibial spacers is not allowed!



41

Remove the trial femoral and tibial components with the extraction instrument for trial prostheses (15-6061/00).

Assembling the Tibial Components



42 + 43

The tibial stems are fixed by means of a tapered connection on the tibial component. It should be ensured that the flanges of the prosthetic stems are inserted into the intended slots. The stem is then fixed to the tibial component with screw assembly. **Screws are only to be tightened hand-tight.**

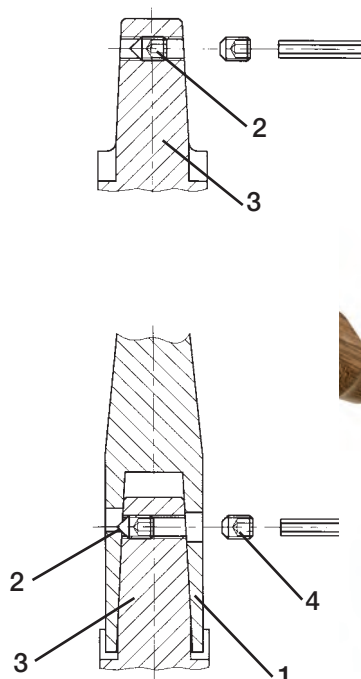
Note: When using cemented modular stems or the monobloc tibial component, the use of centering stars is mandatory. They might not be part of the sterile package for tibial components 16-2817/02, /05 und /07 when indicated.

After the underside of the tibial prosthetic component has been coated with a thin layer of bone cement, the prosthesis is inserted into the tibia with the impactor (16-0018/02).

By tightening the locking screw (2) located in the taper (3) of the tibial respectively femoral component its pointed tip presses the stem (1) firmly onto the taper. A counter screw (4) secures the stem locking screw against loosening. The screw fixation is performed medially. **Screws are only to be tightened hand-tight.**

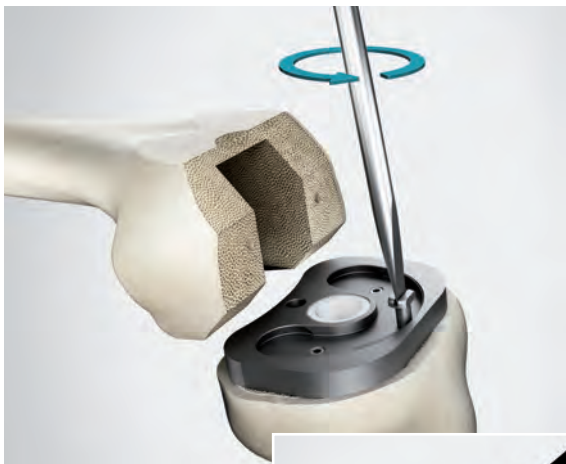
Important Information:

The locking screw (optionally pre-assembled), the counter screw and one replacement screw each are included separately in the packaging.



Counter Screw





44 + 45

Until the bone cement has set, the trial screw remains in the prosthesis thread in order to protect it. After the cement has set, the screw is removed with the screwdriver (322-145/01).

Trial Screw
(flattened screw head)



The femoral prosthetic component coated with bone cement is placed on the femur and gently tapped home with the impactor (317-646/01) until the implant fits snugly (excess bone cement is removed).

Note:

Bone cement must be used to fix femoral segments onto the intracondylar femoral components.



46

After the bone cement has set, the transport lock is removed by loosening the screw joint with the screwdriver (10-5373/01) and withdrawing the lock with slight rotation. The plateau securing screw is removed.

The PE-Plateau is connected to the inserter (15-8035/03).

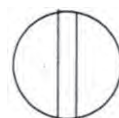


47 + 48

The connector with the rotation axle is placed on the tibial component, and the PE plateau is then inserted and screwed on.

To obtain better access to the screw, the connector is slightly rotated.

Self-locking Screw (round screw head)



Caution!

The self-locking fixation screw may only be used during the final assembly of the plateau. Loosening the fixation screw destroys the screw retention system in the polyethylene plateau, and a new plateau must then be inserted.



49

The connecting component is inserted into the intracondylar slot of the femoral component and the axle lock is removed. Move the component slightly until the axle is heard to click into the joint box.



50 + 51

Check the position of the holes.

When the prosthetic axle is fully expanded, the holes are at the level of the arrow marks. If this is not the case, the connecting and separating forceps (16-0020/01) has to be used to adjust the holes at the correct level.



52 + 53

Following expansion of the prosthetic axle, this must be locked by means of a screw. The screw (enclosed in the pack) is inserted and hand tightened in the connector using the hex screwdriver (64-1181/16).



Situation following assembly.



54 + 55

If use of a hinged knee version is planned, the screws in the tibial plateau must first be removed with the hex screwdriver (64-1181/16).



The connector with hinge axle is placed on the tibial component.

The connector is then screwed in place using the hex screwdriver (10-5373/01) and the PE plateau is inserted.



56 + 57

Connecting the components and loosening the lock are performed as in the rotation knee version.



Driving the plateau fixation screw.



58

Locking the axle by means of a screw.

Optional: Patella Preparation (Patella Resurfacing)

The following operating instructions describe the use of the LINK® Patella Component and assume the use of the instrument set available for this procedure. Both the “**resection of the patella surface**” technique and the “**reaming of the patella surface**” technique are shown.

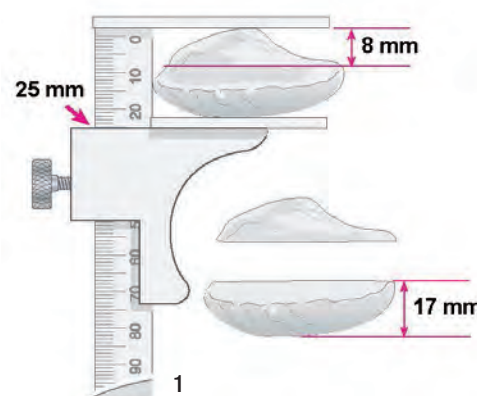
Determination of patella height/patella resection

Using the calliper (65-2000) allows the height of the patella to be determined (1). The dimension is established and an amount corresponding to the size of the selected implant subtracted. The remainder equals the target dimension following resection. A minimal residual dimension of 12 mm should be maintained.

Example:

For a patella size 2 (patella diameter 28 mm), the following calculation is performed:

1. patella 25 mm thick,
2. resection 8 mm,
3. 17 mm remaining patella bone.



Patella Size	Patella-Ø	Patella Height
1	25 mm	7 mm
2	28 mm	8 mm
3	31 mm	9 mm
4	34 mm	10 mm

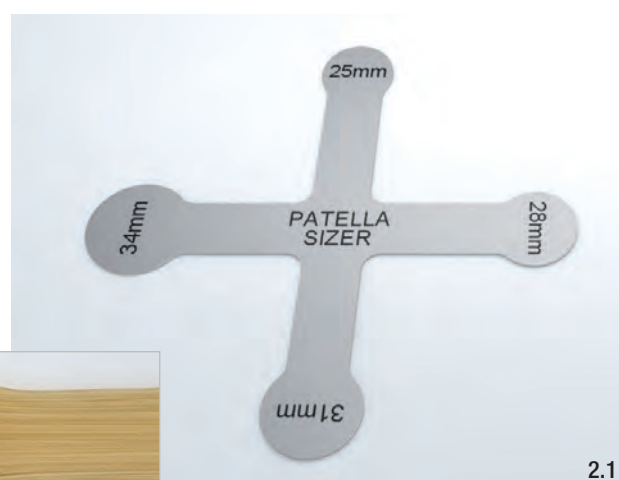
Preparation for Patella Component

The patella surface can be prepared using the onlay technique by resecting with an oscillating saw or by reaming.

Sizing

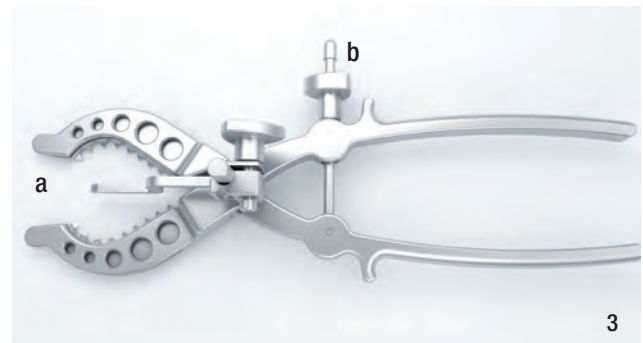
A patella sizing template (340-010) corresponding to the implants is available (2.1)

The template is selected that most adequately covers the articular surface without any overhang. If bone is deficient on the lateral side, the next smaller size is selected, but positioned slightly to the medial side to enhance patellar tracking (2.2).

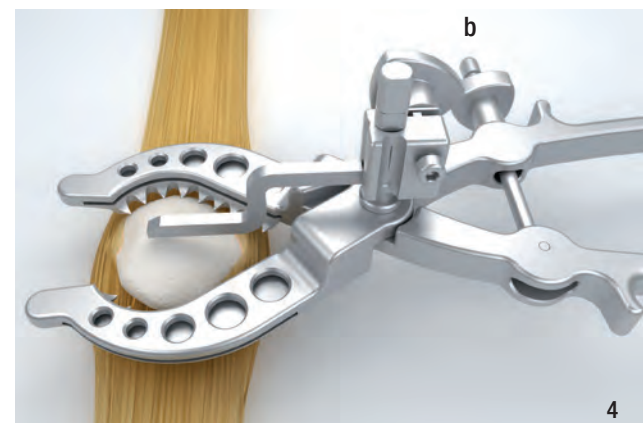


Option: Resection

The height of the bone to be resected can be adjusted using the height stylus (a) on the resection clamp (340-006) (3).



In doing this, it is important to ensure that the remaining patella is sufficiently thick. The patella is held using the toothed jaws. The sectional plane must lie parallel to the extended patellar tendon and the height calliper must lie on the bone. In order to clamp the patella firmly, the resection clamp (340-006) is compressed firmly and fixed using the lateral setting screw (b) (4).



The resection is carried out using an oscillating saw with a 1.27 mm thick sawblade. The saw is guided using the cutting slots of the resection clamp (340-006) (5).

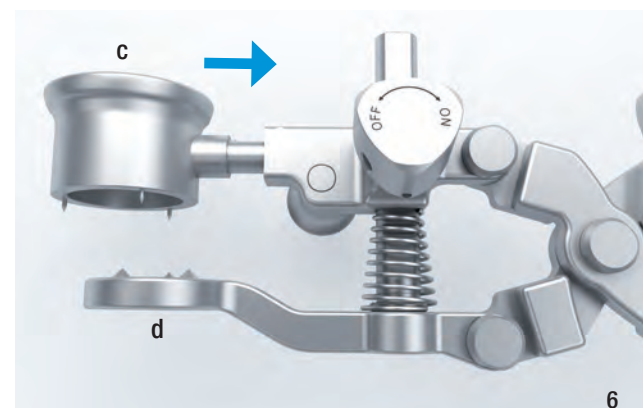


Option: Reaming

Reaming Preparation

The reaming guide (340-025 to 340-034) (c) of the appropriate size is inserted into the patella holding clamp (340-005) (d) (6).

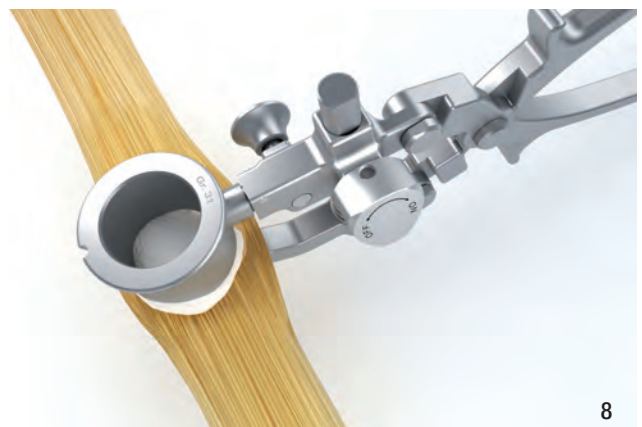
Note: Press the button on the side to insert and remove the guide.



A ratchet can be activated on the patella holding clamp (340-005) with the retaining screw (e). By turning the screw to “ON” the clamp is held shut. By turning the screw to “OFF”, the ratchet is released (7).



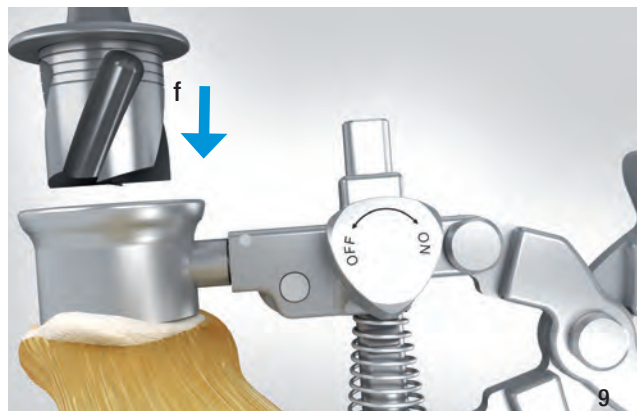
Following removal of the peripheral osteophytes, the patella is clamped with the patella holding clamp (340-005), aiming at the most central position possible (8). Good fixation is achieved when the spikes on the guide grip into the bone.



Reaming

The patella reamer is selected (f) to match the implant size which was previously determined and the corresponding reaming guide (9).

The patella reamer (340-125B to 340-134B) has a Hudson drill B connection for direct fixation with corresponding power tool connection. It is compatible with other machine systems by using an adapter for snap lock chuck.



The patella surface is prepared using the patella reamer (f). Reaming depth is verified by means of the mechanical stop on the reamer. In addition, there are marking grooves at 2 mm intervals. The maximum reaming depth is reached when the reamer collar makes contact with the reaming guide (10).



Fixation Hole Drilling

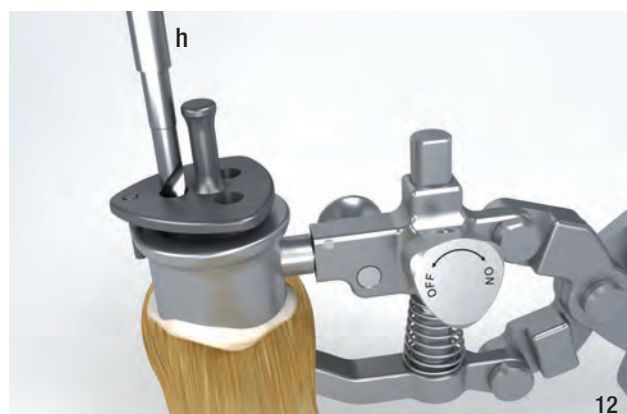
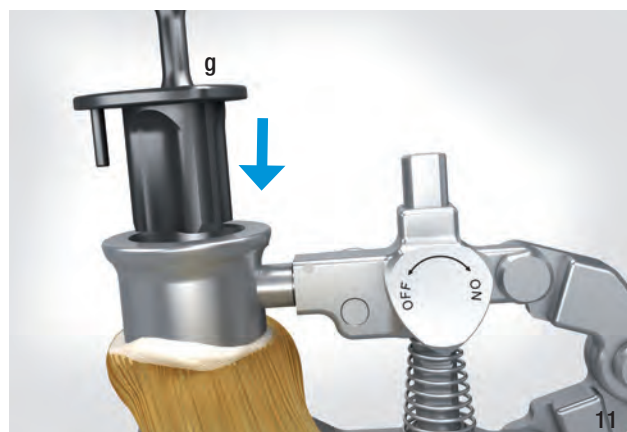
If the saw blade resection option is selected, the patella is to be fixed with the patella holding clamp (340-005) as follows:

- Insert the patella reaming guide into the patella holding clamp
- Ratchet function “ON”
- Clamp the patella firmly

Following description is for options „resection“ as well as „reaming“.

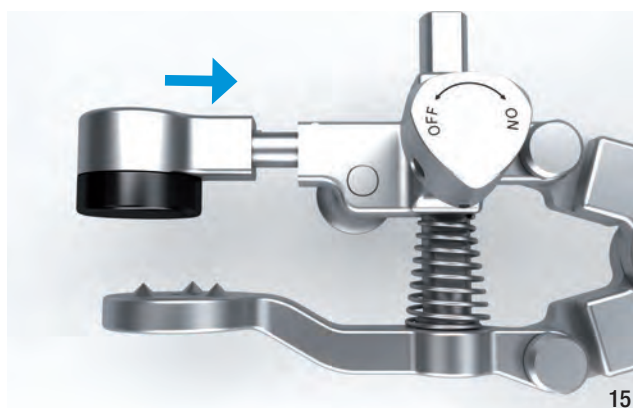
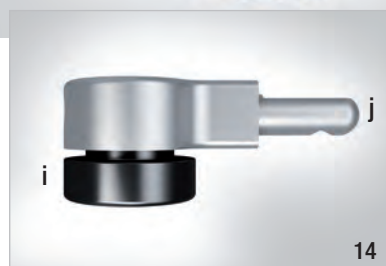
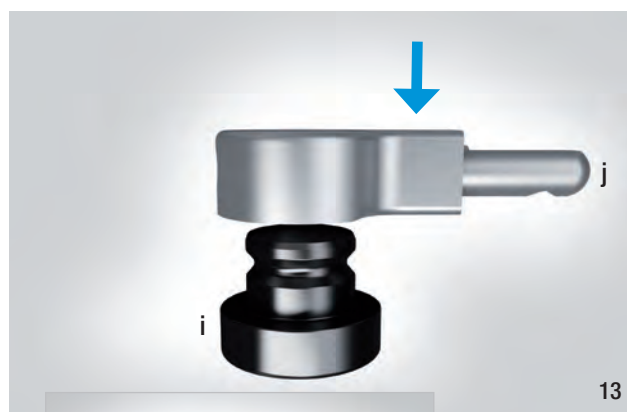
Insert the patella drill guide (340-225 to 340-234) (g) for the anchoring three holes which corresponds to the chosen implant size into the reaming guide (340-025 to 340-034) (11).

In doing so, ensure that the guide pin on the drill guide lies in the opening made for it in the reaming guide. Using the Ø 6.3 mm patella drill (340-306B) (h) three holes guided by the drill guide are drilled. The patella drill is inserted by the until stop (12).



Patella Implantation

The reaming guide is removed from the patella holding clamp (340-005). The patella pusher inserter (340-009) (i) is pressed into the patella pusher attachment (340-007) (j) (13+14) and the pusher inserter is then inserted into the holding clamp (340-005) (15). Appropriate patella tracking can be assessed using patella trial prostheses (340-325 to 340-334).



Following extensive rinsing and removal of all impeding soft tissue, the bone cement is applied to the back of the implant, and the patella component is placed by hand and pressed on using the patella holding clamp (340-005) with the patella pusher inserter (340-009) (**16**).

Caution!

Prepare the bone cement according to the manufacturer's instructions.

Ensure that excess bone cement is completely removed and that no loose particles of bone cement remain in the joint.



Joint Components Endo-Model® SL® for Rotating and Non-Rotating Hinge Knee



Femoral Components, modular, intracondylar
Material: CoCrMo, UHMWPE

Item no.	Size	Side	M/L - A/P
16-2821/21	small (S)	right	63.0 mm - 57.0 mm
16-2821/22	small (S)	left	63.0 mm - 57.0 mm
16-2823/21	medium (M)	right	69.0 mm - 62.0 mm
16-2823/22	medium (M)	left	69.0 mm - 62.0 mm
16-2825/21	large (L)	right	72.5 mm - 65.0 mm
16-2825/22	large (L)	left	72.5 mm - 65.0 mm

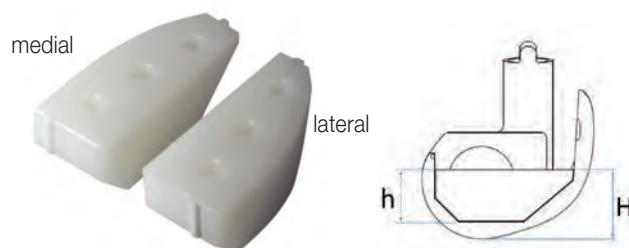


Distal Femoral Replacement Components, modular

Material: CoCrMo, UHMWPE

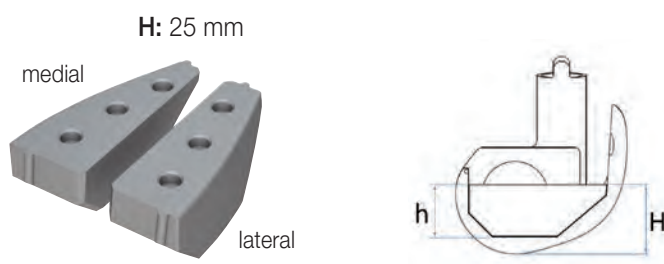
Item no.	Size	Side	M/L - A/P
16-2853/21	small (S)	right	60.0 mm - 57.0 mm
16-2853/22	small (S)	left	60.0 mm - 57.0 mm
16-2855/21	medium (M)	right	65.0 mm - 62.0 mm
16-2855/22	medium (M)	left	65.0 mm - 62.0 mm
16-2857/21	large (L)	right	75.0 mm - 65.0 mm
16-2857/22	large (L)	left	75.0 mm - 65.0 mm

Joint Components Endo-Model® SL® for Rotating and Non-Rotating Hinge Knee



Femoral Segments, cemented,
for femoral components, to replace distal condyles
Material: UHMWPE

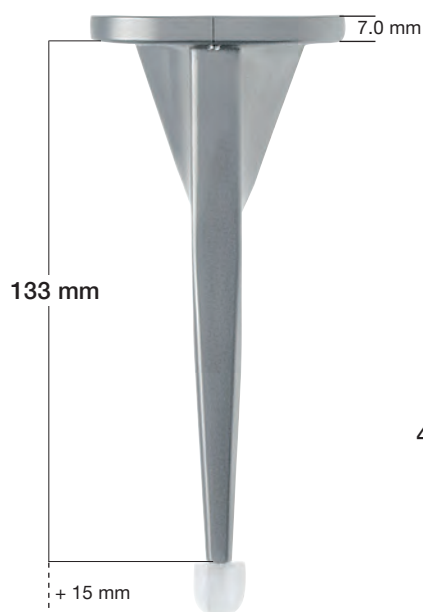
Item no.	Size	Side		h	H
15-8519/01	small	right	medial	18 mm	25 mm
15-8519/02	medium	right	medial	19 mm	25 mm
15-8519/03	large	right	medial	17 mm	25 mm
15-8519/11	small	right	lateral	18 mm	25 mm
15-8519/12	medium	right	lateral	19 mm	25 mm
15-8519/13	large	right	lateral	17 mm	25 mm
15-8520/01	small	left	medial	18 mm	25 mm
15-8520/02	medium	left	medial	19 mm	25 mm
15-8520/03	large	left	medial	17 mm	25 mm
15-8520/11	small	left	lateral	18 mm	25 mm
15-8520/12	medium	left	lateral	19 mm	25 mm
15-8520/13	large	left	lateral	17 mm	25 mm



Femoral Segments, cementless (bone side),
for femoral components, to replace distal condyles,
Material: Tilastan®

Item no.	h	H	Size	Side	
15-8517/01	18 mm	25 mm	small	right	medial
15-8517/02	19 mm	25 mm	medium	right	medial
15-8517/03	17 mm	25 mm	large	right	medial
15-8517/11	18 mm	25 mm	small	right	lateral
15-8517/12	19 mm	25 mm	medium	right	lateral
15-8517/13	17 mm	25 mm	large	right	lateral
15-8518/01	18 mm	25 mm	small	left	medial
15-8518/02	19 mm	25 mm	medium	left	medial
15-8518/03	17 mm	25 mm	large	left	medial
15-8518/11	18 mm	25 mm	small	left	lateral
15-8518/12	19 mm	25 mm	medium	left	lateral
15-8518/13	17 mm	25 mm	large	left	lateral

Joint Components Endo-Model® SL® for Rotating and Non-Rotating Hinge Knee



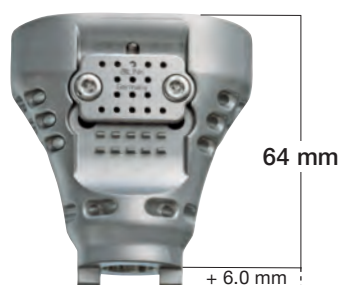
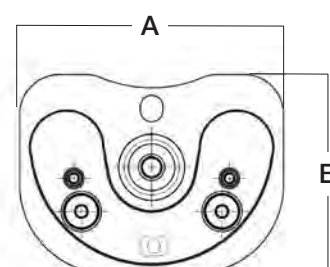
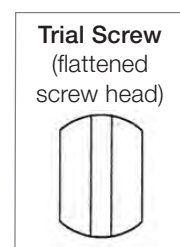
Tibial Components, neutral
Material: CoCrMo, UHMWPE

Item no.	Size	A x B mm
16-2817/02	small (S)	60 x 45
16-2817/05	medium (M)	65 x 45
16-2817/07	large (L)	75 x 48



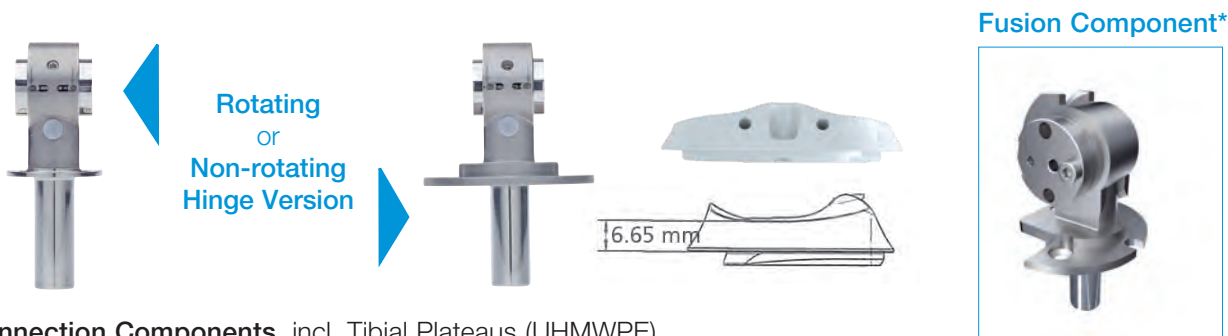
Tibial Components System SL®,
modular, neutral
Material: CoCrMo, UHMWPE

Item no.	Size	A x B mm
16-2817/32	small (S)	60 x 45
16-2817/35	medium (M)	65 x 45
16-2817/37	large (L)	75 x 48



Proximal Tibia Replacement Components, modular, neutral
Material: Tilastan®, UHMWPE

Item no.	Size	A x B mm
16-2849/22	small (S)	60 x 45
16-2849/25	medium (M)	65 x 45
16-2849/27	large (L)	75 x 48



Connection Components, incl. Tibial Plateaus (UHMWPE)

Item no.	Rotating Hinge Material: CoCrMo, UHMWPE		Item no.	Non-rotating Hinge Material: CoCrMo, UHMWPE		Arthrodesis Fusion Version* Material: CoCrMo, UHMWPE
16-2840/02	small (S)	neutral	16-2841/02	small (S)	neutral	Knee fusion option for in-situ femoral and tibial components.
16-2840/05	medium (M)	neutral	16-2841/05	medium (M)	neutral	
16-2840/07	large (L)	neutral	16-2841/07	large (L)	neutral	

* only available as custom-made implant

Proximal Tibial Spacers, Tilastan® and UHMWPE for Endo-Model® SL® Rotating and Non-Rotating Hinge Knee



Proximal Tibia Spacers, full,
for right and left, incl. 2 countersunk screws,
wrench size 2.5 mm, Material: Tilastan®

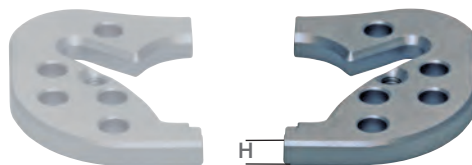
Item no.	Size	H Height	Width
16-2910/05	small	5 mm	60 mm
16-2910/10	small	10 mm	60 mm
16-2910/15	small	15 mm	60 mm
16-2920/05	medium	5 mm	65 mm
16-2920/10	medium	10 mm	65 mm
16-2920/15	medium	15 mm	65 mm
16-2930/05	large	5 mm	75 mm
16-2930/10	large	10 mm	75 mm
16-2930/15	large	15 mm	75 mm

Proximal Tibia Spacers, full,
for right and left, Material: UHMWPE

Item no.	Size	H Height	Width
16-3000/05	small	5 mm	60 mm
16-3000/10	small	10 mm	60 mm
16-3000/15	small	15 mm	60 mm
16-3010/05	medium	5 mm	65 mm
16-3010/10	medium	10 mm	65 mm
16-3010/15	medium	15 mm	65 mm
16-3020/05	large	5 mm	75 mm
16-3020/10	large	10 mm	75 mm
16-3020/15	large	15 mm	75 mm

Important note:

Proximal tibial spacers may not be combined!


Proximal Tibial Spacers, half,

lateral and medial usable, incl. 1 countersunk screw, wrench size 2.5 mm, Material: Tilastan[®]

Item no.	Size	H Height
16-2950/05	small	5 mm
16-2950/10	small	10 mm
16-2950/15	small	15 mm
16-2960/05	medium	5 mm
16-2960/10	medium	10 mm
16-2960/15	medium	15 mm
16-2970/05	large	5 mm
16-2970/10	large	10 mm
16-2970/15	large	15 mm

Important note:

Proximal tibial spacers may not be combined with each other!

Modular Stems, with male taper

Modular Stems* with male taper (for 6 mm flanges)

Material: Tilastan®



Cementless					
Item no.	Stem length L 100 mm	Item no.	Stem length L 130 mm	Item no.	Stem length L 160 mm
15-8524/50	Ø 12/9 mm	15-8523/50	Ø 12/8 mm	15-8522/50	Ø 12/7 mm
15-8524/51	Ø 13/10 mm	15-8523/51	Ø 13/9 mm	15-8522/51	Ø 13/8 mm
15-8524/52	Ø 14/11mm	15-8523/52	Ø 14/10 mm	15-8522/52	Ø 14/9 mm
15-8524/53	Ø 15/12mm	15-8523/53	Ø 15/11 mm	15-8522/53	Ø 15/10 mm
15-8524/54	Ø 16/13mm	15-8523/54	Ø 16/12 mm	15-8522/54	Ø 16/11 mm
15-8524/55	Ø 17/14mm	15-8523/55	Ø 17/13 mm	15-8522/55	Ø 17/12 mm
15-8524/56	Ø 18/15 mm	15-8523/56	Ø 18/14 mm	15-8522/56	Ø 18/13 mm
15-8524/57	Ø 19/16 mm	15-8523/57	Ø 19/15 mm	15-8522/57	Ø 19/14 mm
15-8524/58	Ø 20/17 mm	15-8523/58	Ø 20/16 mm	15-8522/58	Ø 20/15 mm
15-8524/59	Ø 21/18 mm	15-8523/59	Ø 21/17 mm	15-8522/59	Ø 21/16 mm
15-8524/60	Ø 22/19 mm	15-8523/60	Ø 22/18 mm	15-8522/60	Ø 22/17 mm
15-8524/61	Ø 23/20 mm	15-8523/61	Ø 23/19 mm	15-8522/61	Ø 23/18 mm
15-8524/62	Ø 24/21 mm	15-8523/62	Ø 24/20 mm	15-8522/62	Ø 24/19 mm

Modular Stems* with male taper (for 6 mm flanges)

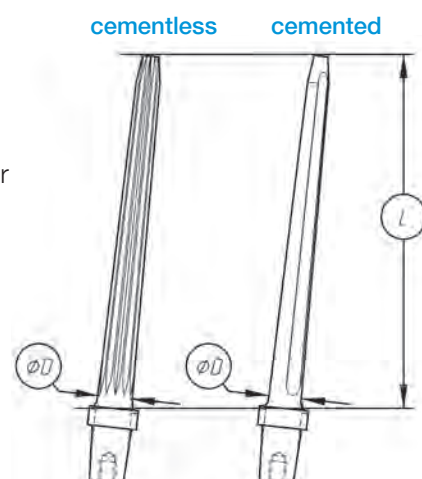
Material: CoCrMo



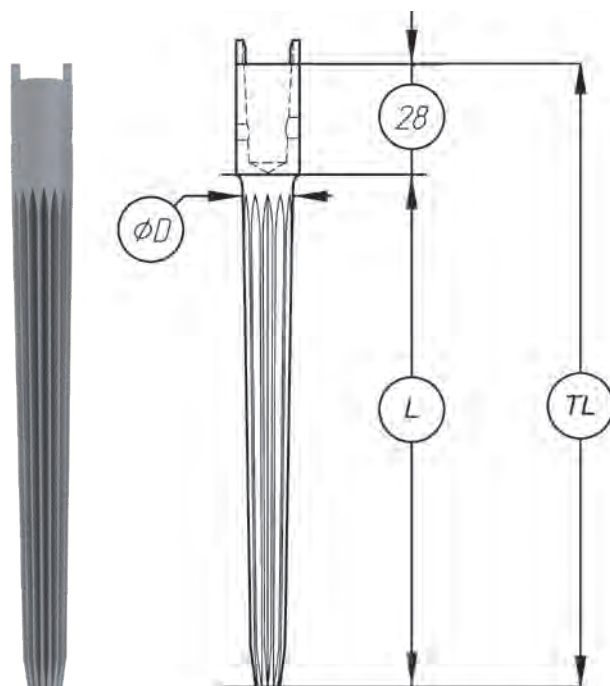
Cemented					
Item no.	Stem length L 100 mm	Item no.	Stem length L 130 mm	Item no.	Stem length L 160 mm
15-8524/40	Ø 12/9 mm	15-8523/40	Ø 12/8 mm	15-8522/40	Ø 12/7 mm
15-8524/42	Ø 14/11 mm	15-8523/42	Ø 14/10 mm	15-8522/42	Ø 14/9 mm
15-8524/44	Ø 16/13 mm	15-8523/44	Ø 16/12 mm	15-8522/44	Ø 16/11 mm

* These modular stems can be combined with:

- coupling device to assemble a proximal partial femur replacement
- femoral joint component, solid, or a stem segment attached to it to assemble a distal partial femur replacement
- coupling device for the diaphyseal spacer
- proximal tibial replacement



Modular Stems, with female taper



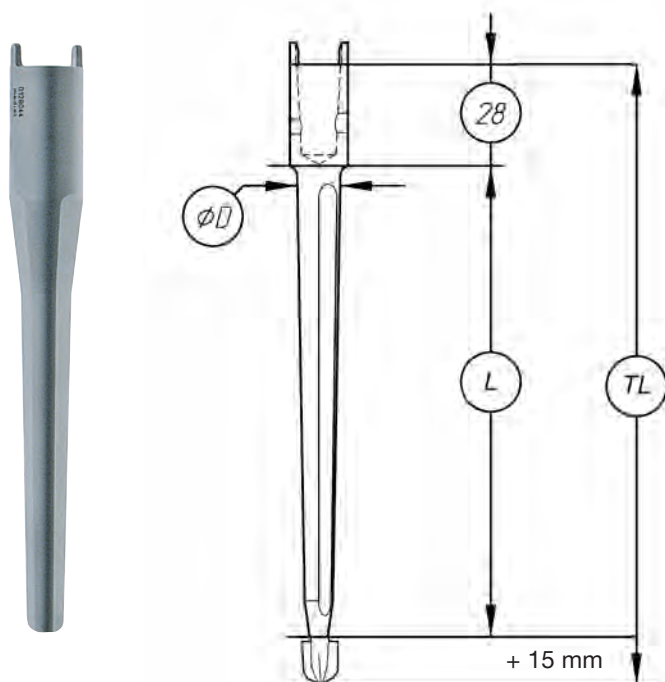
Modular Stems* with female taper (with 6 mm flanges)

Material: Tilastan®

Cementless					
Item no.	Stem length L 100 mm	Total length TL 128 mm	Item no.	Stem length L 130 mm	Total length TL 158 mm
15-8517/50	Ø 12/9 mm		15-8516/50	Ø 12/8 mm	
15-8517/51	Ø 13/10 mm		15-8516/51	Ø 13/9 mm	
15-8517/52	Ø 14/11 mm		15-8516/52	Ø 14/10 mm	
15-8517/53	Ø 15/12 mm		15-8516/53	Ø 15/11 mm	
15-8517/54	Ø 16/13 mm		15-8516/54	Ø 16/12 mm	
15-8517/55	Ø 17/14 mm		15-8516/55	Ø 17/13 mm	
15-8517/56	Ø 18/15 mm		15-8516/56	Ø 18/14 mm	

Item no.	Stem length L 160 mm	Total length TL 188 mm
15-8515/50	Ø 12/7 mm	
15-8515/51	Ø 13/8 mm	
15-8515/52	Ø 14/9 mm	
15-8515/53	Ø 15/10 mm	
15-8515/54	Ø 16/11 mm	
15-8515/55	Ø 17/12 mm	
15-8515/56	Ø 18/13 mm	

Modular Stems, with female taper



Modular Stems*

with female taper (with 6 mm flanges), Material: CoCrMo

Cemented					
Item no.	Stem length L 100 mm	Total length TL 128 mm	Item no.	Stem length L 130 mm	Total length TL 158 mm
15-8527/40	Ø 12/9 mm		15-8526/40	Ø 12/8 mm	
15-8527/42	Ø 14/11 mm		15-8526/42	Ø 14/10 mm	
15-8527/44	Ø 16/13 mm		15-8526/44	Ø 16/12 mm	

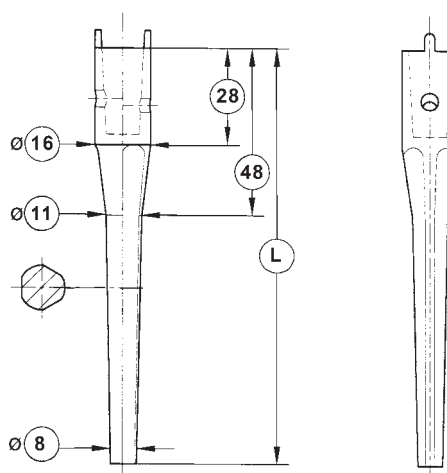
Item no.	Stem length L 160 mm	Total length TL 188 mm
15-8525/40	Ø 12/7 mm	
15-8525/42	Ø 14/9 mm	
15-8525/44	Ø 16/11 mm	

* These modular stems can be combined with: tibial components, neutral, modular (with 6 mm noses)

Modular Stems, short

Material: CoCrMo

Cemented	
Item no.	Stem length L mm
15-2950/01	50
15-2950/02	80



Centering Stars

Centering Stars, height 15 mm
Material: UHMWPE

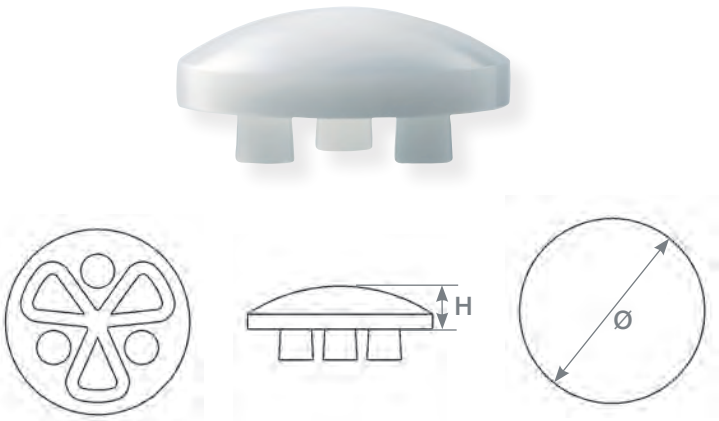
Item no.	Item no.	Ø mm
Set	consisting of:	
15-2975/01	15-2975/12	12
	15-2975/14	14
	15-2975/16	16



LINK® Patella Components
for Endo-Model® SL® Rotating and Non-Rotating Hinge Knee





LINK® Patella Components
Material: UHMWPE




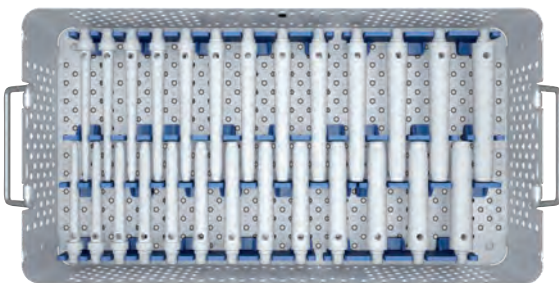
Item no.	Size	Ø mm	Height mm
318-401/25	1	25	7
318-401/28	2	28	8
318-401/31	3	31	9
318-401/34	4	34	10

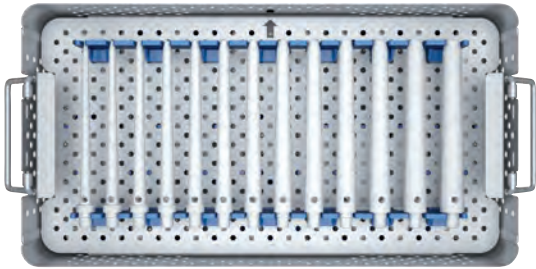
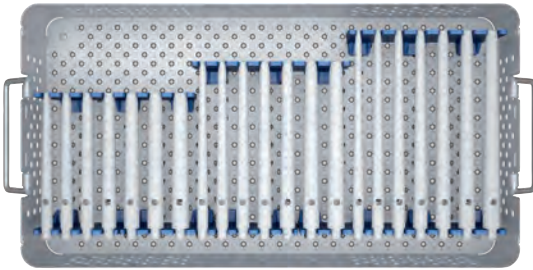
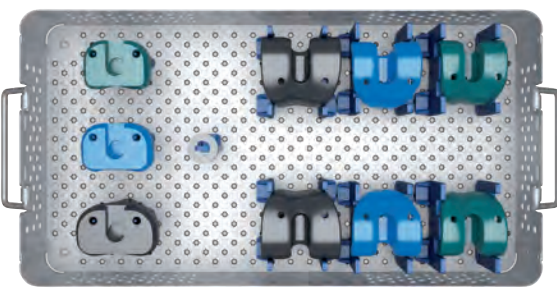
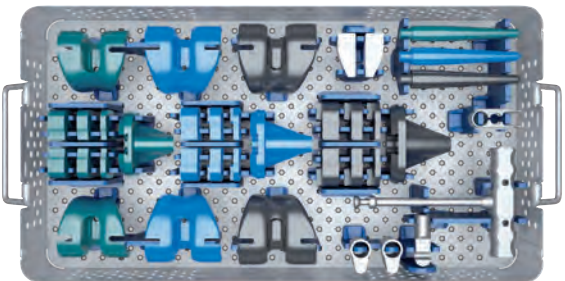


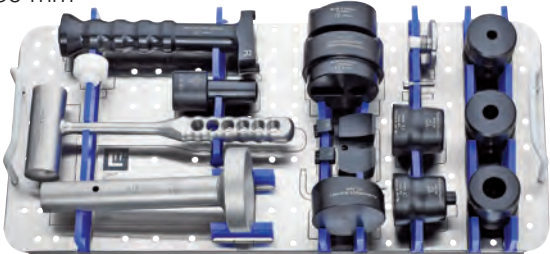
Instrument Set – Overview Containers

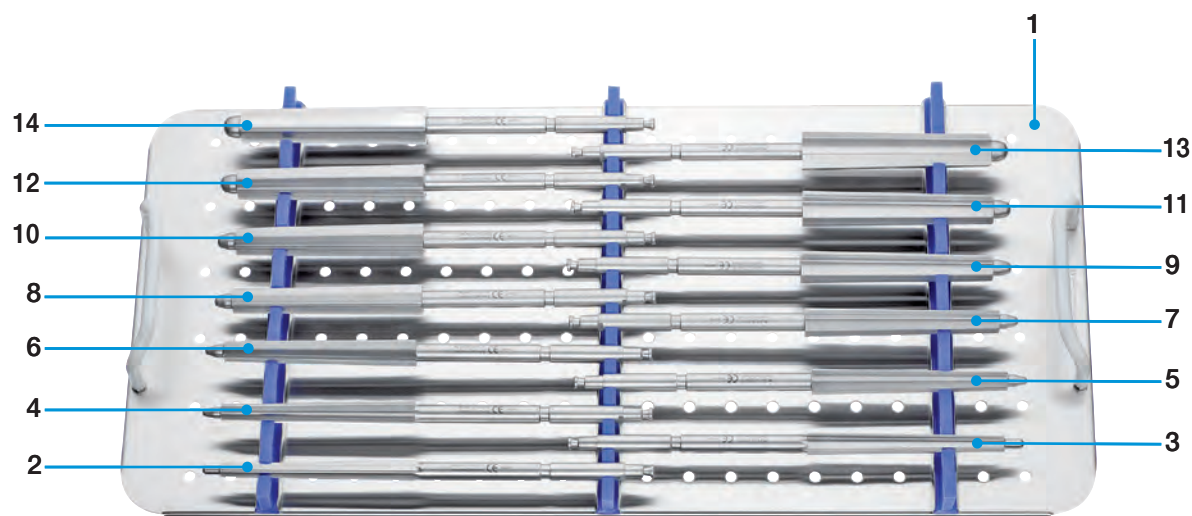
for Endo-Model® SL® Rotating and Non-Rotating Hinge Knee

15-8710/01	Container A	Tapered Reamers 100 mm
consisting of: 05-2001/03 15-8711/01	N11 Standard Container , 575 x 275 x 100 mm Tray , empty, 550 x 265 x 50 mm Tray with detailed instruments see page 48	
15-8720/01	Container B	Tapered Reamers 130 mm
consisting of: 05-2001/03 15-8721/01	N11 Standard Container , 575 x 275 x 100 mm Tray , empty, 550 x 265 x 50 mm Tray with detailed instruments see page 49	
15-8730/01	Container C	Tapered Reamers 160 mm
consisting of: 05-2001/03 15-8731/01	N11 Standard Container , 575 x 275 x 100 mm Tray , empty, 550 x 265 x 50 mm Tray with detailed instruments see page 50	
15-8740/01	Container D	General Instruments
consisting of: 05-2001/03 15-8741/01	N11 Standard Container , 575 x 275 x 100 mm Tray , empty, 550 x 265 x 50 mm Tray with detailed instruments see page 51	

15-8760/01	Container F	Femur I
consisting of: 05-2001/03 15-8761/01	<p>N11 Standard Container, 575 x 275 x 100 mm</p> <p>Tray, empty, 550 x 265 x 50 mm</p> <p>Tray with detailed instruments see page 52</p>	
15-8770/01	Container G	Femur II
consisting of: 15-8771/01	<p>N21 Standard Container, 575 x 275 x 130 mm</p> <p>Tray, empty, 550 x 265 x 50 mm</p> <p>Tray with detailed instruments see page 53</p>	
15-8780/01	Container H	Tibia
consisting of: 05-2002/03 15-8781/01	<p>N21 Standard Container, 575 x 275 x 130 mm</p> <p>Tray, empty, 550 x 265 x 50 mm</p> <p>Tray with detailed instruments see page 54</p>	
15-8790/02	Instrument Set	Trial Stems Femur, 100 + 130 mm
consisting of: 15-8791/02	<p>Instrument Tray, empty, 478 x 253 x 106 mm</p> <p>Tray with detailed instruments see page 55</p>	

15-8790/02	Instrument Set	Trial Stems Femur, 160 mm
consisting of: 15-8791/02	Instrument Tray , empty, 478 x 253 x 106 mm Tray with detailed instruments see page 56	
15-8810/02	Instrument Set	Tibia Trial Stems: 100, 130 und 160 mm
consisting of: 15-8811/02	Instrument Tray , empty, 478 x 253 x 76 mm Tray with detailed instruments see page 57	
15-8840/02	Instrument Set	Trials Distal Femur and Proximal Tibia Replacement
consisting of: 15-8841/02	Instrument Tray , empty, 478 x 253 x 76 mm Tray with detailed instruments see page 58	
15-8820/02	Instrument Set	Femur/Tibia, intracondylar
consisting of: 15-8821/02	Instrument Tray , empty, 478 x 253 x 76 mm Tray with detailed instruments see page 59	

16-0100/00	Container	Assembling Instrument Set
<p>consisting of:</p> <p>05-2002/03</p> <p>16-0100/01</p>	<p>N21 Standard Container, 575 x 275 x 130 mm</p> <p>Tray, empty, 550 x 265 x 50 mm</p> <p>Tray with detailed instruments see page 60</p>	

15-8710/01 Container A – Tapered Reamers 100 mm


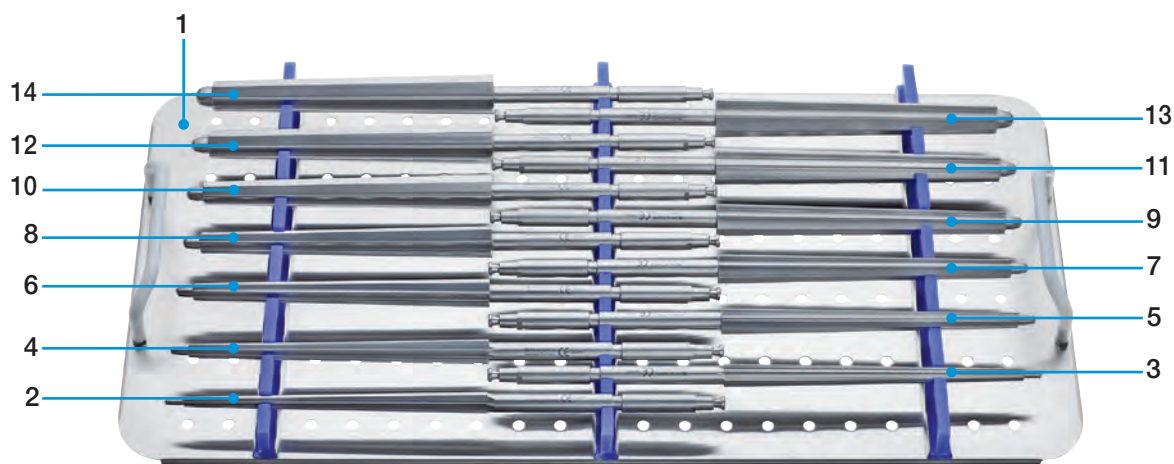
1	05-2001/03 15-8711/01	N11 Standard Container , 575 x 275 x 100 mm (without illustration) Tray , empty, 550 x 265 x 50 mm	
		Tapered Reamers: for prosthesis stems 100 mm, conical, with fitting B: Hudson*	
2	16-5100/12	for: stem-Ø 12 mm	stem length 100 mm
3	16-5100/13	for: stem-Ø 13 mm	stem length 100 mm
4	16-5100/14	for: stem-Ø 14 mm	stem length 100 mm
5	16-5100/15	for: stem-Ø 15 mm	stem length 100 mm
6	16-5100/16	for: stem-Ø 16 mm	stem length 100 mm
7	16-5100/17	for: stem-Ø 17 mm	stem length 100 mm
8	16-5100/18	for: stem-Ø 18 mm	stem length 100 mm
9	16-5100/19	for: stem-Ø 19 mm	stem length 100 mm
10	16-5100/20	for: stem-Ø 20 mm	stem length 100 mm
11	16-5100/21	for: stem-Ø 21 mm	stem length 100 mm
12	16-5100/22	for: stem-Ø 22 mm	stem length 100 mm
13	16-5100/23	for: stem-Ø 23 mm	stem length 100 mm
14	16-5100/24	for: stem-Ø 24 mm	stem length 100 mm

* optional various adapters with different fittings are available (see page 69).

15-8720/01 Container B – Tapered Reamers 130 mm

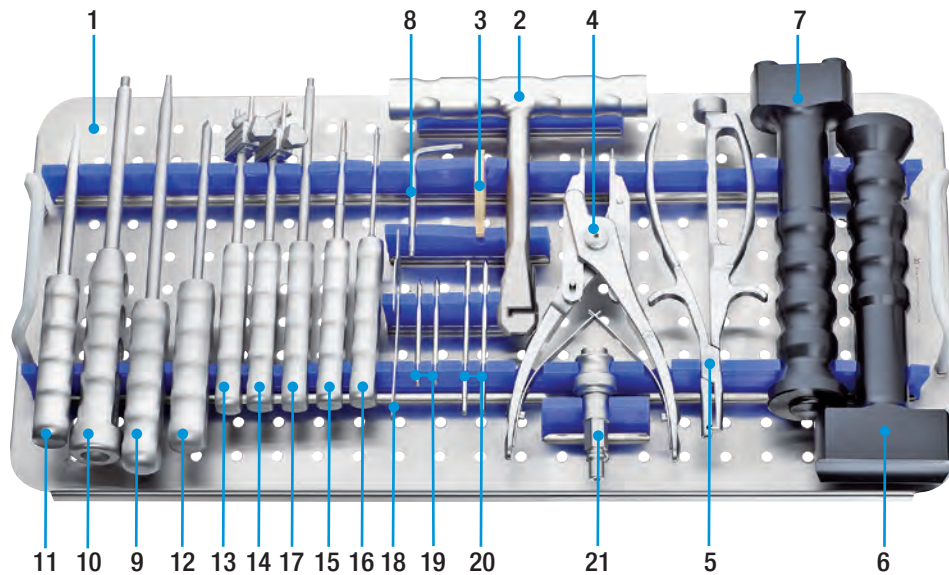

1	05-2001/03 15-8721/01	N11 Standard Container , 575 x 275 x 100 mm (without illustration) Tray , empty, 550 x 265 x 50 mm	
		Tapered Reamers: for prosthesis stems 130 mm, conical, with fitting B: Hudson*	
2	16-5130/12	for: stem-Ø 12 mm	stem length 130 mm
3	16-5130/13	for: stem-Ø 13 mm	stem length 130 mm
4	16-5130/14	for: stem-Ø 14 mm	stem length 130 mm
5	16-5130/15	for: stem-Ø 15 mm	stem length 130 mm
6	16-5130/16	for: stem-Ø 16 mm	stem length 130 mm
7	16-5130/17	for: stem-Ø 17 mm	stem length 130 mm
8	16-5130/18	for: stem-Ø 18 mm	stem length 130 mm
9	16-5130/19	for: stem-Ø 19 mm	stem length 130 mm
10	16-5130/20	for: stem-Ø 20 mm	stem length 130 mm
11	16-5130/21	for: stem-Ø 21 mm	stem length 130 mm
12	16-5130/22	for: stem-Ø 22 mm	stem length 130 mm
13	16-5130/23	for: stem-Ø 23 mm	stem length 130 mm
14	16-5130/24	for: stem-Ø 24 mm	stem length 130 mm

* optional various adapters with different fittings are available (see page 69).

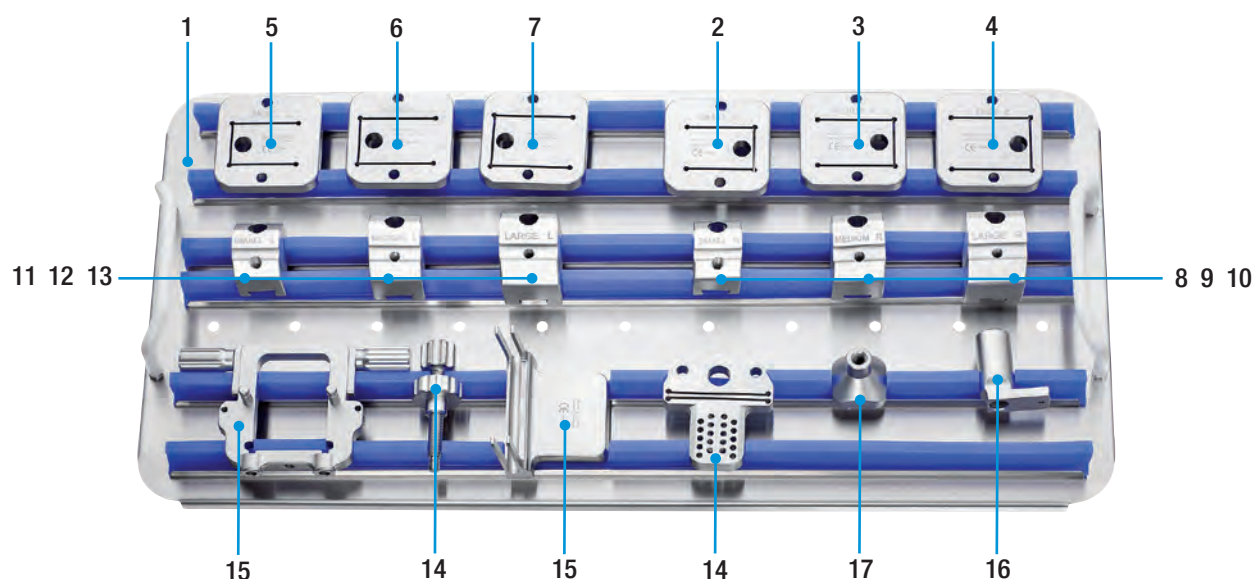
15-8730/01 Container C – Tapered Reamers 160 mm


1	05-2001/03 15-8731/01	N11 Standard Container , 575 x 275 x 100 mm (without illustration) Tray , empty, 550 x 265 x 50 mm	
		Tapered Reamers: for prosthesis stems 160 mm, conical, with fitting B: Hudson*	
2	16-5160/12	for: stem-Ø 12 mm	stem length 160 mm
3	16-5160/13	for: stem-Ø 13 mm	stem length 160 mm
4	16-5160/14	for: stem-Ø 14 mm	stem length 160 mm
5	16-5160/15	for: stem-Ø 15 mm	stem length 160 mm
6	16-5160/16	for: stem-Ø 16 mm	stem length 160 mm
7	16-5160/17	for: stem-Ø 17 mm	stem length 160 mm
8	16-5160/18	for: stem-Ø 18 mm	stem length 160 mm
9	16-5160/19	for: stem-Ø 19 mm	stem length 160 mm
10	16-5160/20	for: stem-Ø 20 mm	stem length 160 mm
11	16-5160/21	for: stem-Ø 21 mm	stem length 160 mm
12	16-5160/22	for: stem-Ø 22 mm	stem length 160 mm
13	16-5160/23	for: stem-Ø 23 mm	stem length 160 mm
14	16-5160/24	for: stem-Ø 24 mm	stem length 160 mm

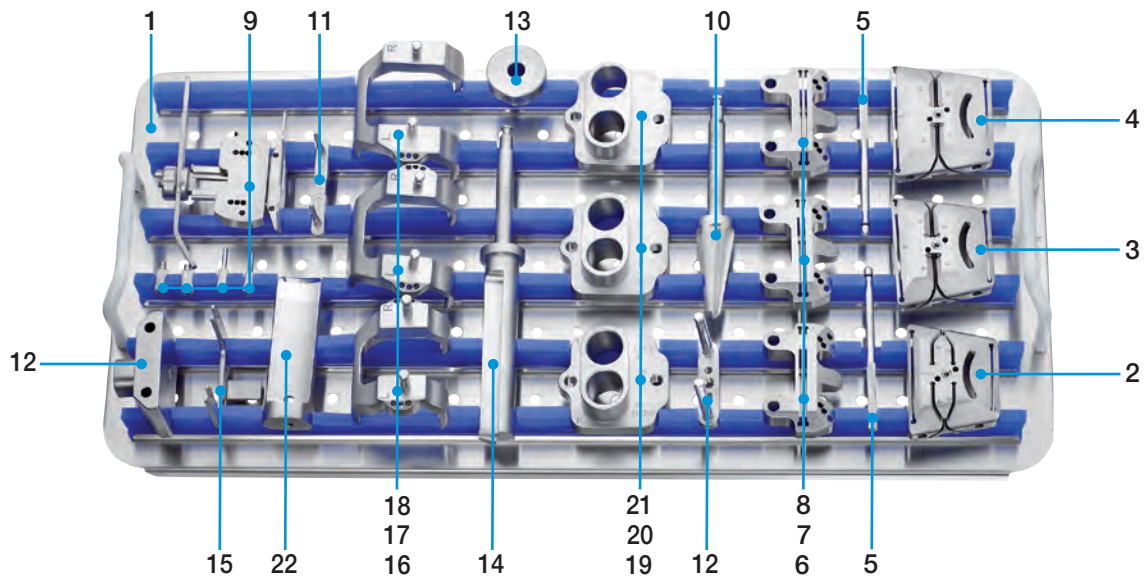
* optional various adapters with different fittings are available (see page 69).

15-8740/01 Container D – General Instruments


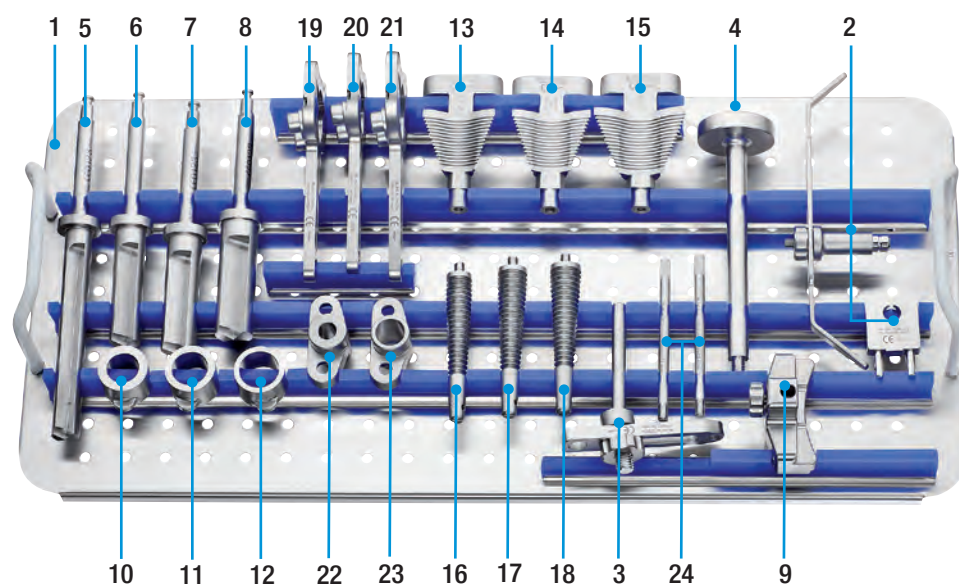
1	05-2001/03 15-8741/01	N11 Standard Container , 575 x 275 x 100 mm (without illustration) Tray , empty, 550 x 265 x 50 mm
2	15-6053/00	T-Handle , with Hudson fitting
3	16-3203/00	Impaction Plate for tapered reamers
4	16-0020/01	Connecting/Disconnecting Forceps , 175 mm
5	317-586	Extraction Forceps for fixation pins, 210 mm
6	317-646/01	Grooved Driver for femoral components, 210 mm
7	16-0018/02	Grooved Driver für tibial components SL®
8	16-0116/01	Hex Screwdriver , wrench size 2.5 mm
9	64-8008/02	Hex Screwdriver , wrench size 3.5 mm, 250 mm
10	15-8516/45	Driver Extractor , for modular stems, 365 mm
11	322-145/01	Screwdriver , blade width 8 mm, 210 mm
12	317-658/01	Bone Awl , with trocar point, 215 mm
13	15-8035/02	Insertion Instrument for PE plateaus Endo-Model® S, M and L
14	15-8035/03	Insertion Instrument for PE plateaus Endo-Model® XS and Endo-Model® SL®
15	10-5373/01	Hex Screwdriver , wrench size 2.5 mm, 180 mm
16	64-1181/16	Hex Screwdriver , wrench size 2.0 mm, 175 mm
17	16-0017/01	Separate Rod M5 , 220 mm
18	317-607/50	Cutting Template
19	317-585/65	Wire Pins , Ø 3 mm, 65 mm (4 ea. included)
20	317-585/95	Wire Pins , Ø 3 mm, 95 mm (4 ea. included)
		Adapters optional with fittings:
21	16-3284/00	Hudson female/AO male
	16-3283/00	Hudson female/Jacobs male (without illustration)
	16-3285/00	Hudson female/Harris male (without illustration)

15-8760/01 Container F – Femur I


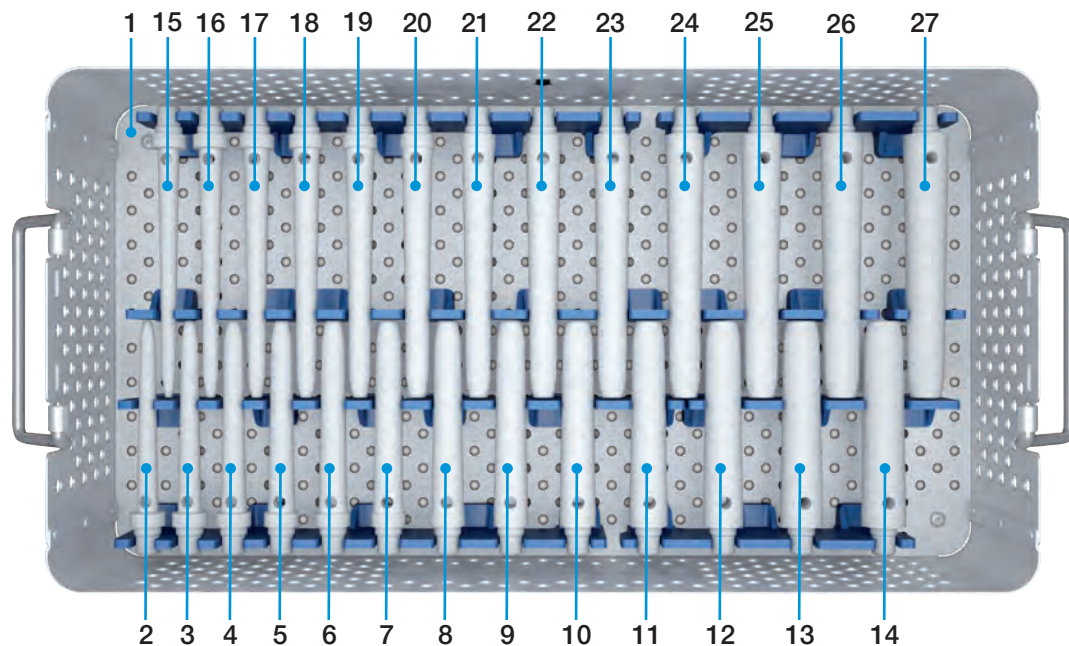
1	05-2001/03 15-8761/01	N11 Standard Container , 575 x 275 x 100 mm (without illustration) Tray , empty, 550 x 265 x 50 mm	
2	Saw Attachments for femoral components SL®		
2	16-3221/01	small S	right
3	16-3223/01	medium M	right
4	16-3225/01	large L	right
5	16-3221/02	small S	left
6	16-3223/02	medium M	left
7	16-3225/02	large L	left
	Femoral Trial Box for femoral components SL®		
8	16-3261/01	small S	right
9	16-3263/01	medium M	right
10	16-3265/01	large L	right
11	16-3261/02	small S	left
12	16-3263/02	medium M	left
13	16-3265/02	large L	left
14	16-3277/00	Revision Cutting Block , distal, for femoral components SL® (2 parts)	
15	16-3278/00	Revision Alignment Gauge , distal, for femoral components SL® (2 parts)	
16	16-3282/00	Alignent Instrument for femoral trial box	
17	16-3271/00	Adapter for femoral trial box	

15-8770/01 Container G – Femur II


1	05-2002/03 15-8771/01	N21 Standard Container , 575 x 275 x 130 mm (without illustration) Tray , empty, 550 x 265 x 50 mm	
		Femur Cutting Blocks for chamfer cuts	
2	16-3250/02	size 2	
3	16-3250/03	size 3	
4	16-3250/04	size 4	
5	317-802/58	Alignment Rod for epicondyles, 100 mm (2 ea. included)	
		Femur Cutting Bblocks for distal cut	
6	16-3228/02	size 2	
7	16-3228/03	size 3	
8	16-3228/04	size 4	
9	16-3276/00	Alignment Instrument , for determination of external rotation (5 parts)	
10	16-3202/00	Drill , conical, to open femoral and tibial cavity, with Hudson fitting	
11	317-802/36	Dove Tail Adapter , neutral, for femoral cutting blocks	
12	16-3275/00	Alignment Instrument for valgus angulation (2 parts)	
13	16-3281/00	Center Sleeve for drill cap femur	
14	16-3206/20	Drill , with stop, with Hudson fitting, Ø 20 mm	
15	16-3279/00	Holding Clamp	
		Condyles Caps Femur	
16	16-3240/02	size 2	right/left
17	16-3240/03	size 3	right/left
18	16-3240/04	size 4	right/left
		Drill Caps Femur , Ø 20 mm	
19	16-3213/02	size 2	right/left
20	16-3213/03	size 3	right/left
21	16-3213/04	size 4	right/left
22	317-802/32	Chisel for patella glide, 80 mm	

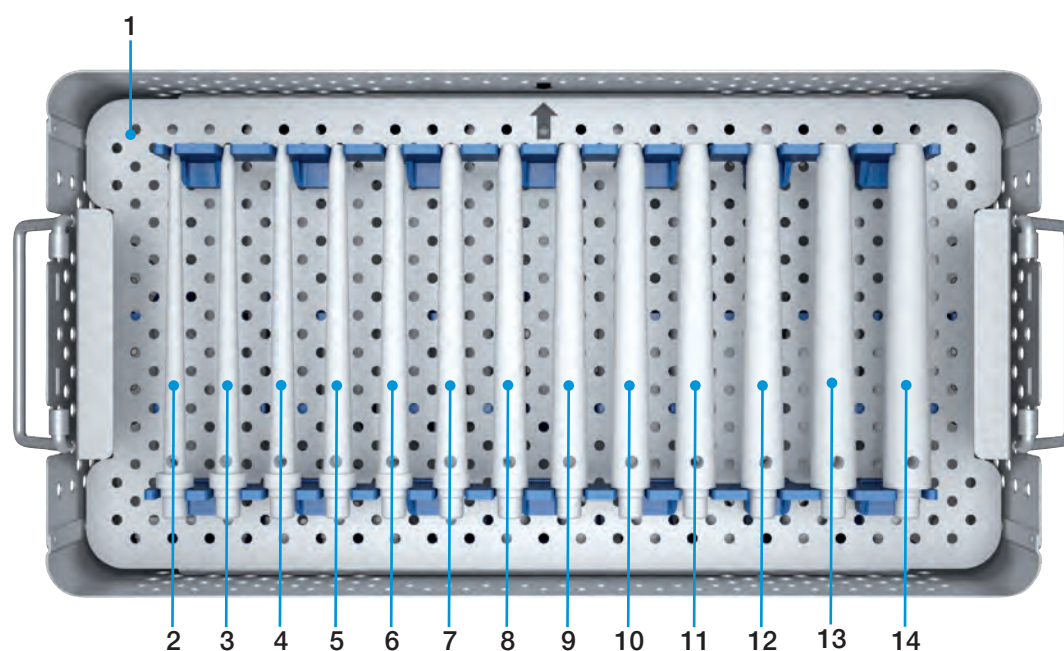
15-8780/01 Container H – Tibia


1	05-2002/03 15-8781/01	N21 Standard Container , 575 x 275 x 130 mm (without illustration) Tray , empty, 550 x 265 x 50 mm
2	317-802/52	Stylus for tibial saw guide, adjustable (2 parts)
3	16-3212/08	Connector , size 8°, for tapered reamer/tibial saw guide
4	16-3197/00	Handle , for tibial compressor/femoral trial box
5	16-3207/16	Drill with stop, for tibial components SL®, ventral side, Ø 16 mm Drill with stop, for tibial components SL®
6	16-3208/18	small S Ø 18 mm
7	16-3208/20	medium M Ø 20 mm
8	16-3208/22	large L Ø 22 mm
9	16-3241/00	Tibial Saw Guide for tibial components SL® Drill Guides for drill templates
10	16-3270/18	small S Ø 18 mm
11	16-3270/20	medium M Ø 20 mm
12	16-3270/22	large L Ø 22 mm Compressors for tibial components SL®
13	16-3199/12	small S
14	16-3199/13	medium M
15	16-3199/14	large L Stem Compressors for tibial components SL®
16	16-3201/02	small S
17	16-3201/03	medium M
18	16-3201/04	large L Drill Templates for tibial components SL®
19	16-3198/12	small S
20	16-3198/13	medium M
21	16-3198/14	large L
22	16-3266/00	Alignment Gauge , tibia, for drill templates
23	16-3267/00	Drill Guide , tibia, for ventral cut, Ø 16 mm
24	16-3211/00	Guide Rod , tibia, for drill template (2 ea. included)
	16-3242/00	Alignment Rod Tibia (without illustration)

15-8790/02 Instrument Set – Femur Trial Stems 100 and 130 mm


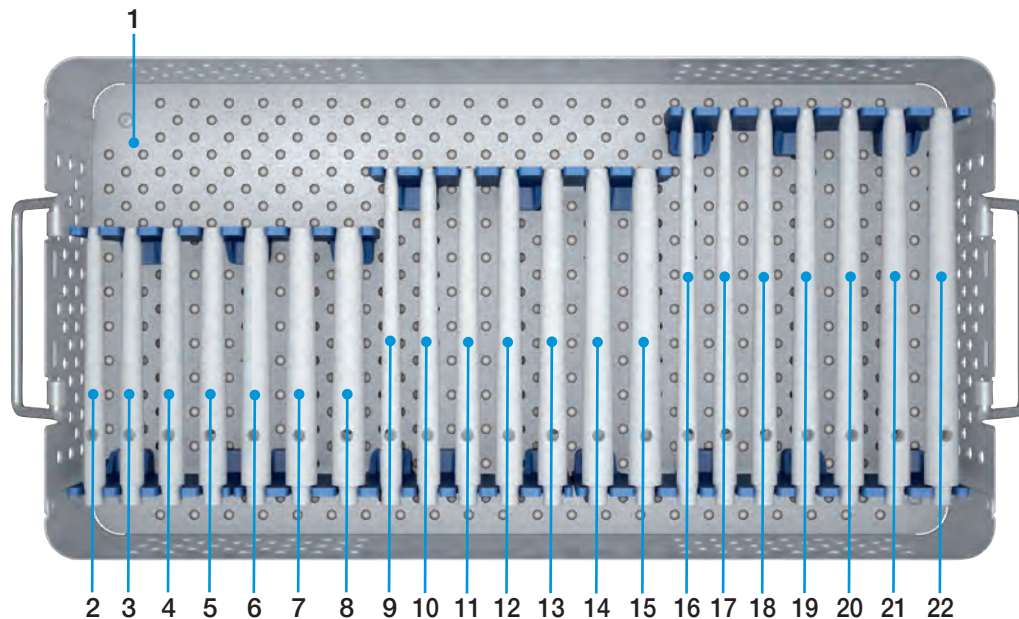
1	15-8791/02	Instrument Tray , empty, 478 x 253 x 106 mm	
		Trial Stems for prosthesis stems 100 mm, conical	
2	16-3101/12	for: stem-Ø 12 mm	stem length 100 mm
3	16-3101/13	for: stem-Ø 13 mm	stem length 100 mm
4	16-3101/14*	for: stem-Ø 14 mm	stem length 100 mm
5	16-3101/15	for: stem-Ø 15 mm	stem length 100 mm
6	16-3101/16*	for: stem-Ø 16 mm	stem length 100 mm
7	16-3101/17	for: stem-Ø 17 mm	stem length 100 mm
8	16-3101/18*	for: stem-Ø 18 mm	stem length 100 mm
9	16-3101/19	for: stem-Ø 19 mm	stem length 100 mm
10	16-3101/20	for: stem-Ø 20 mm	stem length 100 mm
11	16-3101/21	for: stem-Ø 21 mm	stem length 100 mm
12	16-3101/22	for: stem-Ø 22 mm	stem length 100 mm
13	16-3101/23	for: stem-Ø 23 mm	stem length 100 mm
14	16-3101/24	for: stem-Ø 24 mm	stem length 100 mm
		Trial Stems for prosthesis stems 130 mm, conical	
15	16-3131/12	for: stem-Ø 12 mm	stem length 130 mm
16	16-3131/13	for: stem-Ø 13 mm	stem length 130 mm
17	16-3131/14*	for: stem-Ø 14 mm	stem length 130 mm
18	16-3131/15	for: stem-Ø 15 mm	stem length 130 mm
19	16-3131/16*	for: stem-Ø 16 mm	stem length 130 mm
20	16-3131/17	for: stem-Ø 17 mm	stem length 130 mm
21	16-3131/18*	for: stem-Ø 18 mm	stem length 130 mm
22	16-3131/19	for: stem-Ø 19 mm	stem length 130 mm
23	16-3131/20	for: stem-Ø 20 mm	stem length 130 mm
24	16-3131/21	for: stem-Ø 21 mm	stem length 130 mm
25	16-3131/22	for: stem-Ø 22 mm	stem length 130 mm
26	16-3131/23	for: stem-Ø 23 mm	stem length 130 mm
27	16-3131/24	for: stem-Ø 24 mm	stem length 130 mm

* also for cemented stems 12, 14 und 16 mm

15-8790/02 Instrument Set – Femur Trial Stems 160 mm


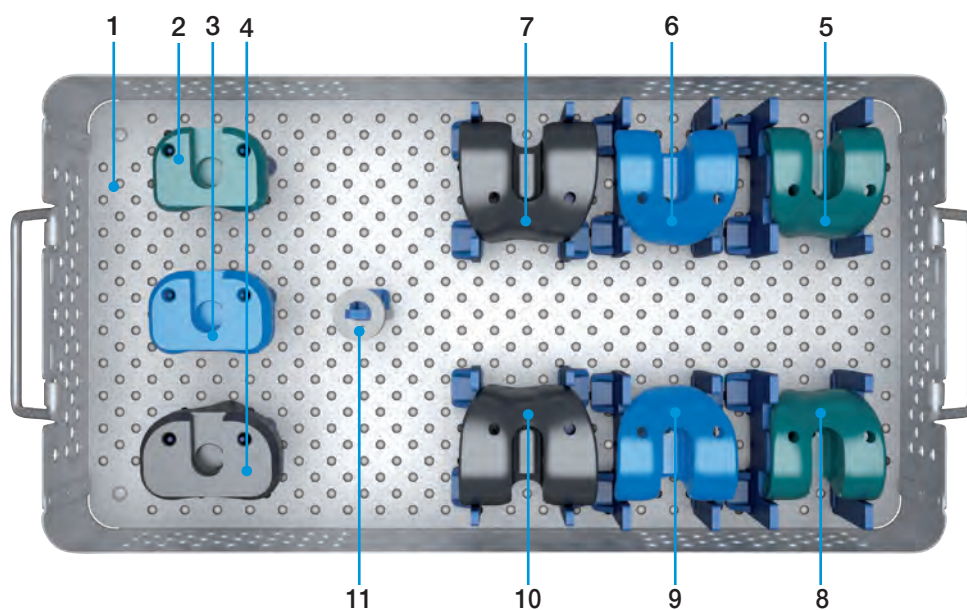
1	15-8791/02	Instrument Tray , empty, 478 x 253 x 106 mm	
		Trial Stems for prosthesis stems 160 mm, conical	
2	16-3161/12	for: stem-Ø 12 mm	stem length 160 mm
3	16-3161/13	for: stem-Ø 13 mm	stem length 160 mm
4	16-3161/14*	for: stem-Ø 14 mm	stem length 160 mm
5	16-3161/15	for: stem-Ø 15 mm	stem length 160 mm
6	16-3161/16*	for: stem-Ø 16 mm	stem length 160 mm
7	16-3161/17	for: stem-Ø 17 mm	stem length 160 mm
8	16-3161/18*	for: stem-Ø 18 mm	stem length 160 mm
9	16-3161/19	for: stem-Ø 19 mm	stem length 160 mm
10	16-3161/20	for: stem-Ø 20 mm	stem length 160 mm
11	16-3161/21	for: stem-Ø 21 mm	stem length 160 mm
12	16-3161/22	for: stem-Ø 22 mm	stem length 160 mm
13	16-3161/23	for: stem-Ø 23 mm	stem length 160 mm
14	16-3161/24	for: stem-Ø 24 mm	stem length 160 mm

* also for cemented stems 12, 14 und 16 mm

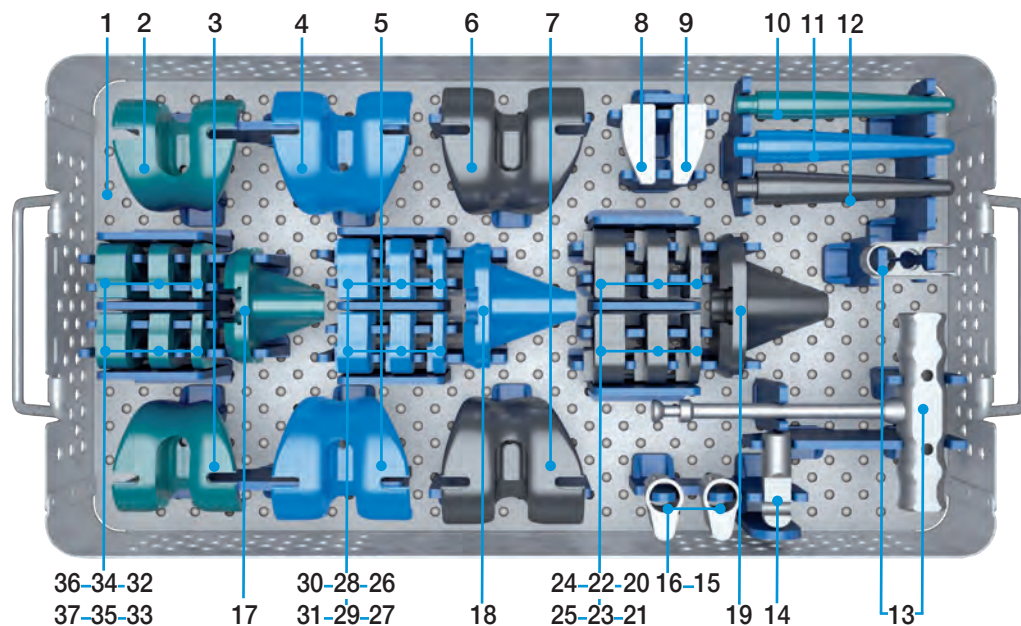
15-8810/02 Instrument Set – Tibial Trial Stems 100, 130 and 160 mm


1	15-8811/02	Instrument Tray , empty, 478 x 253 x 76 mm	
		Trial Stems for modular tibia: 100 mm, conical	
2	16-4101/12	for: stem-Ø 12 mm	for modular tibia 100 mm
3	16-4101/13	for: stem-Ø 13 mm	for modular tibia 100 mm
4	16-4101/14*	for: stem-Ø 14 mm	for modular tibia 100 mm
5	16-4101/15	for: stem-Ø 15 mm	for modular tibia 100 mm
6	16-4101/16*	for: stem-Ø 16 mm	for modular tibia 100 mm
7	16-4101/17	for: stem-Ø 17 mm	for modular tibia 100 mm
8	16-4101/18*	for: stem-Ø 18 mm	for modular tibia 100 mm
		Trial Stems for modular tibia: 130 mm, conical	
9	16-4131/12	for: stem-Ø 12 mm	for modular tibia 130 mm
10	16-4131/13	for: stem-Ø 13 mm	for modular tibia 130 mm
11	16-4131/14*	for: stem-Ø 14 mm	for modular tibia 130 mm
12	16-4131/15	for: stem-Ø 15 mm	for modular tibia 130 mm
13	16-4131/16*	for: stem-Ø 16 mm	for modular tibia 130 mm
14	16-4131/17	for: stem-Ø 17 mm	for modular tibia 130 mm
15	16-4131/18*	for: stem-Ø 18 mm	for modular tibia 130 mm
		Trial Stems for modular tibia: 160 mm, conical	
16	16-4161/12	for: stem-Ø 12 mm	for modular tibia 160 mm
17	16-4161/13	for: stem-Ø 13 mm	for modular tibia 160 mm
18	16-4161/14*	for: stem-Ø 14 mm	for modular tibia 160 mm
19	16-4161/15	for: stem-Ø 15 mm	for modular tibia 160 mm
20	16-4161/16*	for: stem-Ø 16 mm	for modular tibia 160 mm
21	16-4161/17	for: stem-Ø 17 mm	for modular tibia 160 mm
22	16-4161/18*	for: stem-Ø 18 mm	for modular tibia 160 mm

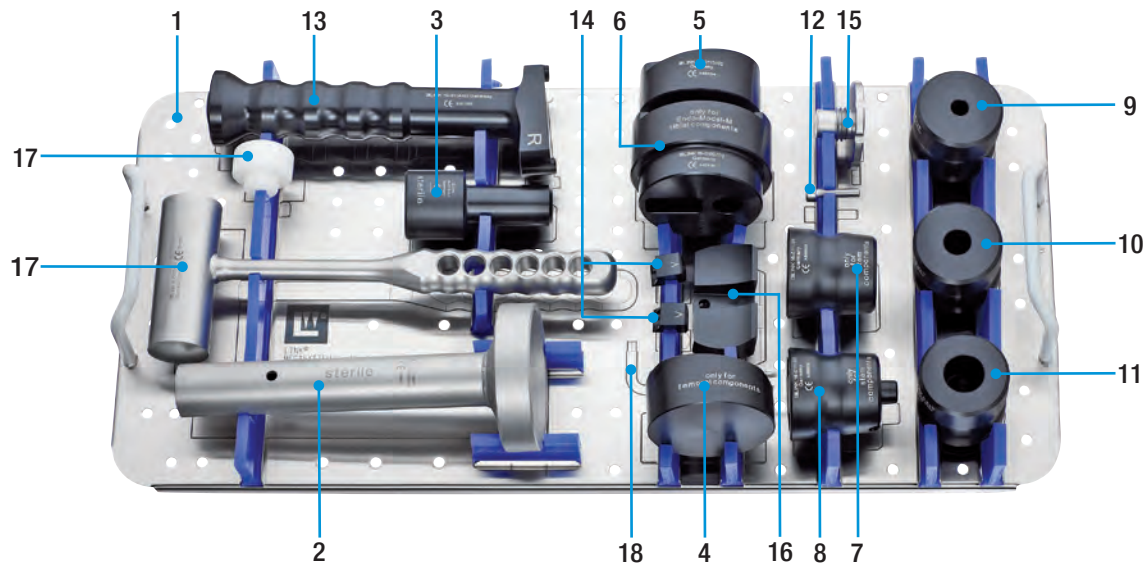
* also for cemented stems 12, 14 und 16 mm

15-8840/02 Instrument Set – Trial Instruments for Distal Femur and Proximal Tibia Replacement


1	15-8841/02	Instrument Tray , empty, 478 x 253 x 76 mm	
		Trial Prosthesis for proximal tibia replacement	
2	16-3180/12	small S	
3	16-3180/13	medium M	
4	16-3180/14	large L	
		Trial Prostheses for distal femur replacement	
5	16-3191/12	small S	left
6	16-3193/12	medium M	left
7	16-3195/12	large L	left
8	16-3191/11	small S	right
9	16-3193/11	medium M	right
10	16-3195/11	large L	right
11	15-6094/00	Trial Support Ring, Ø 28 mm, Height 10 mm	

15-8820/02 Instrument Set – Femur and Tibial Trial Prostheses, intracondylar


1	15-8821/02	Instrument Tray, empty, 550 x 265 x 50 mm		
		Femur Trial Prostheses, intracondylar		
2	16-3181/12	left	small S	
3	16-3181/11	right	small S	
4	16-3183/12	left	medium M	
5	16-3183/11	right	medium M	
6	16-3185/12	left	large L	
7	16-3185/11	right	large L	
8	16-4368/01	Femur Trial Segment, right, size S/M/L, 25 mm height		
9	16-4368/02	Femur Trial Segment, left, size S/M/L, 25 mm height		
10	16-3170/12	Tibial Trial Stems, small S		
11	16-3170/13	Tibial Trial Stems, medium M		
12	16-3170/14	Tibial Trial Stems, large L		
13	15-6061/00	Extraction Instrument for trial prostheses		
14	16-4367/20	Trial Axis for mobile and hinged versions		
15	16-4367/00	Trial Connection Component for hinge knee		
16	16-4367/10	Trial Connection Component for rotational knee		
17	16-3175/12	Tibial Trial Prostheses, intracondylar, small S		
18	16-3175/13	Tibial Trial Prostheses, intracondylar, medium M		
19	16-3175/14	Tibial Trial Prostheses, intracondylar, large L		
		Tibial Trial Spacers, intracondylar		
20	16-4361/01	right	small S	5 mm height
21	16-4362/01	left	small S	5 mm height
22	16-4363/01	right	small S	10 mm height
23	16-4364/01	left	small S	10 mm height
24	16-4365/01	right	small S	15 mm height
25	16-4366/01	left	small S	15 mm height
26	16-4361/02	right	medium M	5 mm height
27	16-4362/02	left	medium M	5 mm height
28	16-4363/02	right	medium M	10 mm height
29	16-4364/02	left	medium M	10 mm height
30	16-4365/02	right	medium M	15 mm height
31	16-4366/02	left	medium M	15 mm height
32	16-4361/03	right	large L	5 mm height
33	16-4362/03	left	large L	5 mm height
34	16-4363/03	right	large L	10 mm height
35	16-4364/03	left	large L	10 mm height
36	16-4365/03	right	large L	15 mm height
37	16-4366/03	left	large L	15 mm height

16-0100/00 Container Assembling Instrument Set


1	05-2001/03 16-0100/01	N11 Standard Container , 575 x 275 x 100 mm (without illustration) Tray , empty, 550 x 265 x 50 mm
2	16-0118/01	Assembling Table: upper part
3	16-0118/03	Assembling Table: silencer
4	16-0110/01	Femur Assembling Plate
5	16-0110/02	Tibia Assembling Plate für Endo-Model® SL® knee prostheses
6	16-0110/03	Tibia Assembling Plate für Endo-Model® knee prostheses
7	16-0111/01	Assembling Protector for female taper
8	16-0111/02	Assembling Protector for male taper
9	16-0112/01	Assembling Protectors , size 1
10	16-0112/02	Assembling Protectors , size 2
11	16-0112/03	Assembling Protectors , size 3
12	16-0113/01	Metal Core for assembling protector, for modular stem (size 1 – 3)
13	16-0114/01	Handle for femur assembling protector
14	16-0114/02	Block , spare part for femur assembling protector 16-0114/01 (2 ea. included)
15	16-0114/04	Screw for assembling protectors femur
16	16-0114/03	Plate , spare part for femur assembling protector 16-0114/01
17	16-0115/01	Mallet , non-resilient, 800g, complete
18	16-0116/02	Hex Screwdriver , wrench size 5 mm, conical
	16-0118/02*	Assembling Table: lower part

* Not included in container 16-0100/00.

Assembling Instruments

for Endo-Model® SL® Rotating and Non-Rotating Hinge Knee



16-0118/01

Assembling Table: Upper Part



16-0118/02

Assembling Table: Lower Part
with fixed base plate



16-0118/03

Assembling Table: Silence



16-0110/01
Femur Assembling Plate



16-0110/02
Tibia Assembling Plate
for Endo-Model® SL® Knee Prostheses

16-0110/03
Tibia Assembling Plate
for Endo-Model® Knee Prostheses



16-0111/01
Assembling Protector for female taper



16-0111/02
Assembling Protector for male taper



Assembling Protectors
16-0112/01 for modular stem, size 1
16-0112/02 for modular stem, size 2
16-0112/03 for modular stem, size 3



16-0113/01
Metal Core for assembling protector,
for modular stem (size 1-3)



16-0114/01

Femur Assembling Protector,
complete



16-0114/02

Block, for femur assembling protector
(Spare part included in 16-0114/01)



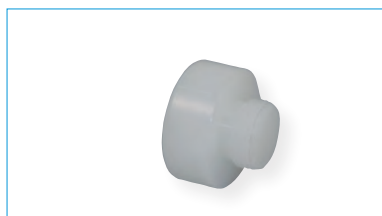
16-0114/03

Plate, for femur assembling protector
(Spare part included in 16-0114/01)



16-0115/01

Mallet, non-resilient proof, 800g,
complete



16-0115/02

Spare Mallet Head
(Spare part included in 16-0115/01)



16-0116/02

Hex Screwdriver,
SW 5 mm

Description of Use: Mounting the Assembling Table

The Assembling Table is mounted before the operation as illustrated below.

The Superior Component (16-0118/01) and the Silencer (16-0118/03) are assembled in sterile condition. The Inferior Component (16-0118/02) with fixed Base Plate may be kept non-sterile.



The Silencer is screwed onto the Inferior Component.



The Superior Component is attached.

In use of the Assembling Table, place foot on base plate to prevent the table from toppling over.

The table is dismantled in reverse order.

Description of Use: Connecting modular Stems to Modular Femoral and Tibial Components

The Femur Assembling Plate (16-0110/01) is placed into the reception of the Superior Component of the Assembling Table (16-0118/01).



According to the chosen size of modular stem, the Metal Core (16-0113/01) is screwed into Assembling Protector Modular Stem, size 1-3 (16-0112/01-03).



Prosthesis stem and femoral joint component are connected by hand (the taper surfaces must be clean and dry) and, observing the 6° valgus alignment of the femoral implant, placed on the



Assembling Plate so that the prosthesis stem is vertical. If not, the implant assembly must be rotated 180°.

As shown in the picture, the components are struck with two powerful blows of the non-resilient mallet (16-0115/01) in order to firmly engage the taper connection (the second blow is a safety measure). The metal face of the Mallet must only be used.



When using Modular Tibial Implants, the Tibial Assembling Plate (16-0110/02 or 16-0110/03) is first placed into the reception of the Superior Table Component.



Proceed according to the assembly of Modular Femoral Implant. Before using the Locking Screws, the taper connections must be connected firmly as described, using the Assembling Instruments.

Description of Use: Connecting Stem Elements

The Assembling Protector for Female Taper (16-0111/01) is placed into the reception of the assembling Table Superior Component.



Attaching the selected Stem Element.



Attaching the Stem Element to be connected and the Assembling Protector for Male Taper (16-0111/02).

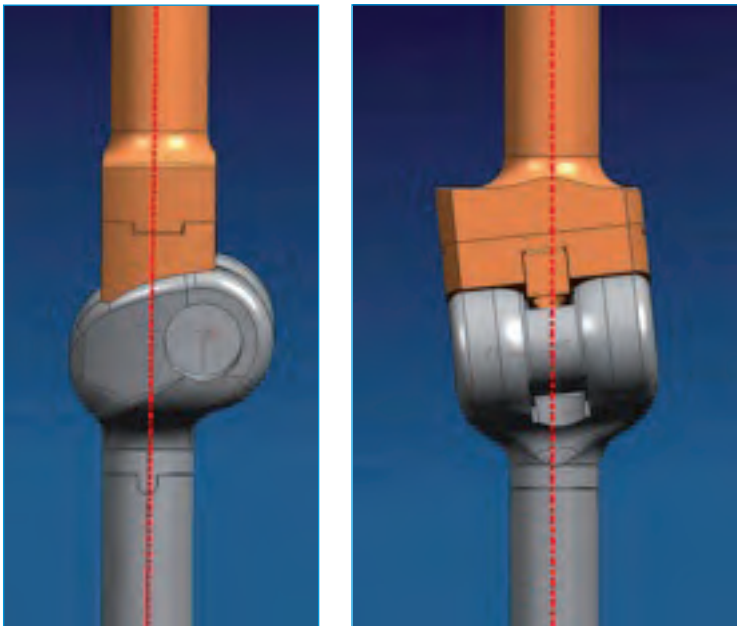


Assemble the elements with two blows of the non-resilient Mallet, using the metal face of the Mallet only.

Use and Mounting of the Femur Assembling Protector

Use

The femur assembling protector is always used if the surgical procedure does not permit assembly of implant components outside the operating field (e.g. connecting femoral joint components to push-through stems). The instrument compensates for the valgus geometry of the knee implant and permits the engagement of the tapers in axial alignment in the transverse and sagittal plane.

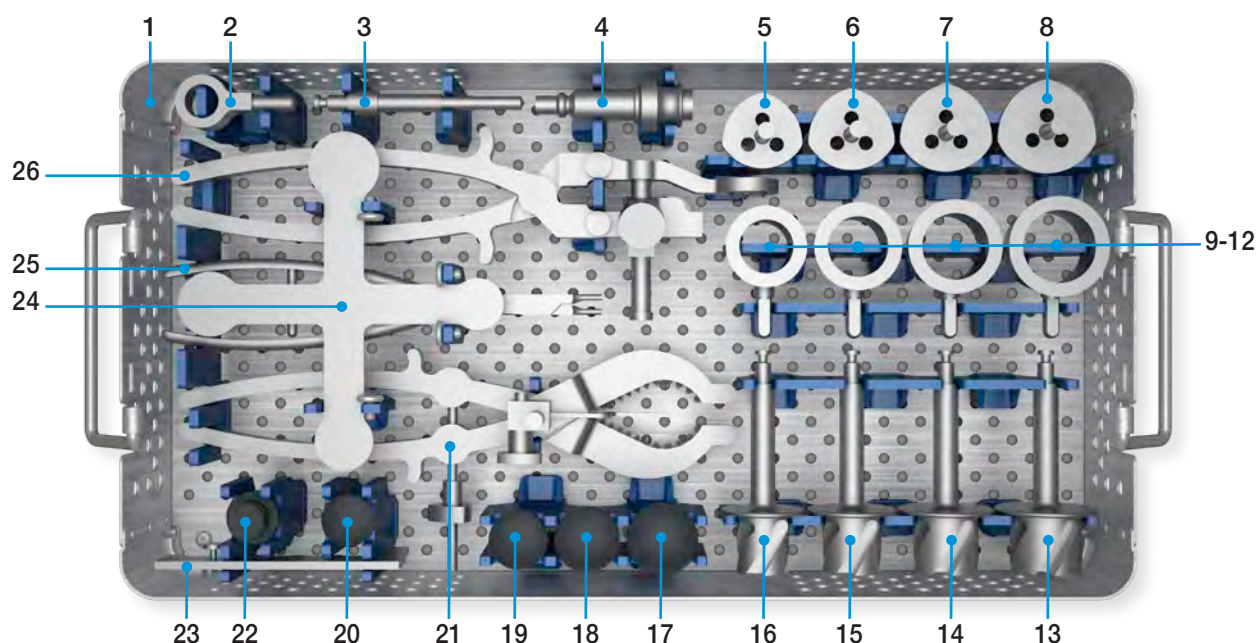


Its use enables taper engagement in axial alignment.

Assembly

The femur assembling protector is mounted as shown in below illustration. First the plate is connected to the handle so that, depending on the side being operated on, the letters “R” for the right side or the letters “L” for the left side are lined up. In the illustration the right side is chosen. Consecutively the block for Assembling Protector is secured using the Hex Screwdriver. The two arrows (of block and plate) must point toward each other.





340-200/01 Instrument Set for Patella Components, 3-pegs for LINK® knee family SL®


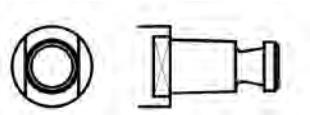
1	340-015/01	Tray, empty, 485 x 253 x 100 mm (incl. lid)
2	340-007	Patella Pusher Attachment
3	340-306B	Patella Drill, Ø 6.3 mm, with Hudson B fitting
4	optional*	Adapter for snap lock chuck, optional
		Patella Drill Guides for patella holding clamp
5	340-225	Size 25 for 318-401/25
6	340-228	Size 28 for 318-401/28
7	340-231	Size 31 for 318-401/31
8	340-234	Size 34 for 318-401/34
		Patella Reaming Guides for patella holding clamp
9	340-025	Size 25 for 318-401/25
10	340-028	Size 28 for 318-401/28
11	340-031	Size 31 for 318-401/31
12	340-034	Size 34 for 318-401/34
		Patella Reamers for patella reaming guides, with Hudson B fitting
13	340-134B	Size 34 for 318-401/34
14	340-131B	Size 31 for 318-401/31
15	340-128B	Size 28 for 318-401/28
16	340-125B	Size 25 for 318-401/25
		Patella Trial Prostheses for Patella Component 3-pegs
17	340-334	Size 34 for 318-401/34
18	340-331	Size 31 for 318-401/31
19	340-328	Size 28 for 318-401/28
20	340-325	Size 25 for 318-401/25
21	340-006	Patella Resection Clamp
22	340-009	Patella Pusher Inserter
23	65-2000	Townley Femur Caliper, 115 mm
24	340-010	Patella Sizing Template
25	15-2042	Inserting Forceps for inserting the manipulating bearings, 215 mm
26	340-005	Patella Holding Clamp

* Adapter optional: 16-3283/00, 16-3284/00, 16-3285/00

Additional Instruments

Adapter for Snap Lock Chuck with Hudson Fitting (B)
Various adapters to enable compatibility with other equipment connections.

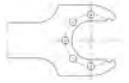



Item no.	Fitting	
16-3283/01	Jacobs Fitting (E)	
16-3284/00	AO Fitting (D)	
16-3285/00	Harris Fitting (C)	



Hudson Fitting (B)
Basic tool connection
for patella instruments



Sawblades,
without offset teeth, 1.24 mm thick

Wide (A) 25 mm	Wide (A) 13 mm	Fitting
317-654/10	317-656/10	Synthes 
317-654/11	317-656/11	Aesculap combi 
317-654/13	317-656/13	Zimmer / Hall Kombi 
317-654/14	317-656/14	Stryker System 4 

Additional Instruments



Revision Drill Guide

for mobile and hinged version, to support the axis decoupling when changing the Endo-Model® SL® connection components

Item no.	complete
16-3295/00	bestehend aus 4 Teilen: Halterung, Bohrerstab, 2 Stäbe mit Griff

Extractor

for modular stems with female taper, taper 12/10 mm (for 3 mm + 6 mm noses)

Item no.	Length
15-0036/81	230 mm

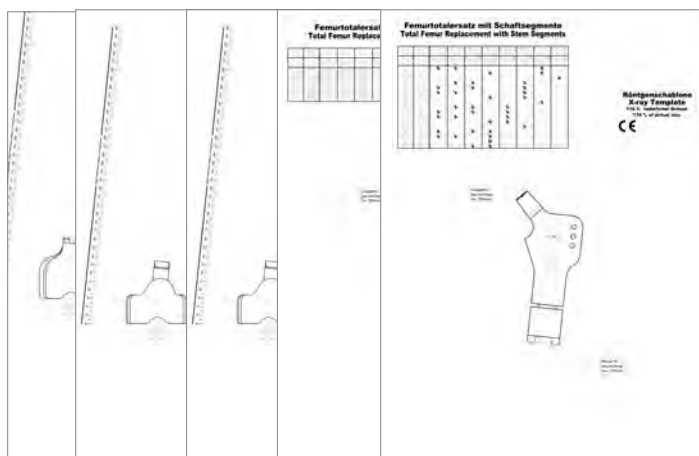
Guide Rod with Slaphammer

(without illustration)

(optional for stem extraction: Guide Rod 317-661 in combination with Driver Extractor 15-8516/45)

Item no.	Length
317-661	365 mm

Accessories



X-ray Templates,

110% of actual size, set = 22 sheets

Item no.	
15-8516/61	for LINK® MEGASYSTEM-C® with SL® knee components

Surface Modifications (available as custom-made prosthesis on request)

- a) Cementless Modular Stems PoroLink[®]*
optional with HX[®]** coating.

* PoroLink[®]: grid blasted titanium surface Ra ~7µm
** HX[®]: CaP = Calcium Phosphate coating

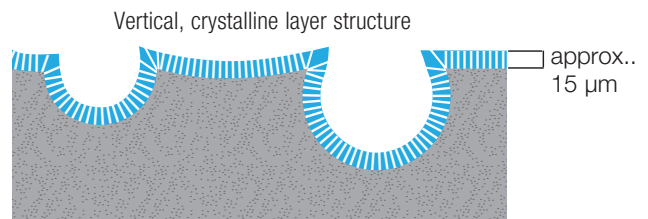


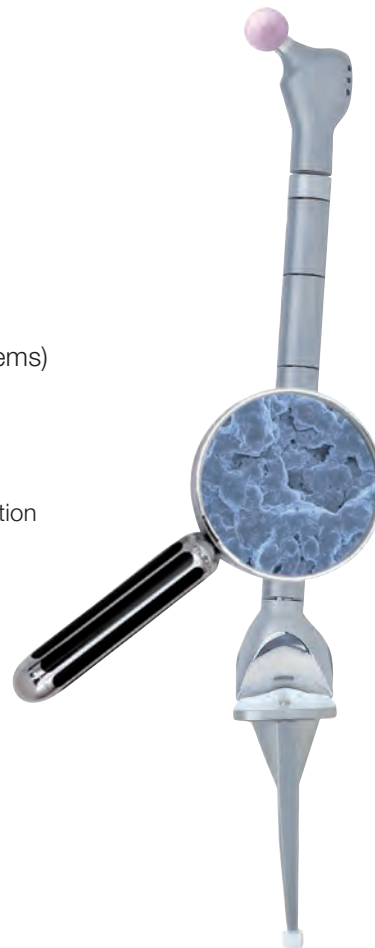
Fig.: PoroLink[®]* with HX[®]** coating.
Schematic cross-section of coated surfaces.



Catalog available on request.

- b) Bone Replacement Components CoCrMo
for metal hypersensitive patients, optional with
LINK PorEx[®]*** Technology.

*** LINK PorEx[®]: TiNbN = Titanium Niobium Nitride



- c) Bone Replacement Components (no intramedullary stems)
with PorAg[®]****.

**** PorAg[®]: antimicrobial surface modification
TiAgN/Ag = titanium silver nitride/silver surface modification



MEGASYSTEM-C®
Tumor and Revision System

- Implants & Instruments
- Surgical Technique



LinkSpacer
PMMA/Gentamicin


Synimed Synergie Ingénierie Médicale S.A.R.L., France
Distributor: Waldemar Link GmbH & Co. KG, Germany



TrabecuLink® Tibial Cones

- Product Rationale
- Surgical Technique

All catalogs available on request.



For more information please register for our LINK Media Library (linkorthopaedics.com)

Please note the following regarding the use of our implants:

1. Choosing the right implant is very important.

The size and shape of the human bone determines the size and shape of the implant and also limits the load capacity. Implants are not designed to withstand unlimited physical stress. Demands should not exceed normal functional loads.

2. Correct handling of the implant is very important.

Under no circumstances should the shape of a finished implant be altered, as this shortens its life span. Our implants must not be combined with implants from other manufacturers.

The instruments indicated in the Surgical Technique must be used to ensure safe implantation of the components.

3. Implants must not be reused.

Implants are supplied sterile and are intended for single use only. Used implants must not be used again.

4. After-treatment is also very important.

The patient must be informed of the limitations of the implant. The load capacity of an implant cannot compare with that of healthy bone!

5. Unless otherwise indicated, implants are supplied in sterile packaging.

Note the following conditions for storage of packaged implants:

- Avoid extreme or sudden changes in temperature.
- Sterile implants in their original, intact protective packaging may be stored in permanent buildings up until the "Use by" date indicated on the packaging.
- They must not be exposed to frost, dampness or direct sunlight, or mechanical damage.
- Implants may be stored in their original packaging for up to 5 years after the date of manufacture. The "Use by" date is indicated on the product label.
- Do not use an implant if the packaging is damaged.

6. Traceability is important.

Please use the documentation stickers provided to ensure traceability.

7. Further information on the material composition is available on request from the manufacturer.

Follow the instructions for use!

Waldemar Link GmbH & Co. KG, Hamburg

All content in this catalog, including text, pictures and data, is protected by law. Every instance of use, whether in part or in whole and which is not permitted by law, is subject to our prior consent. In particular, this applies to the reproduction, editing, translation, publishing, saving, processing, or passing on of content stored in databases or other electronic media and systems, in any manner or form. The information in the catalogs is solely intended to describe the products and does not constitute a guarantee.

The Surgical Technique described has been written to the best of our knowledge and belief, but it does not relieve the surgeon of his/her responsibility to duly consider the particularities of each individual case.

Waldemar Link GmbH & Co. KG

Barkhausenweg 10 • 22339 Hamburg, Germany

Phone +49 40 53995-0 • info@linkhh.de

www.linkorthopaedics.com

