

Endo-Model SL

Rotational and Hinge Knee Prosthesis System

Surgical Technique



C€ 0482

Explanation of Pictograms

	Manufacturer	REF	Item number
MAT	Material (number)	CE	Product meets the applicable requirements, which are regulated in the EU harmonization legislation for the affixing of the CE marking.

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Surgical Technique

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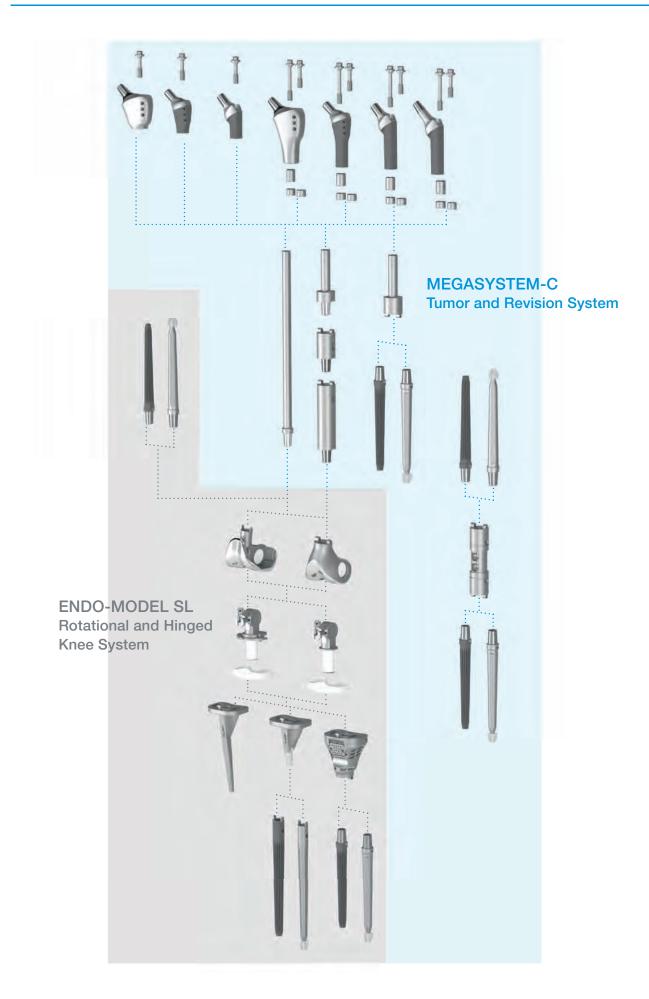
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Important Information

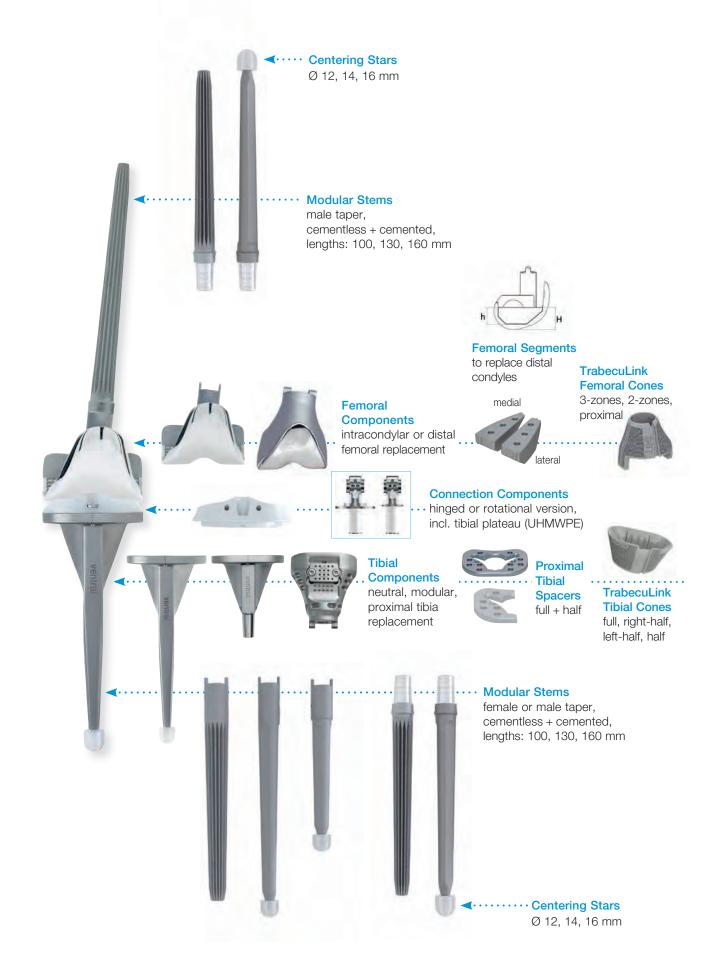
Modular System Overview





Varity of Assembly







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 Prosthesis, 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 revision surgery. 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 (15-6037/00).



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.

CAUTION:

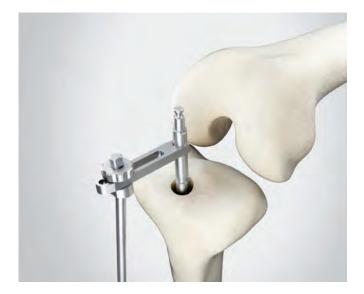
The position of the impaction plate represents the level of the joint line. Using the awls with a power tool is not permitted.





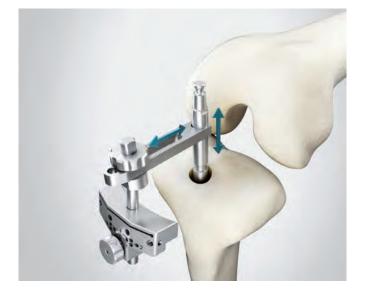
03

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 levelof 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 mmmust be used.





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

Removal of the alignment gauge. The awl \emptyset 19 – 24 mm must also be removed temporarily and then reinstalled.





Attach 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.



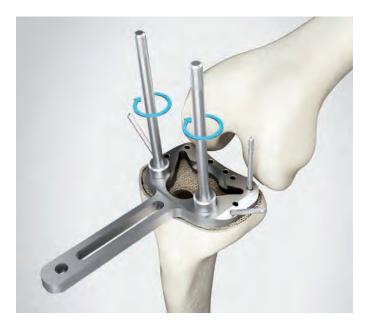
13

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.





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.





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 (15-6037/00).



20 + 21

Mount the awl in the planned length (100 mm, 130 mm or 160 mm) in 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 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.



CAUTION:

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





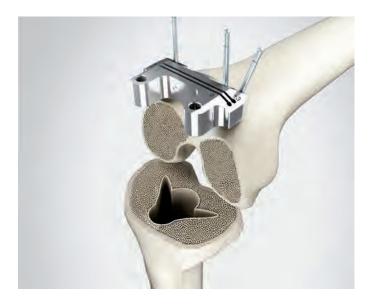
Remove the impaction plates and attach the alignment instrument for valgus angulation (16-3275/00). Ensure that the correct instrument for the right or left side is attached. The word **"Left"** or **"Right"** must face upward.



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.





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.





The femur 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). Two 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 driver extractor 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.





The condyle cap (16-3241/02, /03, /04) is placed on the prepared bone surfaces. The shaft of the awl forms the center.

Through the ventral saw slots the bone can be prepared, matched to the femoral segments in 15 mm, 25 mm or full.



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

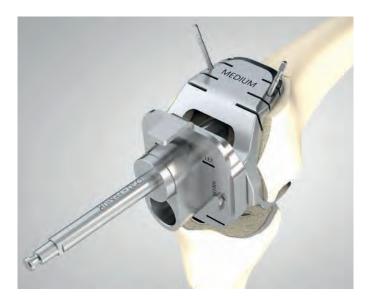
Using the centre sleeve (16-3281/00), the instruments are aligned and centered on the shaft of the awl.





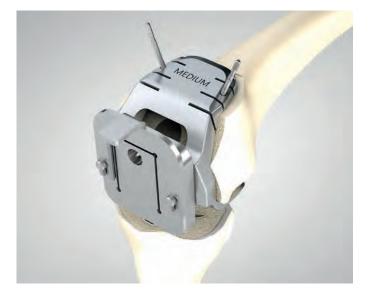
34

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 two wire pins. The center sleeve and awl are removed If necessary, the drill cap must also be removed temporarly and then re-attached again.



35

The drill for femur \emptyset 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.





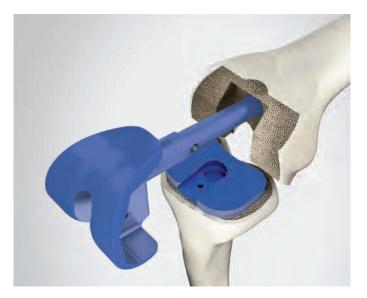
37

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 with femoral trial segments and/or tibial trial spacers. The components are separated by proceeding in reverse order.

INFORMATION:

A combination of serveral 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.



INFORMATION:

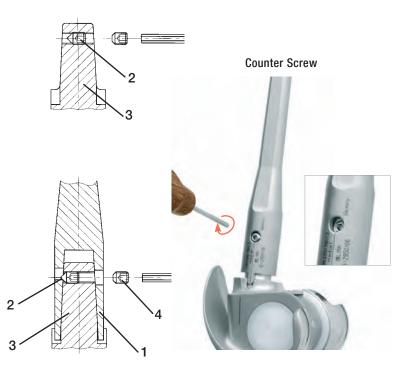
When using cemented modular stems or the monobloc tibial component, the use of centering stars is mandatory. They might not be part of the sterile package for tibial components 16-2817/02, /05 and /07 when indicated.

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.

INFORMATION:

The locking srew (optionally preassembled), the counter screw and one replacement screw each are included separately in the packaging.

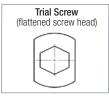






44 + 45

Until the bone cement has set, the trial screw remains in the prosthesis thread in order to protect it (excess bone cement is removed). After the cement has set, the screw is removed with the hex screwdriver (64-8008/02).



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).

INFORMATION:

If the intention was to use a tibial component with pre-assembled PE sleeve and tension plate these parts are no longer required, as they are now form part of the connecting components.

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



46

2

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

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

If required, the PE plateau can be inserted after the connection component (rotating version) has been assembled.





47 + 48

The connector with the rotation axis is put on the tibial component, and the PE plateau with the inserter (15-8035/03) is then inserted and screwed home. The self-locking screw is screwed with the hex screwdriver (64-8008/02). For easier access to the plateau retaining screw, the connector is rotated slightly.



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 + 50

The connecting component is inserted into the intracondylar slot of the femoral component. The tibia is placed as far posteriorly as possible, relative to the femur. Then the U-shaped opening of the securing sleeve is rotated 180° until the opening faces upwards. The securing sleeve can then be pulled upwards along the spring wire, thereby unloading the spring so that it presses the axis halves outwards. The prosthesis axis is inserted into the bushings by slightly lifting the connecting component and carefully moving the joint components. A 2-mm-torque wrench hex screwdriver (15-2546) is used to tighten the securing screw, which

is already in the sleeve. Then the axis lock is removed.

The screw must be tightened until an audible "Click" is heard that indicates that the required torque has been applied, locking the connection component in place.

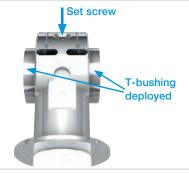




51 + 52

Check the position of the holes. When assembly is complete, the two drill holes in the prosthetic axis are located exactly under the arrow marks. Confirm that the set screw is slightly recessed. If it is not, remove the set screw, deploy the T-bushing with the connecting and separating forceps (16-0020/01 or /02) and reinsert the set screw, with the 2-mm-torque screwdriver (15-2546) until it is recessed.









53 + 54

If use of a non-rotating hinged knee version is planned, the screws in the tibial plateau must first be removed with the hex screwdriver (15-2546). 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.











55

When assembly is complete, the two drill holes in the prosthetic axis are located exactly under the arrow marks. The PE plateau is then screwed home with the hex screwdriver (64-8008/02).

To decouple the connecting component, in the event of a revision, the securing screw is removed, the connecting and separating forceps (16-0020/01 or /02) are inserted, and the prosthetic axis is squeezed together. Then the connecting component can be decoupled ventrally.



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 seleted 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

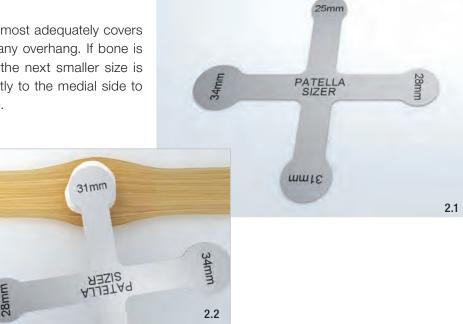


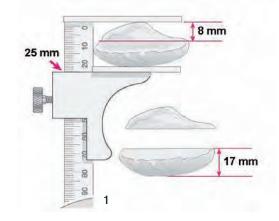
The patella surface can be prepared using the onlay technique by resecting with an oscillating saw or by reaming.

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).

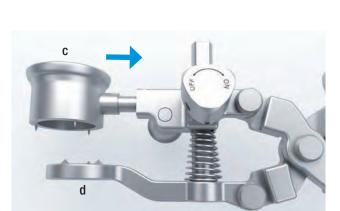
Option: Reaming

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).

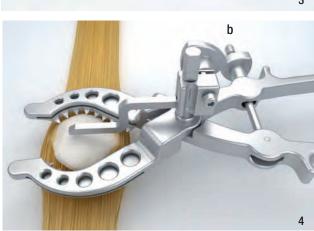
INFORMATION:

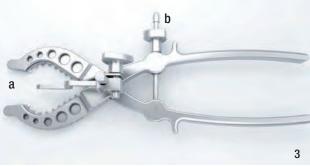
Press the button on the side to insert and remove the guide.



5

6









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 spikeson the guide grip into the bone.

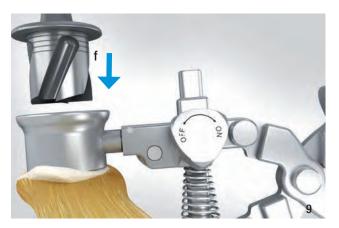


Reaming

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).

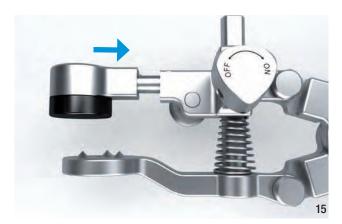
In doing so, ensure that the guide pin on the drill guide lies in the opening made for it in the reaming guide. Using the \emptyset 6.3 mm patella drill (340-306B) (h) three holes guided by the drill guide are drilled. The patella drill is inserted by the until stop (12).





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

All size-specific implants for the Endo-Model SL only allow a combination of the same size (applies from page 32 to page 39)



Femoral Components, intracondylar,

with patellar flange, modular **MAT** CoCrMo, UHMWPE

REF	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





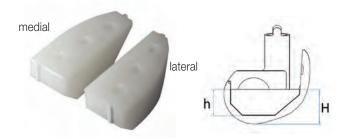
Distal Femoral Replacement Components, slim

modular MAT CoCrMo, UHMWPE

REF	Size	Side	M/L - A/P
16-2853/31	small (S)	right	60.0 mm - 57.0 mm
16-2853/32	small (S)	left	60.0 mm - 57.0 mm
16-2855/31	medium (M)	right	65.0 mm - 62.0 mm
16-2855/32	medium (M)	left	65.0 mm - 62.0 mm
16-2857/31	large (L)	right	75.0 mm - 65.0 mm
16-2857/32	large (L)	left	75.0 mm - 65.0 mm



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

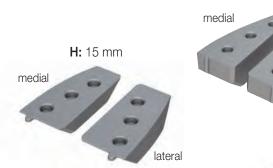


Femoral Segments, cemented,

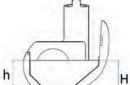
for femoral components, to replace distal condyles

REF	Size	Side		h	н
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





H: 25 mm



lateral

Femoral Segments, cementless, for femoral components, to replace distal condyles, MAT Tilastan

REF	h	н	Size	Side	
15-8570/01	8 mm	15 mm	small	right	medial
15-8570/02	9 mm	15 mm	medium	right	medial
15-8570/03	7 mm	15 mm	large	right	medial
15-8570/11	8 mm	15 mm	small	right	lateral
15-8570/12	9 mm	15 mm	medium	right	lateral
15-8570/13	7 mm	15 mm	large	right	lateral
15-8571/01	8 mm	15 mm	small	left	medial
15-8571/02	9 mm	15 mm	medium	left	medial
15-8571/03	7 mm	15 mm	large	left	medial
15-8571/11	8 mm	15 mm	small	left	lateral
15-8571/12	9 mm	15 mm	medium	left	lateral
15-8571/13	7 mm	15 mm	large	left	lateral
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



Femoral Segments, full, cementless

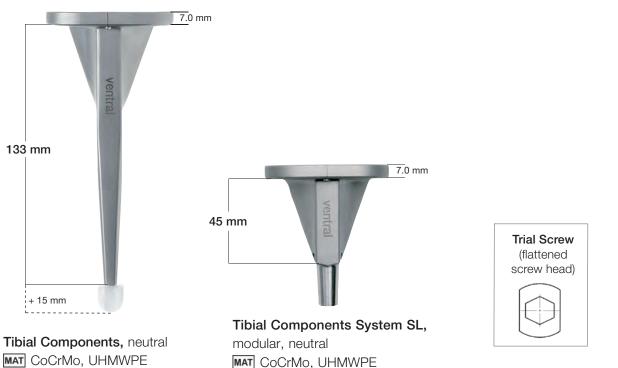
for femoral components, to replace distal condyles, MAT Tilastan

REF	Size	Side	
15-8572/01	small	right	medial
15-8572/02	medium	right	medial
15-8572/03	large	right	medial
15-8572/11	small	right	lateral
15-8572/12	medium	right	lateral
15-8572/13	large	right	lateral
15-8573/01	small	left	medial
15-8573/02	medium	left	medial
15-8573/03	large	left	medial
15-8573/11	small	left	lateral
15-8573/12	medium	left	lateral
15-8573/13	large	left	lateral

Implants



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



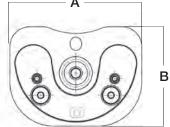
REF	Size	A x B mm
16-2817/02	small (S)	60 x 45
16-2817/05	medium (M)	65 x 45
16-2817/07	large (L)	75 x 48

MAT CoCrMo, UHMWPE $\mathbf{A} \times \mathbf{B}$ REF Size mm 16-2817/32 small (S) 60 x 45 16-2817/35 medium (M) 65 x 45

75 x 48

16-2817/37 large (L)







Proximal Tibia Replacement Components, modular, neutral MAT Tilastan, UHMWPE

REF	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 Plateau (UHMWPE)

REF	Rotating Hin MAT CoCrMc UHMWPE	•	REF	Non-rotating MAT CoCrMc UHMWPE		Arthrodesis Fusion Version* MAT CoCrMo, UHMWPE
16-2840/02	small (S)	neutral	16-2841/02	small (S)	neutral	Knee fusion option
16-2840/05	medium (M)	neutral	16-2841/05	medium (M)	neutral	for in-situ femoral and
16-2840/07	large (L)	neutral	16-2841/07	large (L)	neutral	tibial components.

* 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,

Т

for right and left, incl. 2 countersunk screws, wrench size 2.5 mm, \fbox{MAT} Tilastan

REF	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, MAT UHMWPE

REF	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 with each other!





Proximal Tibial Spacers, half,

lateral and medial usable, incl. 1 countersunk screw, wrench size 2.5 mm, \fbox{MAT} Tilastan

REF	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!

Replacement Sets

Sparepart Kit Bearing Boxes for Endo-Model SL

MAT UHMWPE

REF	Side	Size
16-2011/21	right	small
16-2011/22	left	small
16-2013/21	right	medium
16-2013/22	left	medium
16-2015/21	right	large
16-2015/22	left	large





Modular Stems, with male taper



Modular Stems* with	ı <u>male</u> taper	(for 6 mm	flanges)
MAT Tilastan			

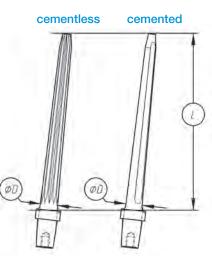
Cementless						
	Stem length		Stem length		Stem length	
REF	L 100 mm	REF	L 130 mm	REF	L 160 mm	
15-8524/50	Ø 12/9 mm	15-8523/50	Ø 12/8 mm	15-8522/50	Ø 12/7 mm	
15-8524/51	Ø 13/10 mm	15-8523/51	Ø 13/9 mm	15-8522/51	Ø 13/8 mm	
15-8524/52	Ø 14/11mm	15-8523/52	Ø 14/10 mm	15-8522/52	Ø 14/9 mm	
15-8524/53	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 <u>male</u> taper (for 6 mm flanges)

	Cemented						
REF	Stem length L 100 mm	REF	Stem length L 130 mm	REF	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:

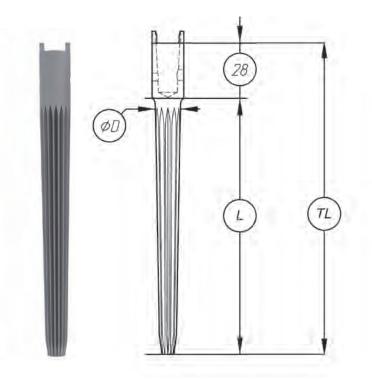
- coupling device to assemble a proximal partial femur replacement
- femoral joint component, solid, or a stem segment attached to it to assemble a distal partial femur replacement
- coupling device for the diaphyseal spacer
- proximal tibial replacement







Modular Stems, with female taper



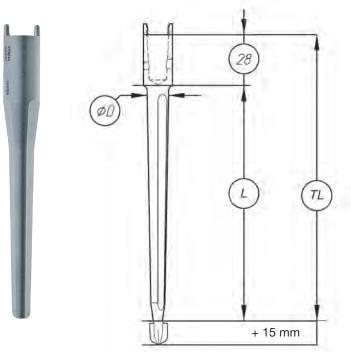
Modular Stems* with <u>female</u> taper (with 6 mm flanges) **MAT** Tilastan

Cementless					
REF	Stem length L 100 mm	Total length TL 128 mm	REF	Stem length L 130 mm	Total length TL 158 mm
15-8517/50	Ø 12/9 mm		15-8516/50	Ø 12/8	3 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/-	l1 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 18/1	5 mm	15-8516/56	Ø 18/-	I4 mm

REF	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*

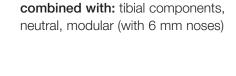
with <u>female</u> taper (with 6 mm flanges), MAT CoCrMo

Cemented					
REF	Stem length L 100 mm	Total length TL 128 mm	REF	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

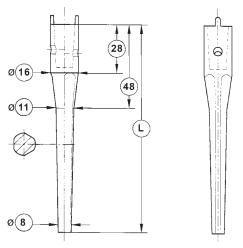
REF	Stem length L 160 mm	Total length TL 188 mm
15-8525/40	Ø 12/7 mm	
15-8525/42	Ø 14/9 mm	
15-8525/44	Ø 16/11 mm	

Modular Stems, short MAT CoCrMo

Cemented		
REF	Stem length L mm	
15-2950/01	50	
15-2950/02	80	



* These modular stems can be





Centering Stars

Centering Stars, height 15 mm MAT UHMWPE

REF	ltem no.	Ø mm
Set consisting of:		
15-2975/01	15-2975/12	





Ø 16 mm

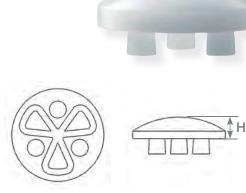
LINK Patella Components

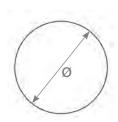
for Endo-Model SL Rotating and Non-Rotating Hinge Knee

LINK Patella Components

MAT UHMWPE

REF	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

15-8710/02	Instrument Set	Tapered Reamers: 100 mm
consisting of: 15-8711/02	Instrument Tray, empty 478 x 253 x 76 mm	
	Tray with detailed instruments see page 48	
15-8720/02	Instrument Set	Tapered Reamers: 130 mm
consisting of: 15-8721/02	Instrument Tray, empty 478 x 253 x76 mm	
	Tray with detailed instruments see page 49	
15-8730/02	Instrument Set	Tapered Reamers: 160 mm
consisting of: 15-8731/02	Instrument Tray, empty 478 x 253 x76 mm Tray with detailed instruments see page 50	
	Instrument Cot	General Instruments I
15-8740/02	Instrument Set	
15-8740/02 consisting of: 15-8741/02	Instrument Set Instrument Tray, empty 478 x 253 x76 mm	



15-8760/02	Instrument Set	Femur I
consisting of: 15-8761/02	Instrument Tray, empty 478 x 253 x 76 mm	
	Tray with detailed instruments see page 52	
15-8770/02	Instrument Set	Femur II
consisting of: 15-8771/02	Instrument Tray, empty 478 x 253 x 76 mm	
	Tray with detailed instruments see page 53	
15-8780/02	Instrument Set	Tibia
consisting of: 15-8781/02	Instrument Tray, empty 478 x 253 x 76 mm	
	Tray with detailed instruments see page 54	
15-8790/02	Instrument Set (1)	Femur Trial Stems: 100 and 130 mm
consisting of: 15-8791/02	Instrument Tray, empty, below 478 x 253 x 106 mm	
	Tray with detailed instruments see page 55	



15-8790/02	Instrument Set (2)	Femur Trial Stems, 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	Tibial 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	
15-8820/03	Instrument Set	Trial Prostheses: Femur/Tibia, intracondylar
consisting of: 15-8821/03	Instrument Tray, empty, 478 x 253 x 76 mm Tray with detailed instruments see page 58	
15-8840/03	Instrument Set Trial Instrume	nts: 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 59	



16-0100/02	Instrument Set	Assembling Instrument Set
consisting of:		
16-0100/03	Instrument Tray, empty	
	478 x 253 x 86 mm	
	Tray with detailed instruments see page 60	



15-8710/02 Instrument Set – Tapered Reamers: 100 mm



1	15-8711/02	Instrument Tray, empty, 478 x 253 x 76 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	



15-8720/02 Instrument Set – Tapered Reamers: 130 mm



1	15-8721/02	Instrument Tray, empty, 478 x 253 x 76 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				

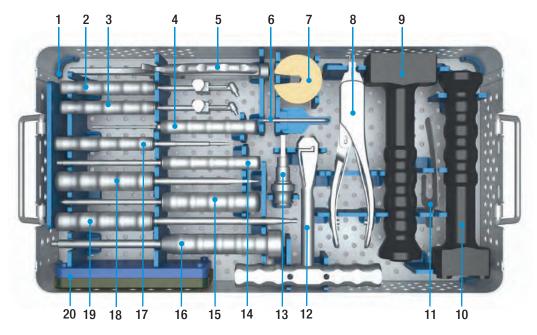


15-8730/02 Instrument Set – Tapered Reamers: 160 mm



1	15-8731/02	Instrument Tray, empty, 478 x 253 x 76 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				





15-8740/02 Instrument Set – General Instruments I

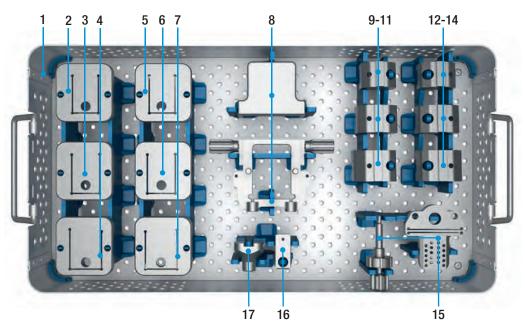
1	15-8741/02	Instrument Tray, empty, 478 x 253 x 76 mm						
2	15-8035/02	Insertion Instrument for PE-Plateaus Endo-Model S, M, L						
3	15-8035/03	nsertion Instrument for PE-Plateau Endo-Model XS + Endo-Model SL						
4	15-2546*	lex Screwdriver, torque limiting, hex 2.0 mm, 175 mm (for implant version V02)						
5	317-586	Extraction Forceps for fixation pins, 210 mm						
6	16-0116/01	Hex Screwdriver, wrench size 2.5 mm						
7	16-3203/00	Impaction Plate for tapered reamers						
8	16-0020/01**	Connecting/Disconnecting Forceps, 175 mm						
9	317-646/01	Grooved Driver for femoral components, 210 mm						
10	16-0018/02	Grooved Driver für tibial components SL						
11	317-607/50	Cutting Template						
12	15-6053/00	T-Handle, with Hudson fitting						
13	16-3283/01	Adapter with fitting: Hudson female/AO male						
		Fittings optional (not included in Instrument Set):						
	16-3284/00	Hudson female/Jacobs male						
	16-3285/00	Hudson female/Harris male						
14	16-0017/01	Separate Rod M5, 220 mm						
15	317-658/01	Bone Awl, with trocar point, 215 mm						
16	15-8516/45	Driver Extractor, for modular stems, 365 mm						
17	10-5373/01	Hex Screwdriver, wrench size 2.5 mm, 180 mm						
18	322-145/01	Screwdriver, blade width 8 mm, 210 mm						
19	64-8008/02	Hex Screwdriver, wrench size 3.5 mm, 250 mm						
	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)						
	319-602/30	Sterilizing Box						

Optional:

- * 64-1181/16 Hex Screwdriver, wrench size 2.0 mm, 175 mm (for implant version V01)
- ** 16-0020/02 Connection/Disconnecting Forceps, slim, 175 mm



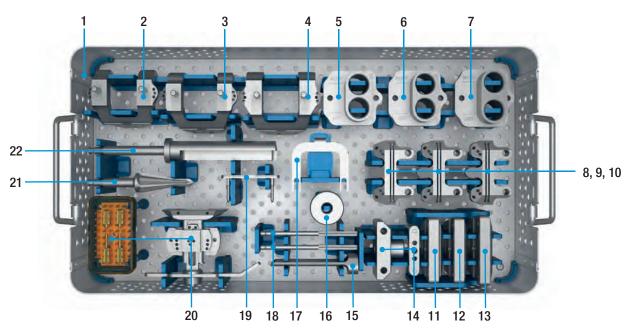
15-8760/02 Instrument Set – Femur I



1	15-8761/02	Instrument Tray	Instrument Tray, empty, 478 x 253 x 76 mm					
		Saw Attachmen	ts for femoral components SL					
2	16-3221/01	small right						
3	16-3223/01	medium M	right					
4	16-3225/01	large L	ge L right					
5	16-3221/02	small S	small S left					
6	16-3223/02	medium M	medium M left					
7	16-3225/02	large L	large L left					
8	16-3278/00	Revision Alignn	nent Gauge, distal, for femoral components SL (2 parts)					
		Femoral Trial B	ox for femoral components SL					
9	16-3261/01	small S	right					
10	16-3263/01	medium M	right					
11	16-3265/01	large L	right					
12	16-3261/02	small S	left					
13	16-3263/02	medium M	left					
14	16-3265/02	large L	left					
15	16-3277/00	Revision Cuttin	Revision Cutting Block, distal, for femoral components SL (2 parts)					
16	16-3282/00	Aligment Instru	ment for femoral trial box					
17	16-3271/00	Adapter for fem	oral trial box					



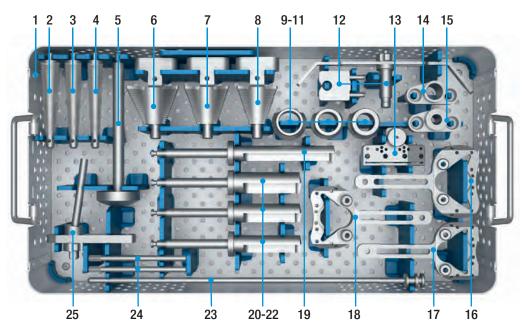
15-8770/02 Instrument Set – Femur II



1	15-8771/02	Instrume	nt Tray, empty, 478 x 253 x 76 mm					
		Condyle (Caps Femur					
2	16-3241/02	size 2	size 2 right/left					
3	16-3241/03	size 3	size 3 right/left					
4	16-3241/04	size 4	right/left					
		Drill Caps	Femur, Ø 20 mm					
5	16-3213/02	size 2	right/left					
6	16-3213/03	size 3	right/left					
7	16-3213/04	size 4	right/left					
		Femur Cu	tting Blocks for distal cut					
8	16-3228/02	size 2						
9	16-3228/03	size 3	size 3					
10	16-3228/04	size 4						
		Femur Cu	Femur Cutting Blocks for chamfer cuts					
11	16-3250/02	size 2						
12	16-3250/03	size 3						
13	16-3250/04	size 4	size 4					
14	16-3275/00	Alignmen	t Instrument for valgus angulation (2 parts)					
15	317-802/32	Chisel for	patella glide, 80 mm					
16	16-3281/00	Center SI	eeve for drill cap femur					
17	16-3279/00	Holding C	Clamp					
18	317-802/58	Alignmen	Alignment Rod, for epicondyles, 100 mm (2 parts)					
19	317-802/36	Dove Tail	Dove Tail Adapter, neutral, for femoral cutting blocks					
20	16-3276/00	-	t Instrument, for determination of external rotation (5 parts)					
	319-601/30	-	Box incl. screws, for Alignment Instrument 16-3276/00					
21	15-6037/00		cal, to open femoral and tibial cavity, with Hudson fitting					
22	16-3206/20	Drill with	stop, with Hudson fitting, Ø 20 mm					

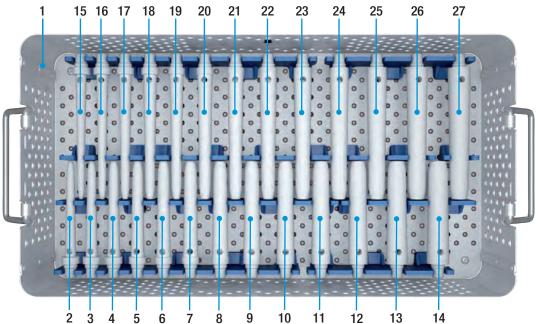


15-8780/02 Instrument Set - Tibia



1	15-8781/02	Instrument Tray, empty, 478 x 253 x 76 mm					
		Stem Compressors for tibial components SL					
2	16-3201/04	large L					
3	16-3201/03	medium M					
4	16-3201/02	small S					
5	16-3197/00	Handle, for tibial compressor/femoral trial box					
		Compressors for tibial components SL					
6	16-3199/14	large L					
7	16-3199/13	medium M					
8	16-3199/12	small S					
		Drill Guides for drill templates					
9	16-3270/22	large L					
10	16-3270/20	medium M					
11	16-3270/18	small S					
12	317-802/52	Stylus for tibial saw guide (2 parts), adjustable					
13	16-3241/00	Tibial Saw Guide for tibial components SL					
14	16-3267/00	Drill Guide, tibia, for ventral cut, Ø 16 mm					
15	16-3266/00	Alignment Gauge, tibia, for drill templates					
		Drill Templates for tibial components SL					
16	16-3198/14	large L					
17	16-3198/12	medium M					
18	16-3198/13	small S					
19	16-3207/16	Drill with stop, for tibial components SL, ventral side, Ø 16 mm					
		Drill with stop, for tibial components SL					
20	16-3208/22	large L Ø 22 mm					
21	16-3208/20	medium M Ø 20 mm					
22	16-3208/18	small S Ø 18 mm					
23	16-3242/00	Alignment Rod Tibia					
24	16-3211/00	Guide Rod, tibia, for drill template (2 ea. included)					
25	16-3212/08	Connector, size 8°, for tapered reamer/tibial saw guide					





15-8790/02 Instrument Set (1) – Femur Trial Stems 100 and 130 mm

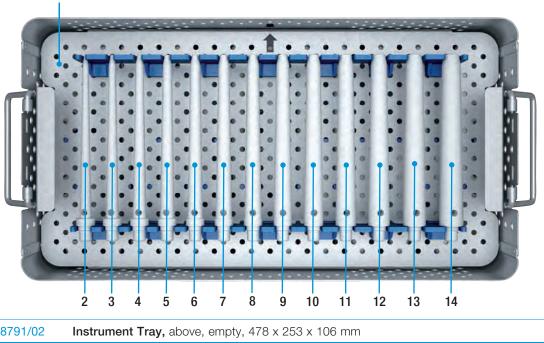
		234507	8 9 10 11 12	13 14					
1	15-8791/02	1/02 Instrument Tray, below, empty, 478 x 253 x 106 mm							
		Trial Stems, conical, for p	prosthesis stems 100 mm						
2	16-3101/12	for: stem-Ø 12 mm	stem length 100 mm						
3	16-3101/13	for: stem-Ø 13 mm	stem length 100 mm						
4	16-3101/14*	for: stem-Ø 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, conical, for p	prosthesis stems 130 mm						
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 27	16-3131/23	for: stem-Ø 23 mm	stem length 130 mm						
21	16-3131/24	for: stem-Ø 24 mm	stem length 130 mm						

* also for cemented stems 12, 14 und 16 mm

1



15-8790/02 Instrument Set (2) – Femur Trial Stems 160 mm

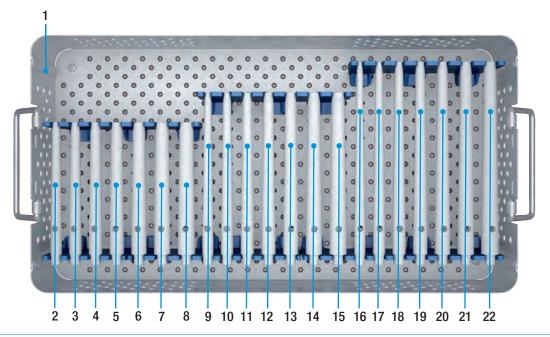


1	15-8791/02	Instrument Tray, above, empty, 478 x 253 x 106 mm						
		Trial Stems, conical, for prosthesis stems 160 mm						
2	16-3161/12	for: stem-Ø 12 mm	stem length 160 mm					
3	16-3161/13	for: stem-Ø 13 mm	stem length 160 mm					
4	16-3161/14*	for: stem-Ø 14 mm	stem length 160 mm					
5	16-3161/15	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



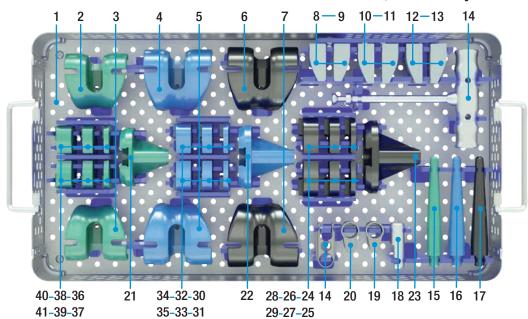
15-8810/02 Instrument Set – Tibial Trial Stems 100, 130 and 160 mm



1	15-8811/02	Instrument Tray, empty,	Instrument Tray, empty, 478 x 253 x 76 mm				
		Trial Stems, conical, for n	nodular tibia: 100 mm				
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, conical, for n	nodular tibia: 130 mm				
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, conical, for n	nodular tibia: 160 mm				
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



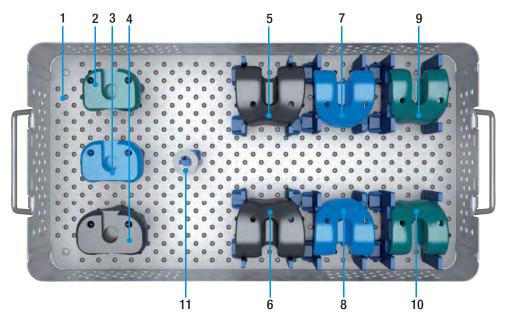


15-8820/03 Instrument Set -Trial Prostheses: Femur/Tibia, intracondylar

1	15-8821/03	Instrum	ent Tray, empty, 478	x 253 x 76 mm		
		Femur T	rial Prostheses, intra	acondylar		
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			
		Femur T	rial Segments			
8	16-4368/12	left	S/M/L	15 mm high		
9	16-4368/11	right	S/M/L	15 mm high		
10	16-4368/02	left	S/M/L	25 mm high		
11	16-4368/01	right	S/M/L	25 mm high		
12	16-4368/22	left	S/M/L	full		
13	16-4368/21	right	S/M/L	full		
14	15-6061/00	Extracti	on Instrument for tria	Il prostheses (2 parts)		
15	16-3170/12	Tibial Tr	rial Stems, small S			
16	16-3170/13	Tibial Tr	rial Stems, medium M	1		
17	16-3170/14	Tibial Tr	rial Stems, large L			
18	16-4367/20		s for mobile and hing			
19	16-4367/00		nnection Componen			
20	16-4367/10	Trial Co	nnection Componen	t for rotational knee		
21	16-3175/12	Tibial Tr	ial Prostheses, intrac	condylar, small S		
22	16-3175/13	Tibial Tr	ial Prostheses, intrac	condylar, medium M		
23	16-3175/14	Tibial Tr	ial Prostheses, intrac	condylar, large L		
		Tibial Tr	ial Spacers			
24	16-4361/03	right	large L	5 mm high	black	
25	16-4362/03	left	large L	5 mm high	black	
26	16-4363/03	right	large L	10 mm high	black	
27	16-4364/03	left	large L	10 mm high	black	
28	16-4365/03	right	large L	15 mm high	black	
29	16-4366/03	left	large L	15 mm high	black	
30	16-4361/02	right	medium M	5 mm high	blue	
31	16-4362/02	left	medium M	5 mm high	blue	
32	16-4363/02	right	medium M	10 mm high	blue	
33 34	16-4364/02	left	medium M	10 mm high	blue	
34	16-4365/02 16-4366/02	right left	medium M medium M	15 mm high 15 mm high	blue blue	
35 36	16-4361/01	right	small S	5 mm high	green	
30	16-4362/01	left	small S	5 mm high	green	
38	16-4363/01	right	small S	10 mm high	green	
39	16-4364/01	left	small S	10 mm high	green	
40	16-4365/01	right	small S	15 mm high	green	
41	16-4366/01	left	small S	15 mm high	green	
					3.00	



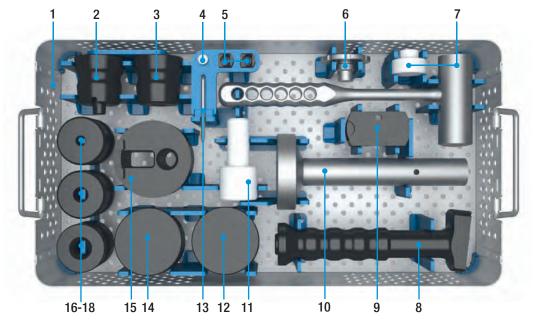
15-8840/03 Instrument Set – Trial Instruments: Distal Femur and Proximal Tibia Replacement



1	15-8841/02	Instrument ⁻	Instrument Tray, empty, 478 x 253 x 76 mm						
		Trial Prosth	esis for proxir	nal tibia r	eplacement				
2	16-3180/12	small S	green						
3	16-3180/13	medium M	blue						
4	16-3180/14	large L	black						
		Trial Prostheses for distal femur replacement, slim, X-ray opaque							
5	16-3195/21	large L	rig	jht	black				
6	16-3195/22	large L	lef	ť	black				
7	16-3193/21	medium M	rig	jht	blue				
8	16-3193/22	medium M	lef	ť	blue				
9	16-3191/21	small S	rig	jht	green				
10	16-3191/22	small S	lef	ť	green				
11	15-6094/00	Trial Suppo	rt Ring , Ø 28	mm, Hei	ght 10 mm				

	Optional: X-	Optional: X-ray opaque Trial Prostheses					
	Femoral Tria	al Prosthe	eses, intra	acondylar, X-ray	ay opaque		
16-3181/21	small S		right	green			
16-3181/22	small S		left	green			
16-3183/21	medium M		right	blue			
16-3183/22	medium M		left	blue			
16-3185/21	large L		right	black			
16-3185/22	large L		left	black			
	Tibial Trial F	Prosthese	s, intraco	ndylar, X-ray op	ppaque		
16-3175/22	small S	green					
16-3175/23	medium M	blue					
16-3175/24	large L	black					
	Trial Prosth	eses for p	roximal til	bia replacement	nt, X-ray opaque		
16-3180/22	small S	green					
16-3180/23	medium M	blue					
16-3180/24	large L	black					





16-0100/02 Instrument Set – Assembling Instrument Set

1	16-0100/03	Instrument Tray, empty, 478 x 253 x 86 mm			
2	16-0111/02	Assembling Protector for male taper			
3	16-0111/01	Assembling Protector for female taper			
4	16-0114/04	Screw for assembling protectors femur			
5	16-0114/02	Block, spare part for femur assembling protector 16-0114/01 (2 ea. included)			
6	16-0113/01	Metal Core for assembling protector, for modular stem (size 1 – 3)			
7	16-0115/01	Mallet, non-resilient, 800g, complete			
8	16-0114/01	Handle for femur assembling protector			
9	16-0114/03	Plate, spare part for femur assembling protector 16-0114/01			
10	16-0118/01	Assembling Table: upper part			
11	16-0118/03	Assembling Table: silencer			
12	16-0110/02	Tibia Assembling Plate für Endo-Model SL knee prostheses			
13	16-0116/02	Hex Screwdriver, wrench size 5 mm, conical			
14	16-0110/01	Femur Assembling Plate			
15	16-0110/03	Tibia Assembling Plate für Endo-Model knee prostheses			
16	16-0112/01	Assembling Protectors, size 1			
17	16-0112/02	Assembling Protectors, size 2			
18	16-0112/03	Assembling Protectors, size 3			
	16-0118/02*	Assembling Table: lower part*			

* Not included in container 16-0100/02



Assembling Instruments

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



16-0118/01

Assembling Table: Upper Part



16-0118/02

AssemblingTable: Lower Part with fixed base plate



16-0118/03 Assembling Table: Silence

Instruments







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



A CONTRACT OF STREET

16-0111/02

Assembling Protector for male taper



Assembling Protectors16-0112/01for modular stem, size 116-0112/02for modular stem, size 216-0112/03for modular stem, size 3



16-0113/01

Metal Core for assembling protector, for modular stem (size 1-3)

Instruments





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).



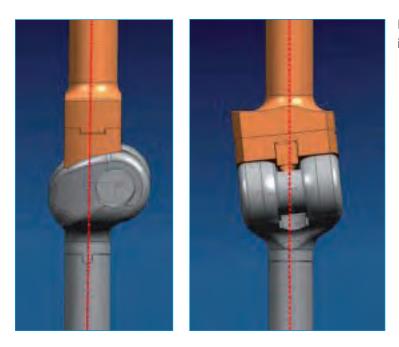
Assemble the elements with two blows of the nonresilient Mallet, using the metal face of the Mallet only.



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.

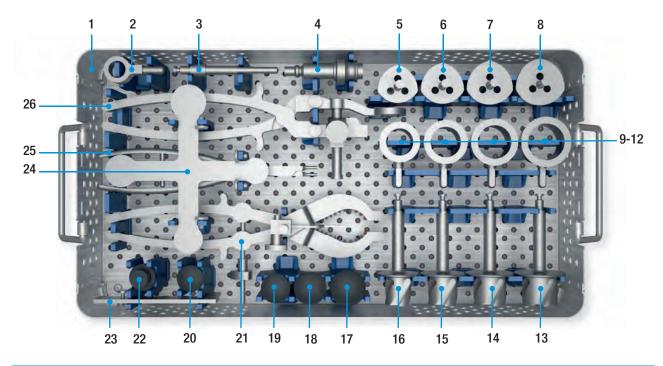
Assembly

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 choosen. Consecutively the block for Assembling Protector is secured using the Hex Screwdriver. The two arrows (of bock 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 Pusł	ner 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 Rese	ection Clamp		
22	340-009	Patella Push	ner Inserter		
23	65-2000	Townley Fe	mur 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 Hold	ling Clamp		

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

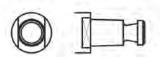


Additional Instruments

Adapter for Snap Lock Chuck

Various adapters to enable compatibility with other equipment connections.

REF	Fitting		
16-3283/01	Jacobs Fitting (E)		
16-3284/00	AO Fitting (D)	Charles .	
16-3285/00	Harris Fitting (C)		



Hudson fitting Basic tool connection for patella instruments



Sawblades,

without offset teeth, 1.24 mm thick

REF Wide (A) 25 mm	REF 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



H

Revision Drill Guide

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

REF	complete		
16-3295/00	consisting of 4 parts: bracket, drill rod, 2 rods with handle		

Extractor

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

REF	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)

REF	Length
317-661	365 mm

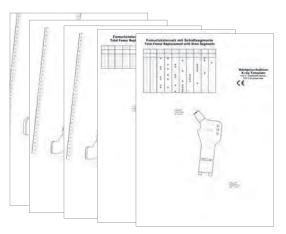
Hex Screwdriver,

hex 2.5 mm, 90°

REF	Length
16-1230/00	139 mm

O 2,5mm

Accessories



X-ray Templates,

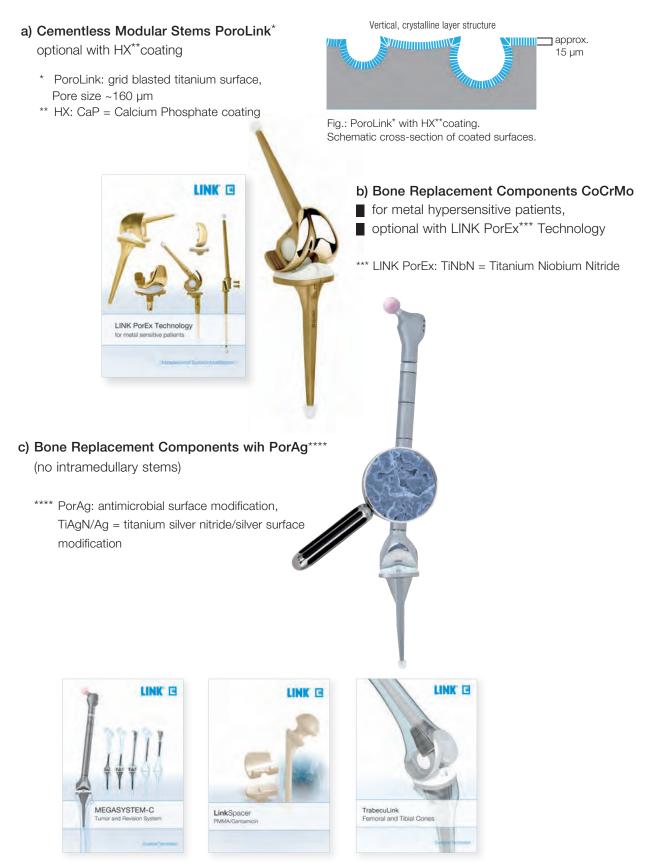
110% of actual size, set = 22 sheets

REF	
15-8516/62	for LINK MEGASYSTEM-C with SL knee components

Additional Information



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





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



Endo-Model SL Rotational and Hinge Knee System	Product	Rotational version	Hinged version
General Indications			
Mobility limiting diseases, fractures or defects of the knee joint, distal femur or proximal tibia which cannot be treated by conservative or osteosynthetic procedures		Х	Х
Indications			
Primary and secondary osteoarthritis		Х	Х
Rheumatoid arthritis		Х	Х
Revision after primary or revision total knee replacement		Х	Х
Bone necroses which won't compromise the successful implantation of a hinged total knee endoprosthesis		Х	X
Varus and valgus deformity with contracture or laxity of the medial or lateral stabilizers		Х	X
Extreme cases of varus/valgus deformities (20-30°), rheumatoid arthritis, muscular deficiency and any kind of genu laxum		-	X
Oncological and revision surgery in lower limb (in conjunction with Megasystem-C)		Х	Х
Contraindications			
Acute or chronic infections, local and systemic insofar as they compromise the successful implantation of a hinged total knee endoprosthesis		х	X
Allergies to (implant) materials		Х	Х
Distinctive muscular, nerve, vascular or other diseases which put the affecte limb at risk	əd	Х	X
Insufficient bone integrity which prevents a stable anchorage of the prosthe	sis	Х	Х

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.



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

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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.

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