

Aesculap[®] e.motion[®]

Knee Endoprosthesis System
Manual operation technique



Aesculap Orthopaedics



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Preoperative planning

The e.motion knee system provides radiographic templates which help the surgeon in determining the following parameters:

- Angle between anatomic and mechanical femur axes
- Resection height of the intact tibia joint surface
- Entry point of the intramedullary alignment rod
- Implant sizes
- Position of the osteophytes

The following x-ray images are required to conduct the radiographic analysis:

- Knee joint in AP projection: knee extended, centred over the distal patella.
- Knee joint in lateral projection: knee in 30° flexion, centred above the distal patella.
- Image of the whole leg in supported monopodal stance.
- Patella-tangential image: knee in 30° flexion, caudocranial radiation, centred over the distal patella.

The e.motion x-ray templates must be used.

The angle between the mechanical and anatomic femur axes is measured with the combination template for axis measurements. The center of the joint, the joint line and the mechanical femur axis can be measured and drawn onto the ray slide. To determine the tibia resection, the template showing representations of tibial implants is superimposed over, and aligned with, the x-ray image. The resection height is given at a 10-20 mm (FP, CS) and 10-24 mm (PS) graduation. A complete set of radiographic templates is provided for the preoperative determination of the appropriate implant sizes. The localization of the osteophytes facilitates their removal, improving the mobility of the joint.

The results of preoperative planning should be documented in the patient's file.



Positioning the tibia cutting block

The extramedullary alignment system for the tibia cutting block is brought into position parallel to the tibia axis. The rotational alignment is carried out with the extension of the malleolary clamp, which is oriented to the second metatarsal bone.

The alignment instrument offers the possibility of adjusting the tibia cutting block in all planes:

- Height adjustment **A**
- Alignment in the sagittal plane **B**
- Varus-valgus alignment **C**

The surgeon can freely define the adjustment of the cutting block, according to the individual patient's requirements. However, we recommend a perpendicular position in relation to the mechanical tibia axis, both in the frontal and in the sagittal plane.

A Height adjustment

The resection height is determined in preoperative planning. The aim is to remove any defect on the tibial joint surface as completely as possible in order to create a bed for the tibia plateau on intact bone. Following this, the measured value is set on the stylus, which is then introduced into the cutting slot. The extramedullary alignment instrument is then lowered by pulling the lever **1** until the stylus comes into contact with a point corresponding to the joint line.

B Alignment of the sagittal plane

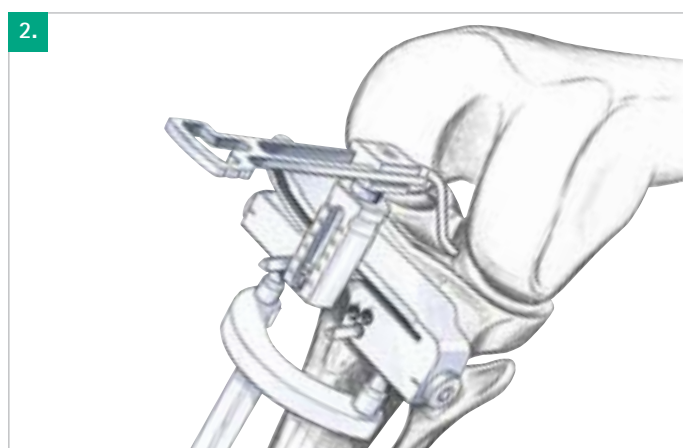
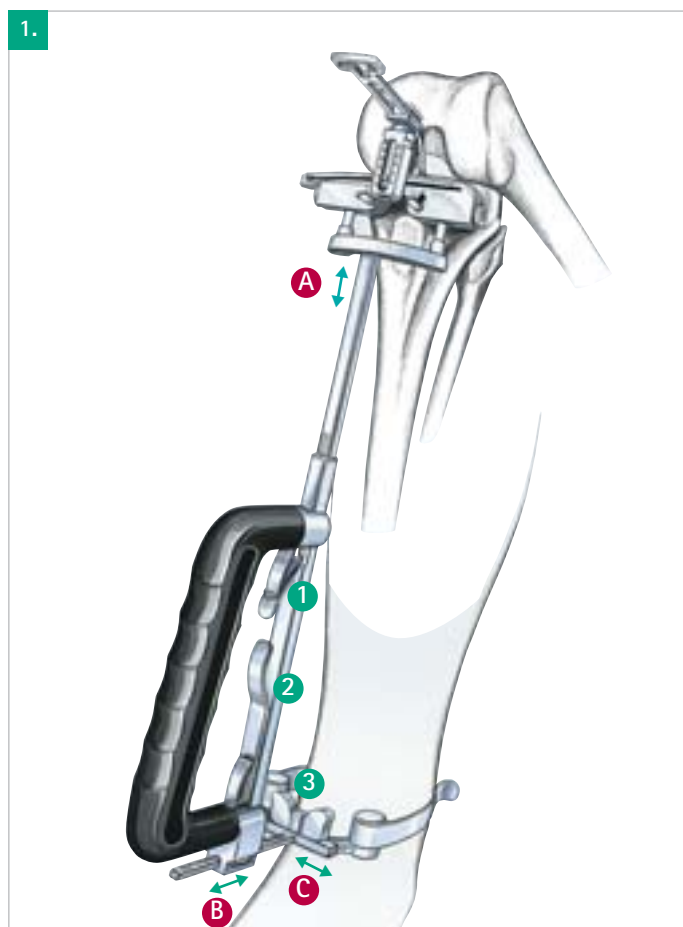
The alignment of the sagittal plane (parallel to the mechanical axis) is achieved by pulling the lever. **2** The distance between the lines on the malleolary clamp corresponds to a posterior slope of 1° for a tibia length of 40 cm.

■ Please note:

The e.motion tibial implant has
a 3° posterior slope in the plateau

C Varus-valgus alignment

Lifting the lever, **3** the slide in the malleolary clamp can be shifted towards mediolateral. The distance between the lines on the scale corresponds to a 1° alteration for a tibia length of 40 cm.



Resecting the tibia plateau

The cutting block is fixed to the bone with four threaded pins as follows. Two headless threaded pins are introduced into the drill holes marked "0". Two other threaded pins with head are then inserted in the convergent holes to secure the cutting block against movement during resection.

After removing the extramedullary alignment instrument, the resection is performed with a 1.27 mm thick saw blade. Extreme care must be taken at this step, since the posterior cruciate ligament must not be damaged in case of implantation of e.motion FP components. The resection is normally performed at a slope of 0°.

While the two convergent pins are removed, the two headless pins remain in place to allow a later corrective resection of 2 or 4 mm, if necessary.



Checking the tibial implant size

To ensure that compatible sizes are chosen for the implant components, the size of the tibial implant is determined, with the help of the tibia preparation plateaus, at this point of the operation. The correct implant size should offer a complete coverage of the resected plane and avoid any overhang. The femoral implant should be smaller, the same size or one size bigger than the tibial implant.

- Please note:
Size of Meniscal Components correspond to size of femoral component.

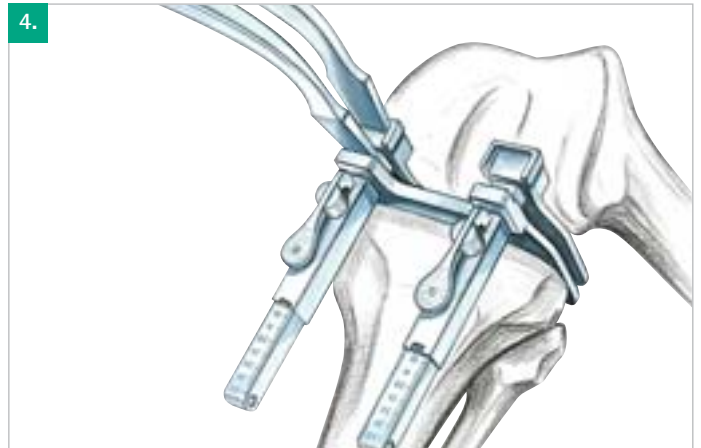
	F2	F3	F4	F5	F6	F7	F8
T1							
T2							
T3							
T4							
T5							
T6							
T7							
T8							

Combinations of implant sizes (tibia / femur) that are not possible

Possible combinations of implant sizes (tibia / femur)

Measuring the extension and flexion gaps

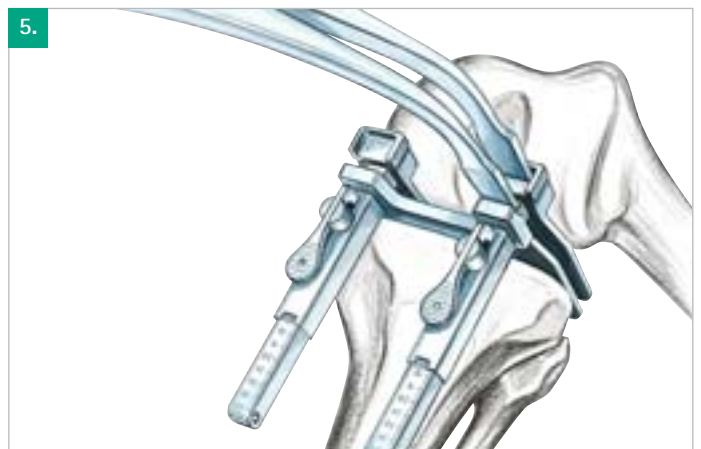
After resection of the tibia plateau, the ligament tension must be checked. The osteophytes on the tibia head and on the femoral condyles must be completely removed. The measurement makes it possible to calculate the resection height on the distal femur. The height to aim at is the thickness of the chosen implant size on the intact condyle.



The size of the flexion and extension gaps are read medially and laterally at the distractor in 90° flexion and 0° extension. The number read is the one on the movable shoe level with the end of the sleeve.

In cases of a medio-lateral asymmetry (of more than 3 mm), ligament release should be performed on the narrower side (medial for varus deformities, lateral for valgus deformities). Following the ligament release the flexion and extension gaps should be re-measured, and the release procedure repeated if necessary. A mediolateral difference of 2 mm is acceptable.

■ **Please note:**
External rotation of the femoral component influences the medio-lateral gaps in flexion.



The size of the flexion and extension gaps are read medially and laterally. Different solutions are possible to compensate a difference between extension and flexion gap. The extension gap can be adjusted to the flexion gap through changing the distal femoral resection height by up to + or -2 mm. The flexion gap can be adjusted to the extension gap by choosing a smaller or larger femoral implant (preferable as the joint line is preserved).

Further possibilities exist in building up the defective distal femoral condyle e.g. with bone.



Distal femur resection

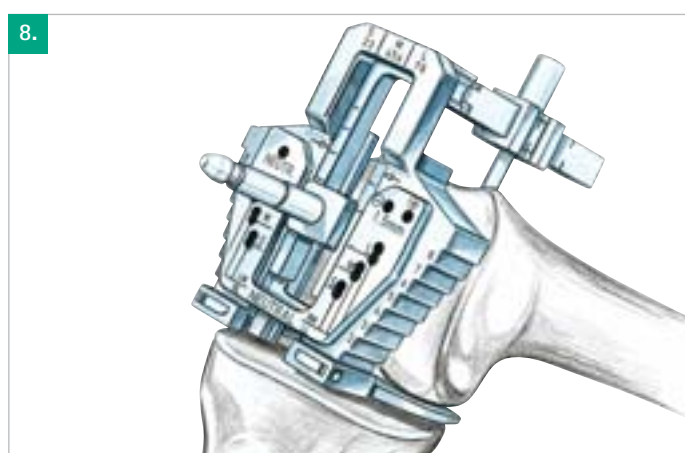
The entry point into the femoral medullary cavity is prepared using a broach, according to preoperative planning.

The medullary canal is opened up with a Ø 9 mm drill. The Ø 8 mm intramedullary femur rod has been specially designed for minimizing the risk of embolisms. The rod is carefully introduced into the medullary cavity using the T-shaped handle.



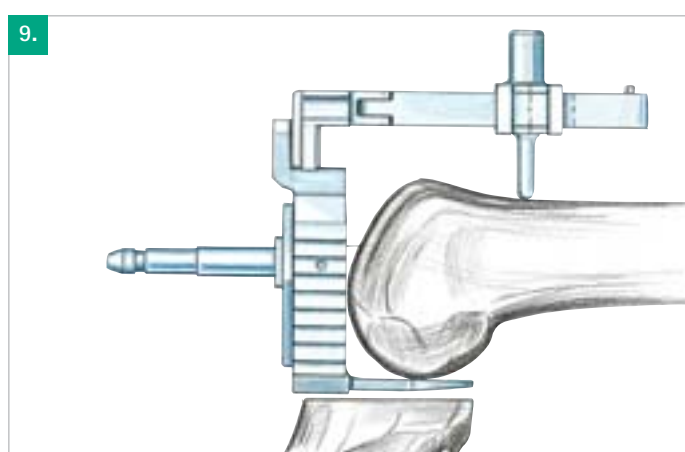
To adjust the varus / valgus angle into the femur orientation block, the appropriate angle block (5°, 6°, 7°, 8° or 9°) is introduced in the block according to preoperative planning. For a left leg, the side marked with an "L" must be the upper side; for a right leg, the side marked "R" must be on top.

The femur stylus is introduced into the block from the upper side; the mounted instrument is then fixed to the femur by means of the intramedullary rod. The posterior condylar plates of the orientation block must touch the posterior condyles and at least one distal condyle.



The femur stylus must touch the anterior cortex. The point palpated at the antero-lateral cortex defines the anterior resection height and the position of the end of the femur shield. Using the graduation on the stylus, the appropriate size of the femoral implant can be defined with reference to the anterior shield. The stylus scale only corresponds to the correct sizing before the distal resection has been performed.

A scale on the frontal side of the instrument shows the respective prosthesis size with reference to the AP dimension of the femur. The appropriate implant size with regard to the mediolateral width of the bone can be read from graduations on the outer ends of the femur orientation block.



Distal femur resection

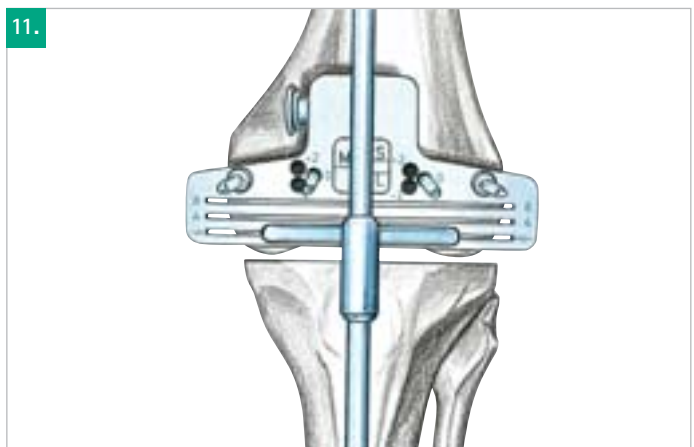
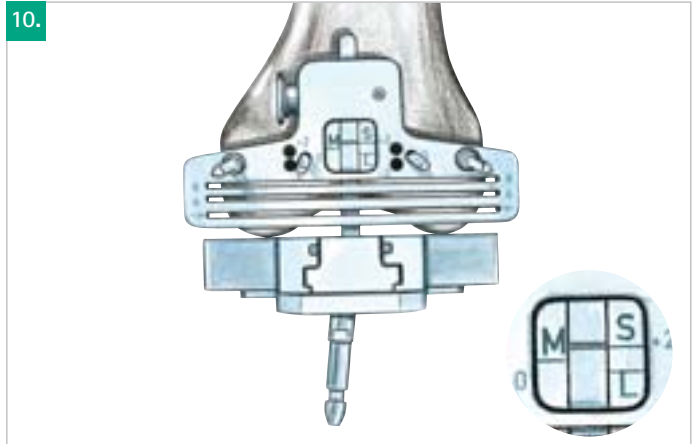
Having determined the appropriate femoral implant size, the anterior femur stylus is replaced by the holding device of the distal cutting block. The distal block is attached to the holding device in position S, M or L, depending on the implant size. (S = implant sizes 2 and 3; M = 4, 5 and 6; L = 7 and 8). This adjustment defines the thickness of the distal resection which is 7.0 mm for size group S, 8.5 mm for size group M and 10.0 mm for size group L.

The distal cutting block must touch at least one anterior condyle. The block is then fixed onto the bone by two headless threaded pins inserted into the holes marked "0". If a different resection becomes necessary, the holes marked "-2" and "+2" and the cutting slit marked "4" and "8" allow to adapt the thickness of the distal resection by shifting the cutting block.

The stability of the fixation is ensured by screwing in two pins with head in the convergent holes on both sides of the cutting block.

The holding system and the intra-medullary femur rod are removed, leaving only the cutting block fixed to the bone.

The axis can be checked by placing the measuring rod holder into the slit on the femoral cutting block. The measuring rod with the socket for the second measuring rod can then be inserted into the holder and the second measuring rod fixed into the socket. Before carrying out the distal resection with 1.27 mm saw blade in the cutting slit marked by arrows, the resection depth must be checked with a cutting check device. The tibia protection plate prevents damage to the tibia plateau.



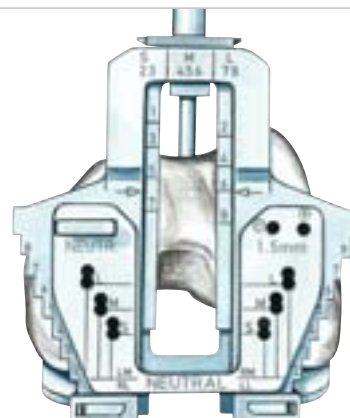
Determining the size of the femoral implant

After the distal resection, the femur orientation block with anterior stylus is placed on the distal resection surface, and the posterior condylar plates are brought into contact with the posterior condyles. The stylus must be positioned on the same point at the anterior cortex where it was prior to the distal resection.

The final size of the femoral implant is defined, according to the size readings of the AP and the mediolateral scale. If the result is a full size (e.g. 6), the fixation key must be positioned in the hole marked as "neutral". If it is an intermediate size (e.g. 5, 4), the fixation key may be positioned in one of the two holes marked "+" or "-" -1.5 mm in order to prevent anterior femoral undercutting or any protrusion of the cortex. If "+" is chosen, the position of the drill holes for the fixation pins for the APC sawing block is shifted towards anterior by 1.5 mm; if "-" is chosen, the same shift is applied towards posterior.

The box measure of the femoral implants increases by only 3 mm from one size to the next, resulting in a maximum compromise of 1.5 mm when choosing an intermediate size.

12.

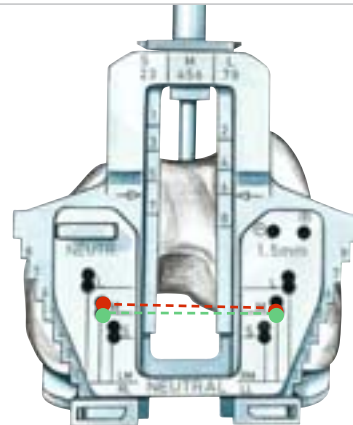


Rotational alignment of the femur component

The two holes for the fixation pins of the APC cutting block can be drilled with the 2 drills with stop Ø 3.2 mm (NE333R) parallel to the posterior condyles by choosing the lower holes (green points) for the respective size group (S, M or L).

To achieve an external rotation of 3°, the surgeon may choose the lower hole on the lateral side and the upper hole on the medial side (red points).

13.



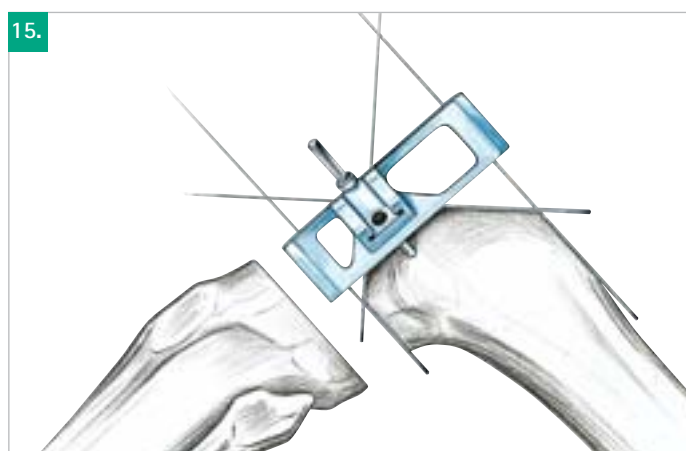
Completing the femur resection

The pegs of the APC cutting block fixed in the two drill holes prepared previously, according to the femoral implant size chosen. The "ANT" marking for the anterior resection on the APC cutting block is visible. After this, the APC sawing block is fixed to the femur with the two universal handles and with two headed pins in the convergent holes.

Care must be taken that the APC sawing block is in good contact with the distal resection surface.



The cutting depth check plate is used for verifying the positioning and depth of the cuts prior to resection. It is advisable to use the tibia protection plate to avoid damaging the tibia plateau. The four femur resections are carried out with a 1.27 mm saw blade inserted into the cutting slits. The chamfer cuts are performed after the anterior and posterior cuts.

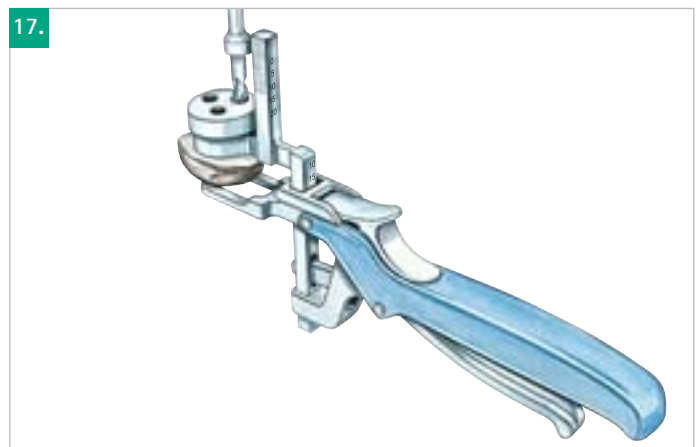
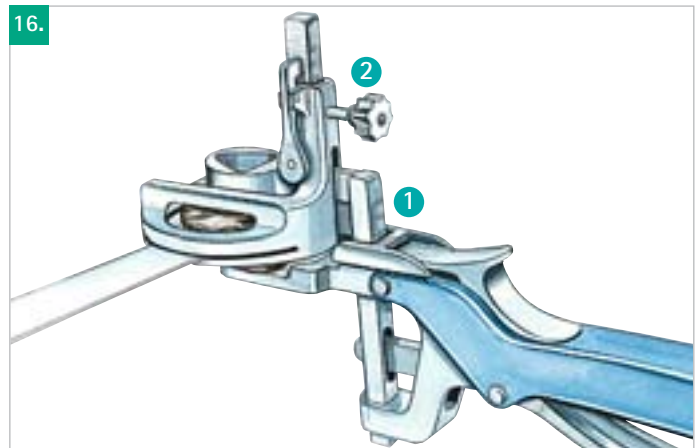


Patella preparation

- 1 With the patella forceps the thickness of the patella is measured.
- 2 After implanting the patella component, this thickness should not be exceeded. The chosen resection height can be adjusted at the forceps by means of a graduated scale.

The resection is performed through the cutting slit. The patella implant is chosen either with regard to the optimum bony coverage of the patella, or it is guided by the thickness of the resection performed. The original thickness should not be exceeded after implantation of the patella rear surface.

After resection the saw attachment is removed and the triple drilled sleeve is attached to the forceps. The holes for the pegs of the patella implant are drilled. The patella size is determined by using the trial patella implants as position and size of the patella pegs are the same for all sizes.

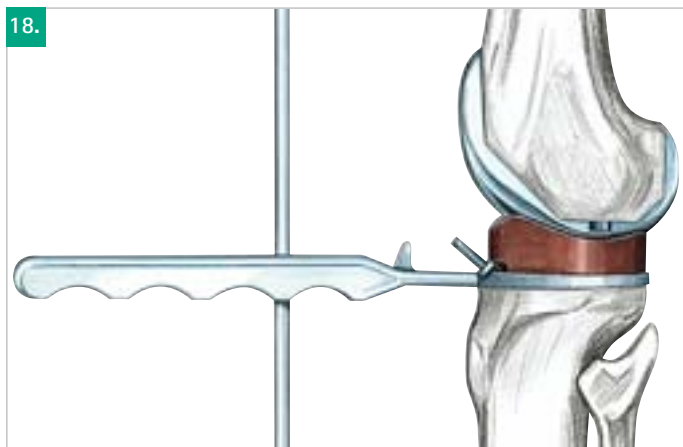


Trial reposition

Initially, the tibia preparation plateau is coupled to the universal handle and positioned on the resection plane. After a meniscus component with size reference to the femur component is fixed on the tibia preparation plateau. Finally the femur implant is inserted and aligned mediolaterally.

Following this procedure it is advisable to test the entire joint function with the patella in its anatomical position or with a trial patella implant.

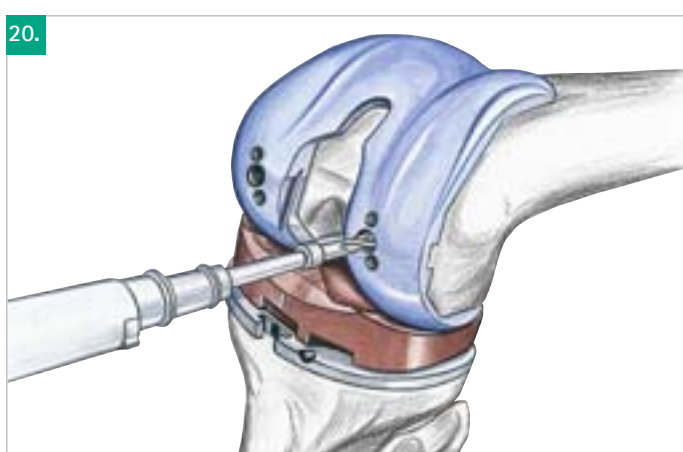
Alignment should be checked in flexion and extension by again inserting the extramedullary measuring rods into the handle attached to the tibial plateau. The position of the measuring rod is checked in relation to the midpoint of the femoral head and the ankle joint.



The rotational alignment of the tibia plateau is carried out with reference to the anterior marking. The latter should point to the medial third of the anterior tuberosity. The rotational alignment can also be carried out in a functional manner, with reference to the femur component, by moving the unfixed tibia plateau from extension to flexion. Internal rotation should be avoided under all circumstances.



Following the functional test of the joint, and the inspection of the medio-lateral position of the femoral trial implant, the peg holes of the final femoral implant can be drilled.



Tibia preparation

The tibia plateau is fixed in the desired position with short threaded pins with head. The correct size is chosen considering the best bony coverage of the resected plain. Additional stabilization is provided with the universal handle. The cylindrical drill guide with the desired diameter is positioned on the tibia preparation plateau. The holding clamp stabilizes the guide in position during the drilling process. The drill guides as well as the drills are available in diameter 10, 12, 14 and 16 mm.

Tibia preparation without stem extension

The selection of the correct diameter of drill and drill guide depends on the size of the final tibia implant and the diameter of its wing stem.

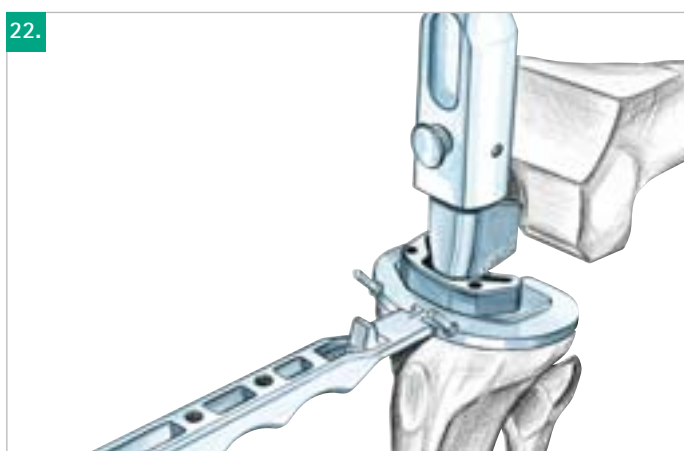
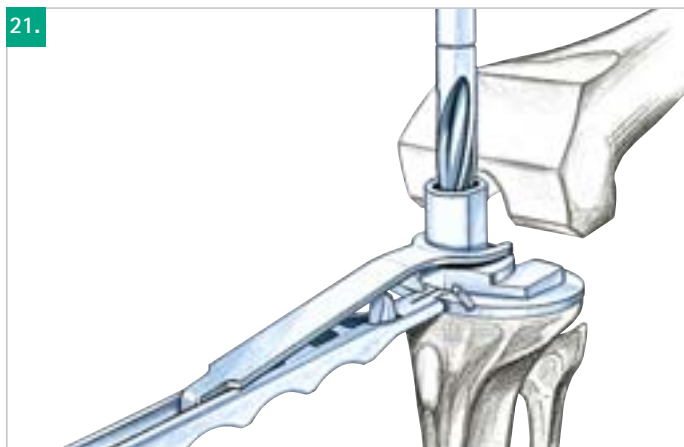
Size Group S	(T1, T2, T3)	wing stem diameter of 12 mm
Size Group M	(T3, T4, T5)	wing stem diameter of 14 mm
Size Group L	(T7, T8)	wing stem diameter of 16 mm

On the modular drills the first mark corresponds to the wing stem length. Please drill until the first mark is in line with the upper border of the drill guide.

Tibia preparation with stem extension

In case of a tibia preparation with the use of extension stems, the choice of the correct drill and drill guide diameter depends on the wing and the extension stem diameter. The e.motion product portfolio offers extension stems in 4 diameters (10, 12, 14 and 16 mm) and 3 different lengths (Short: 52 mm, Middle 92 mm und Long 132 mm).

For the preparation of the wing stem (until the first mark) the diameter of drill and drill guide should correspond to the implant stem size. In cases where the intramedullary canal diameter is smaller, a drill with a smaller diameter is chosen to prepare the canal for the extension stem implantation. The drills have 4 different marks showing the wing stem, small, middle and long stem length.

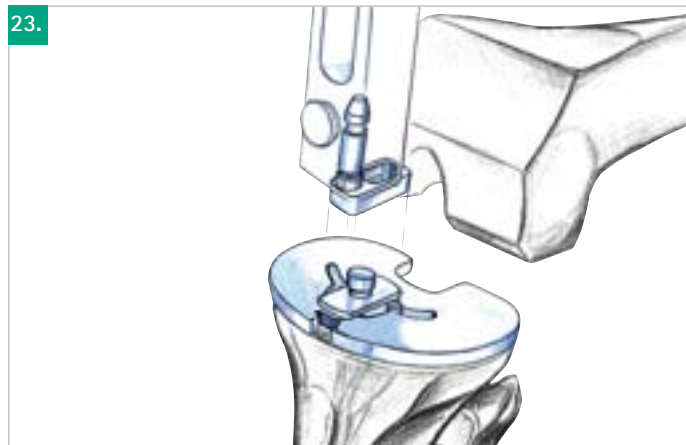


■ Please note:

There is full interchangeability of stems and obturators between the sizes with reference to the thread connection.

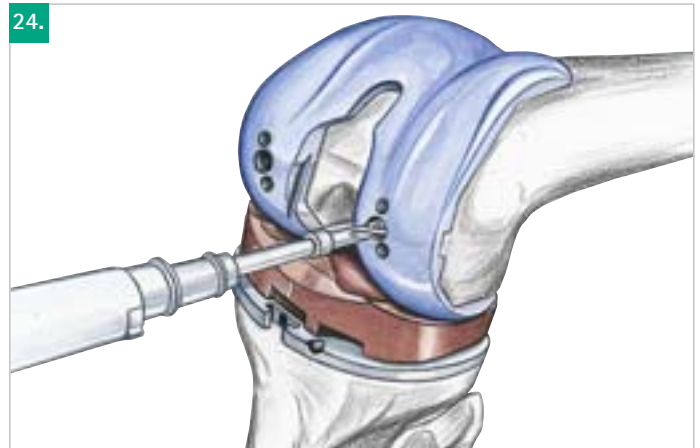
For the preparation of the wings the size of the correct chisel is chosen and the corresponding chisel guide is positioned on the tibia preparation plateau. The wing chisel are available for the size group S (1, 2, 3), size group M (4, 5, 6) and size group L (7, 8).

After the tibia preparation the modular trial implant can be inserted. The trial components consists of the trial plateau being suitable for left or right knees and the wing stem with the possibility to connect trial extension stem if needed. The modular trial implant is inserted with the help of an adapter that is fixed to the impaction handle.



Posterior stabilized PS version

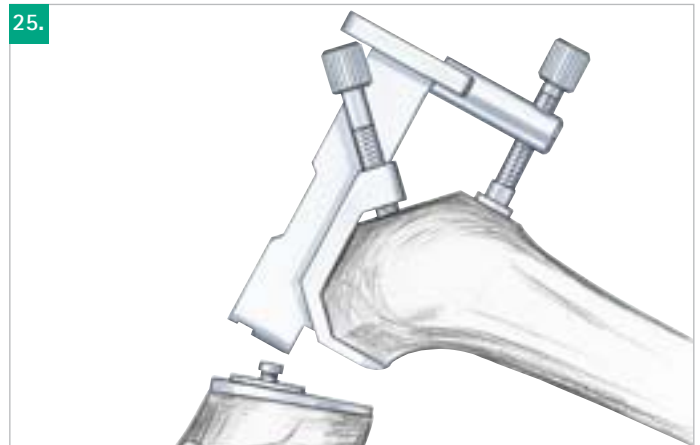
The holes drilled through the trial femur implant are used to fix the box preparation base to the femur. There is one box preparation base available for each femoral size group (S= 2, 3 M= 4, 5, 6 L = 7, 8). Before putting the base in place, the femur trial implant must be removed.



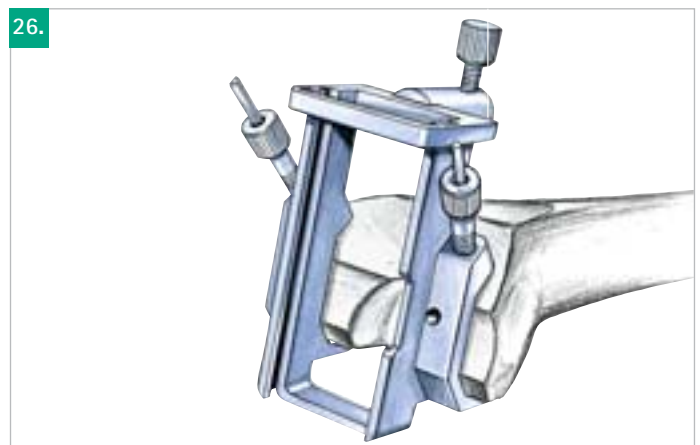
In case of the use of a posterior stabilized implant the medial-lateral orientation of the trial femur implant in the trial reposition step is decisive for the box preparation.

■ Please note:

For a proper alignment of the post-cam mechanism the final preparation of the tibia is recommended before the trial reposition and box preparation step.



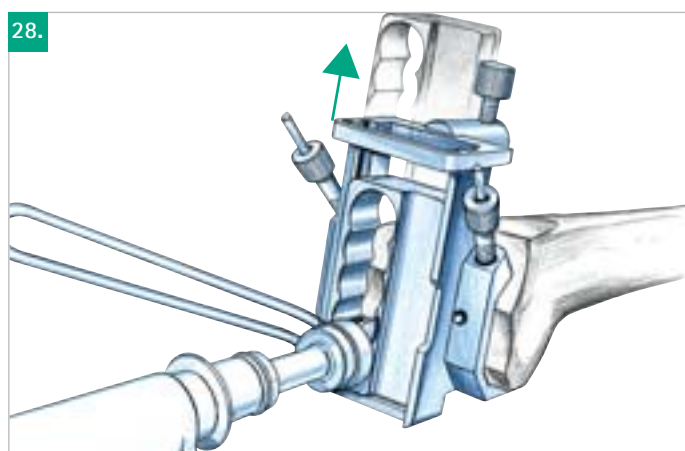
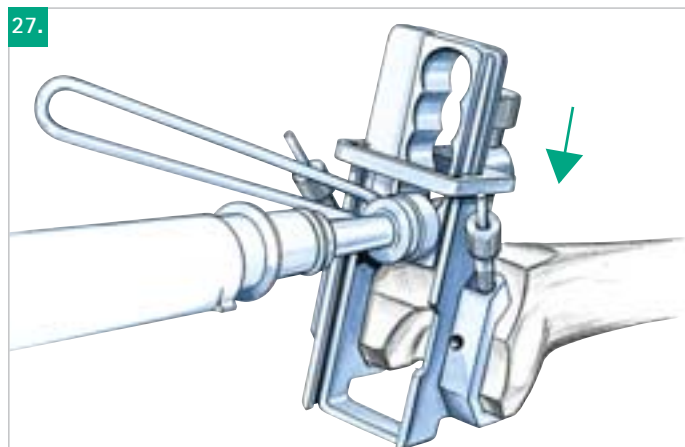
For fixation of the base to the femur the ventral clamping screw is fixed to the anterior cortex. Additional stability is obtained by introducing 2 convergent pins on the medial and lateral side of the base.



Milling templates are available for every femur size individually. The correct milling template is chosen and inserted into the preparation base. The milling procedure is done in two stages starting on the left stage. After performing the first stage, the milling template is removed, turned and inserted in the preparation base again. With the help of a guiding sleeve and the 15 mm diameter cutter the box preparation can be performed directly cutting through the bone. In the case of hard or sclerotic bone several holes can be drilling before guiding the cutter directly through the prepared stage to smoothen the borders.

■ Please note:

When preparing and smoothening the box, the cutter must be guided in a clockwise direction.



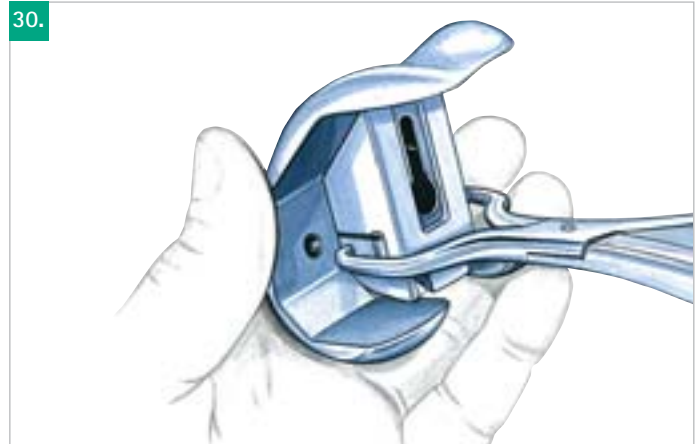
For verification of the proper intercondylar preparation of the femur, the appropriated box is connected to the trial femur implant with the help of a special forceps or by pressing it in the foreseen space by hand.



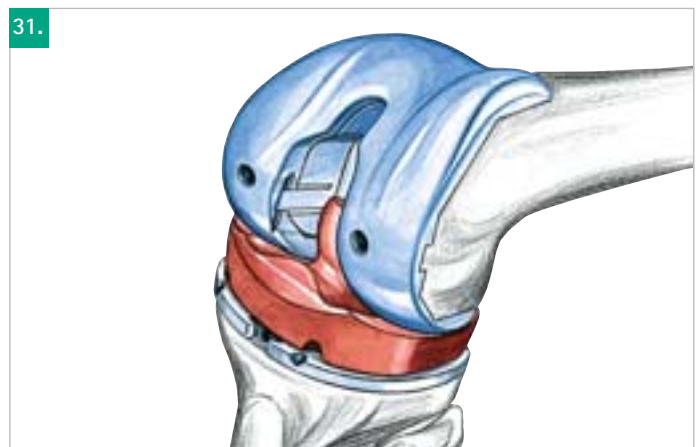
Posterior stabilized PS version

For the removal of the box from the femur trial implant the forceps is used.

After putting the trial femur and meniscal components implants in place, the quality of the preparation and the joint stability are checked in a functional test. The trial implants consists of the modular tibia implant (plateau, wing stem, extension stem if needed), the femur trial implant with connected box and the trial meniscal component in the desired height with post.



The trial PS meniscal components are available in 10, 12 and 14 mm height. 2 additional complementary plate of 6 mm each can be connected to the trial PS meniscal components to reach the remaining heights of the available final PS meniscal component range.



meniscal component	complementary plate	trial component
10	1 x 6	16
12	1 x 6	18
14	1 x 6	20
10	2 x 6	22
12	2 x 6	24

Final Implantation

The e.motion femoral and tibial implants can be implanted with or without cement as desired. The surgeon makes the decision according to the bone quality of the patient.

Because of the precision and congruence of the resection surfaces and the implants, only a small amount of cement should be used. This is particularly important in the posterior regions of the femur and tibia implants e.motion FP and PS as well around the box in the case of e.motion PS to prevent cement getting into the periarticular gap.

■ **Please note:**

In order to prevent third-body wear, any cement residues must be removed thoroughly.

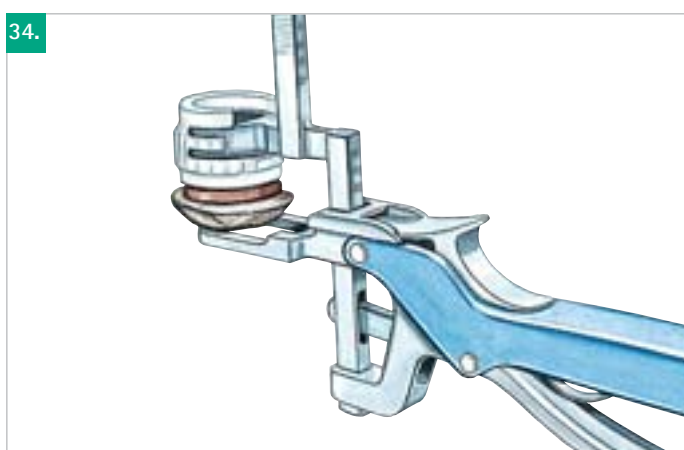
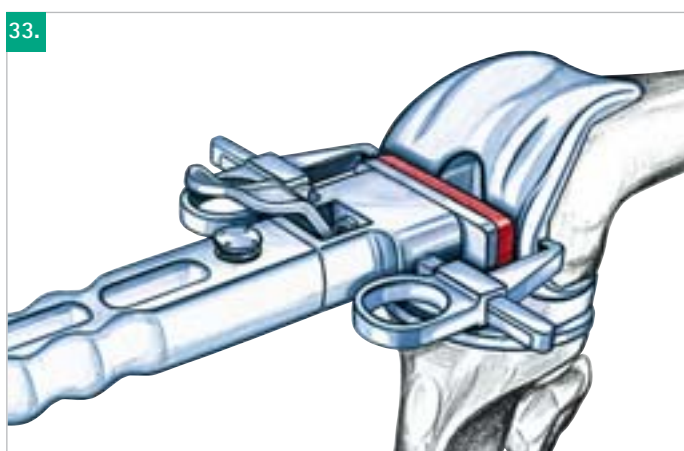
The following implant sequence is recommended in the case of e.motion FP:

- Tibia implant
- Meniscal component
- Femoral component
- Patella implant

The tibia final implant is connected to the holding device and brought precisely into the predefined position using the impaction handle. The final positioning is done with the tibia impactor. After the meniscal component is placed in position.

With the same holding device using the correct insert, the femur final implant is brought into alignment and implanted. The femoral impactor is used to knock the implant into place. In case of no stem application for cemented PS femurs a closing clip for the femur box is available by request to prevent the leakage of cement into the joint articulation area (Art. No.: NB700P).

The patella is implanted using the patella preparation forceps and the concave plastic cap, which allows good transmission of forces during the cement hardening process.



Final Implantation

The following implant sequence is recommended in the case of e.motion PS / CS:

- Tibia implant
- Femoral component
- Meniscal component
- Patella implant

The modular tibia implant e.motion PS is connected to a obturator screw or an extension stem if needed. Screw and stem are fixed to the wing stem of the final implant with the tightening key in the appropriated diameter (10, 12, 14 or 16 mm). Additionnally the bearing pin packed together with the meniscal component is screwed into the tibia plateau and fixed with a torque wrench applying 10 Nm. A scale at the instrument show the amount of force that is used.

In order to facilitate the handling, the tibia trial gliding surface mounted on the universal handle can be used as holding device for the final tibia implant assembling.

- **Please note:**
The bearing pin is fixed to the tibia plateau with a torque wrench applying 10 Nm.

After the femoral implant is put into place with the help of the holding device and then the meniscal component.

35.



36.



Overview trial implants

Femoral trial implant

37.

Adapter to the trial stems for use with the impaction handle

Trial wing stem for each size group available

"Click"-box for left and right

Trial gliding surface
e.motion FP H10, 12, 14, 16, 18, 20

or

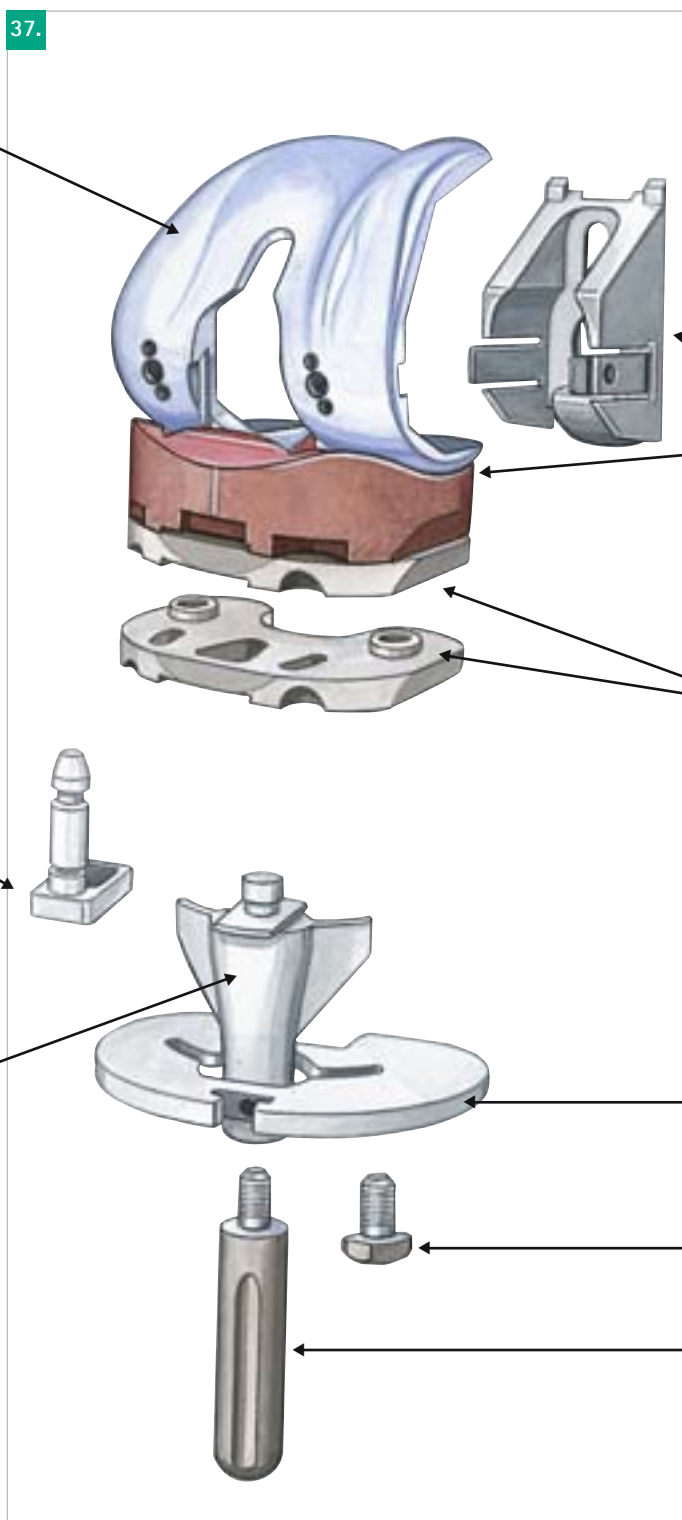
Trial gliding surface
e.motion PS H10, 12, 14, 16, 20, 24

Complementary plates (6 mm) to get every height available in the portfolio in combination with the gliding surfaces

Tibia preparation plateau

Tibia obturator screw

Tibia trial stem

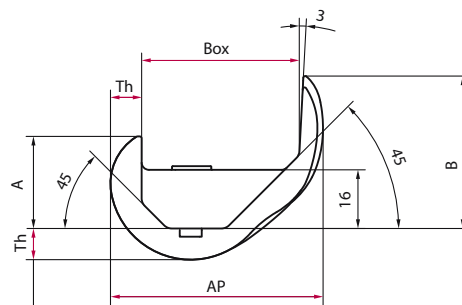
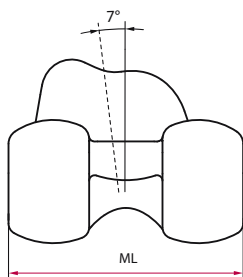


e.motion implant sizes

e.motion FP and PS implant sizes

Femoral component

The table gives an overview on the most important dimensions of the e.motion femoral implants

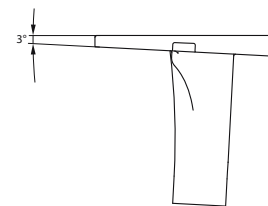
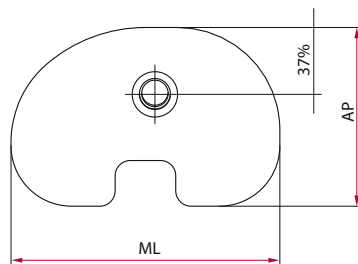
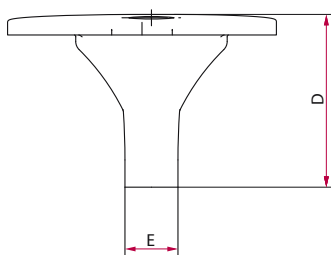


Measurements in (mm)

Size	ML	AP	Box	Th	Trochlear Depth
F2 L / R	56	49.9	37	7	4
F3 L / R	60	53.8	40	7	4.5
F4 L / R	64	58	43	8.5	4.5
F5 L / R	68	61.8	46	8.5	5
F6 L / R	72	65.6	49	8.5	5
F7 L / R	76	69.7	52	10	5.5
F8 L / R	80	73.8	55	10	6

Tibial component

The table gives an overview on the most important dimensions of the e.motion tibial implants



Measurements in (mm)

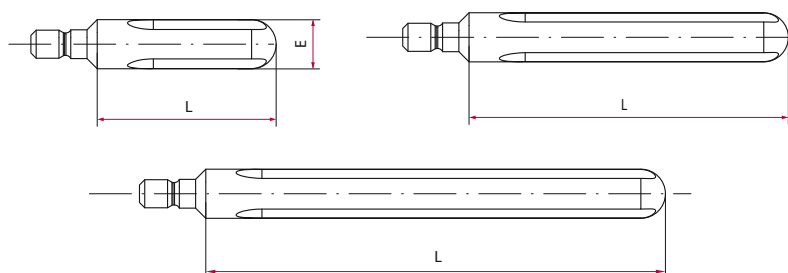
Size	ML	AP	AP / ML	D	E
T1 L / R	59	38	0.64	40	12
T2 L / R	63	41	0.65	40	12
T3 L / R	67	44	0.66	40	12
T4 L / R	71	47	0.66	45	14
T5 L / R	75	50	0.67	45	14
T6 L / R	79	53	0.67	45	14
T7 L / R	83	56	0.67	50	16
T8 L / R	87	59	0.68	50	16

Tibia extension stem

The table gives an overview on the most important dimensions of the e.motion tibia extension stems

Measurements in (mm)

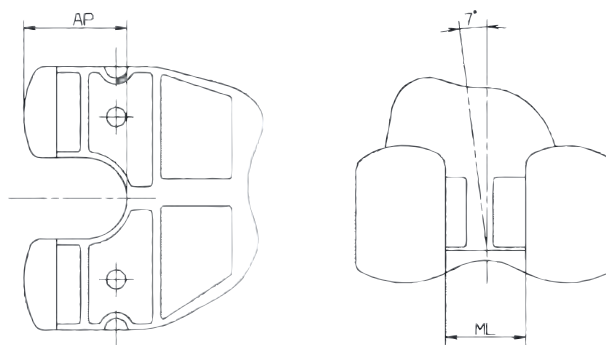
Size	L	D
Short	52	10, 12, 14, 16
Middle	92	10, 12, 14, 16
Long	132	10, 12, 14, 16



AP- / ML-Dimensions [mm] of the e.motion® femoral implants for necessary application of intra medullary nails

Measurements in (mm)

	AP	ML
F2	19.89	18
F3	22.23	19
F4	24.36	20
F5	26.64	21
F6	28.8	22
F7	31.05	23
F8	33.40	25

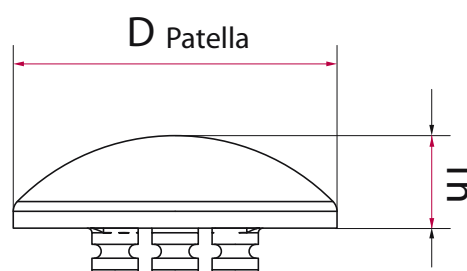


Patella component

The table gives an overview on the most important dimensions of the e.motion patella implants

Measurements in (mm)

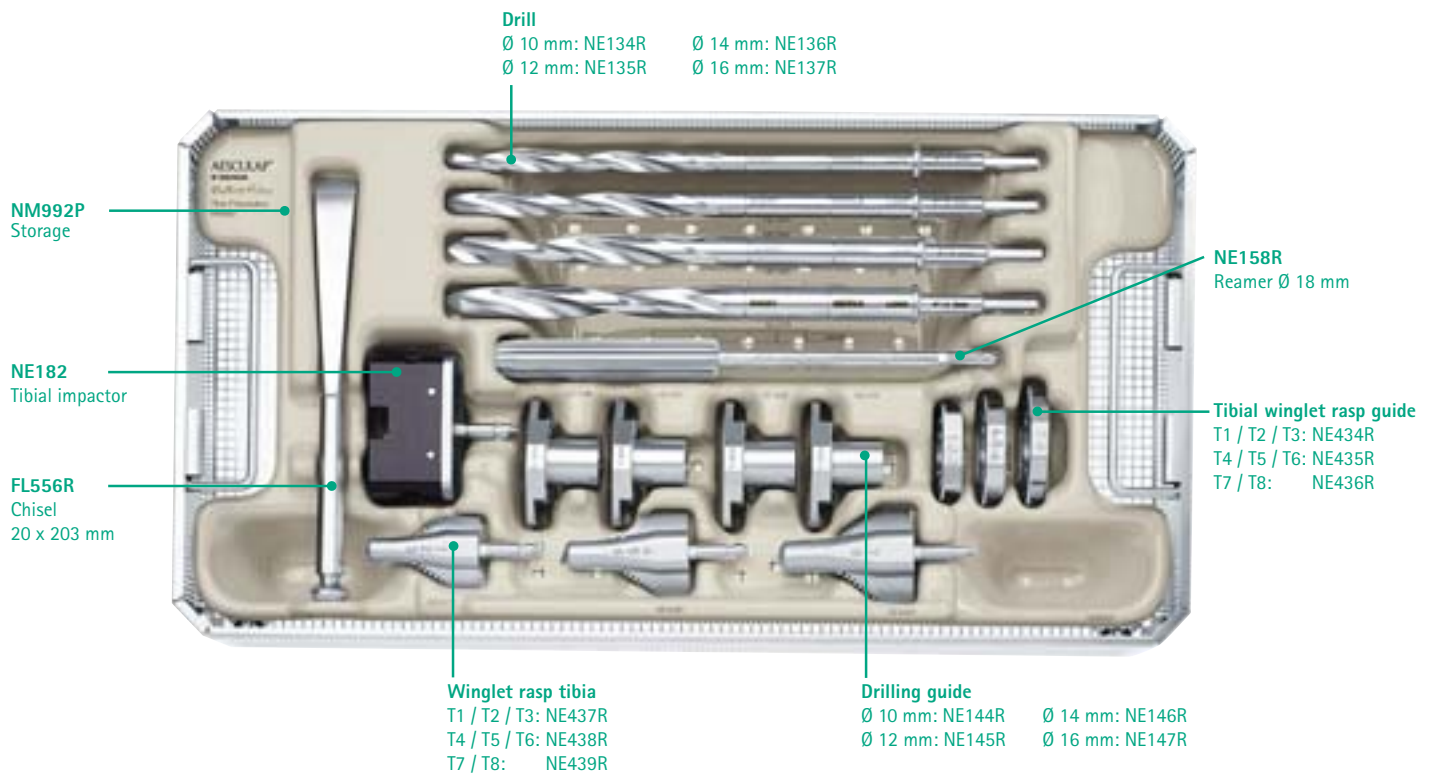
Size	D	Th
1	26	7
2	29	8
3	32	9
4	35	10
5	38	11
6	41	12



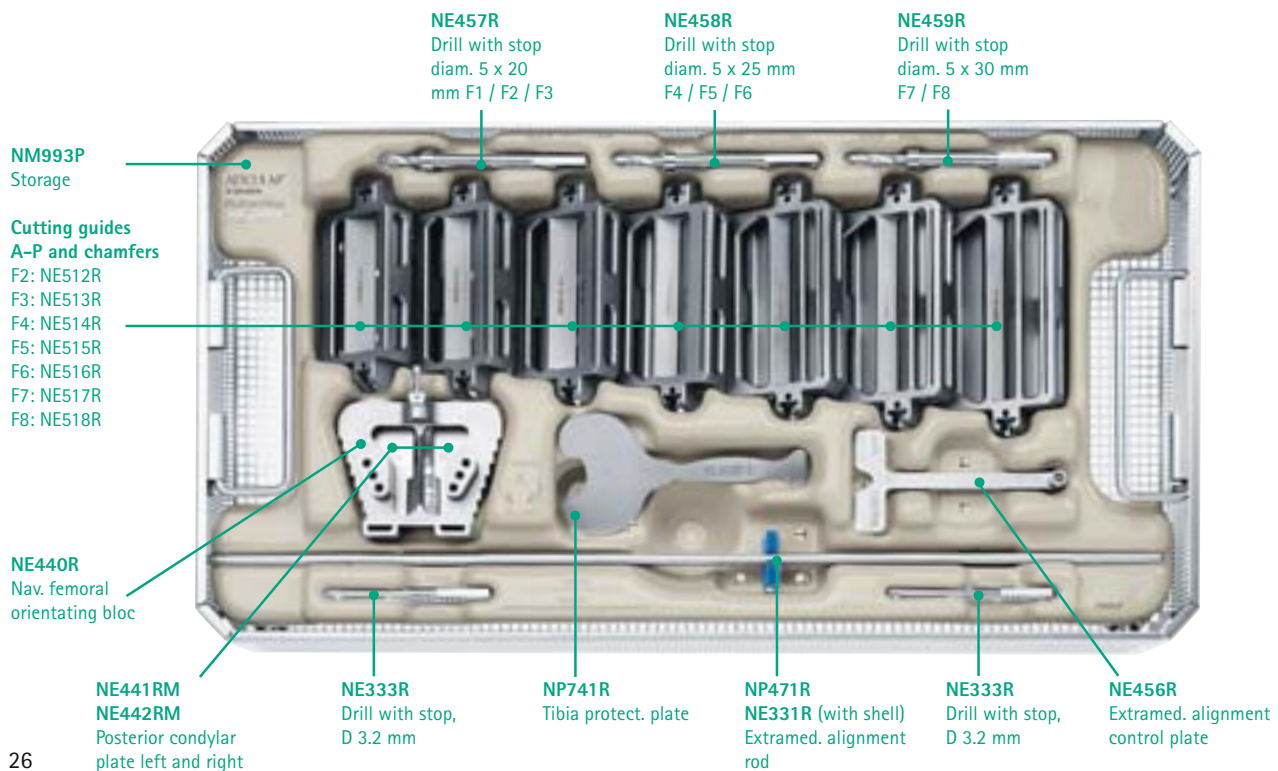
Instrumentarium

NE410 e.motion® FP

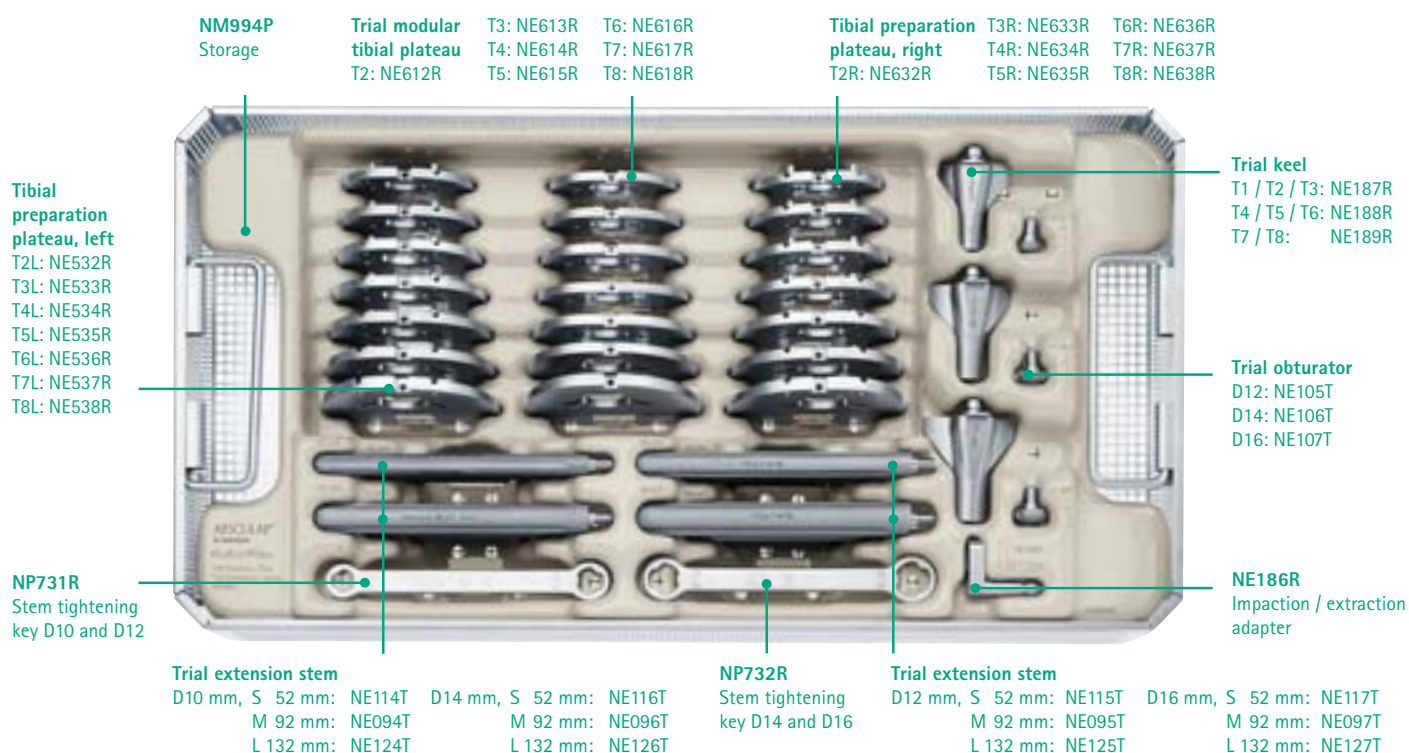
NM982 Tibia Preparation Instruments



NM983 Femur Preparation Instruments



NM984 Trial Tibia Components (left / right)

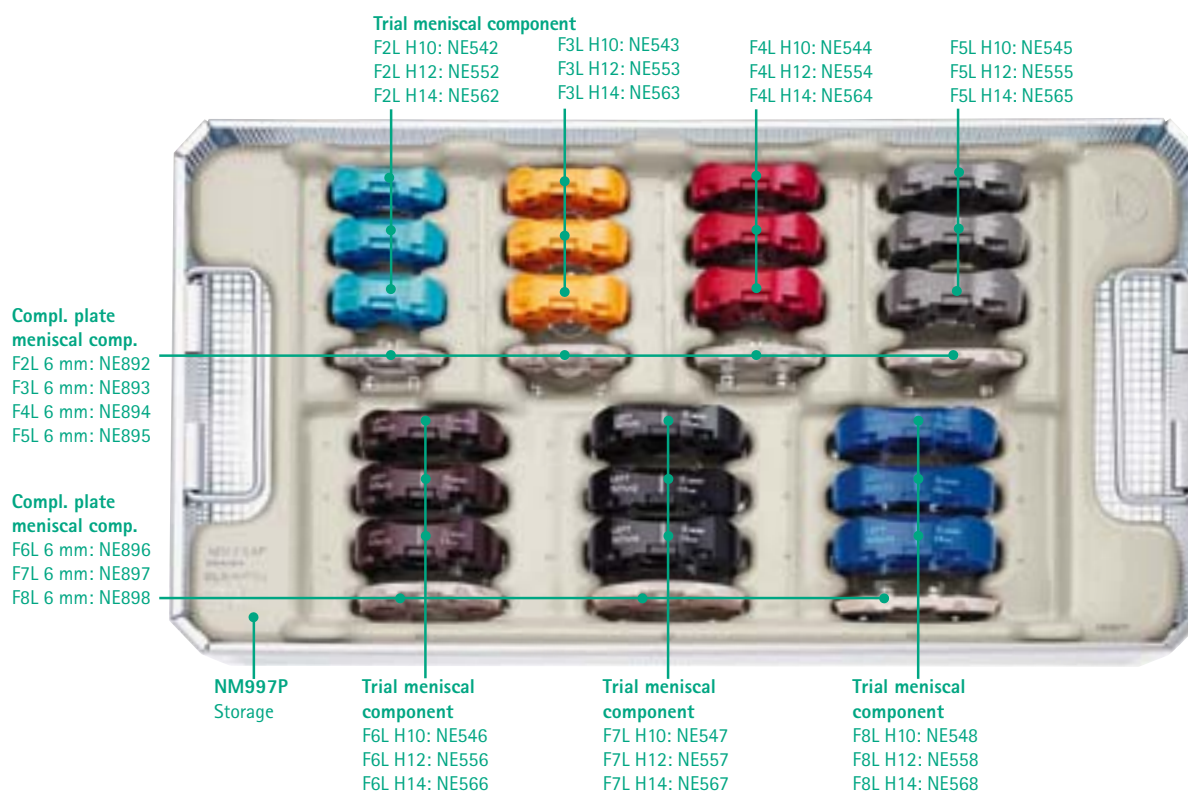


NM984 Trial Tibia Components (left / right)

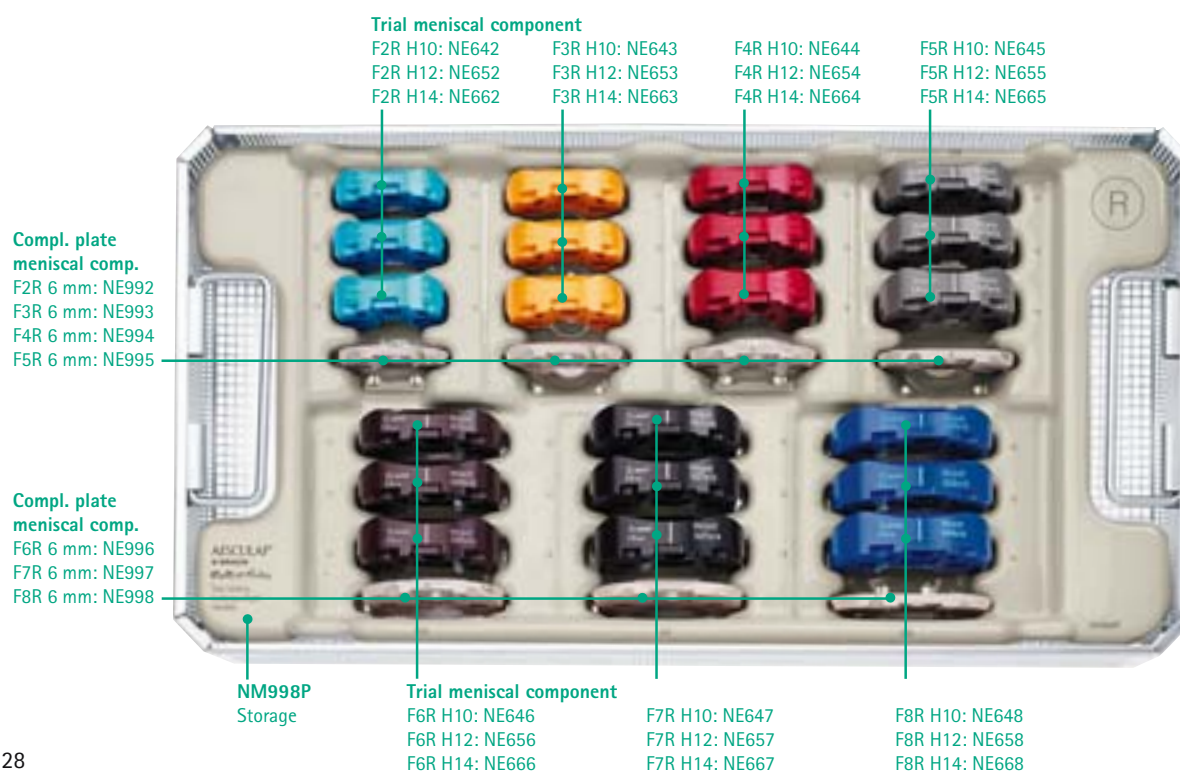


Instrumentarium

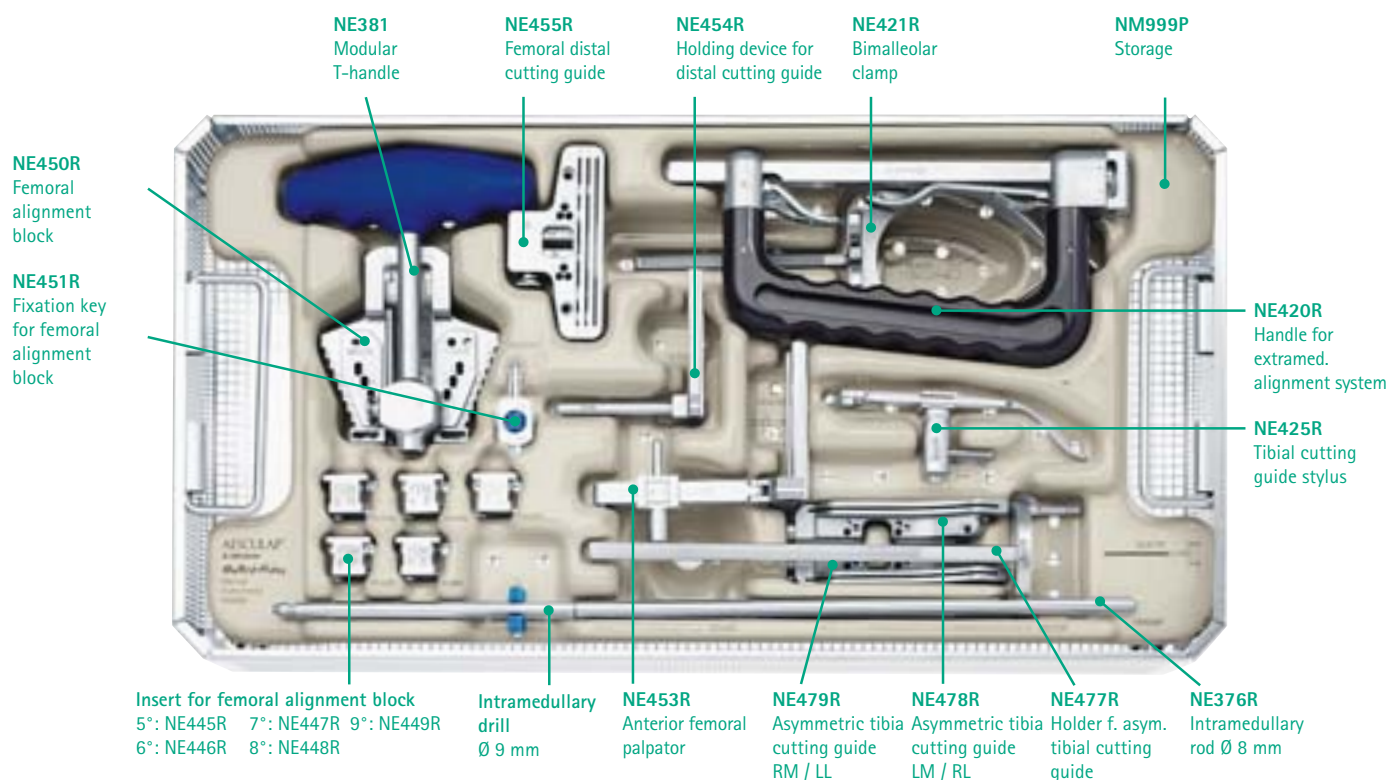
NM987 Trial Meniscal Components left



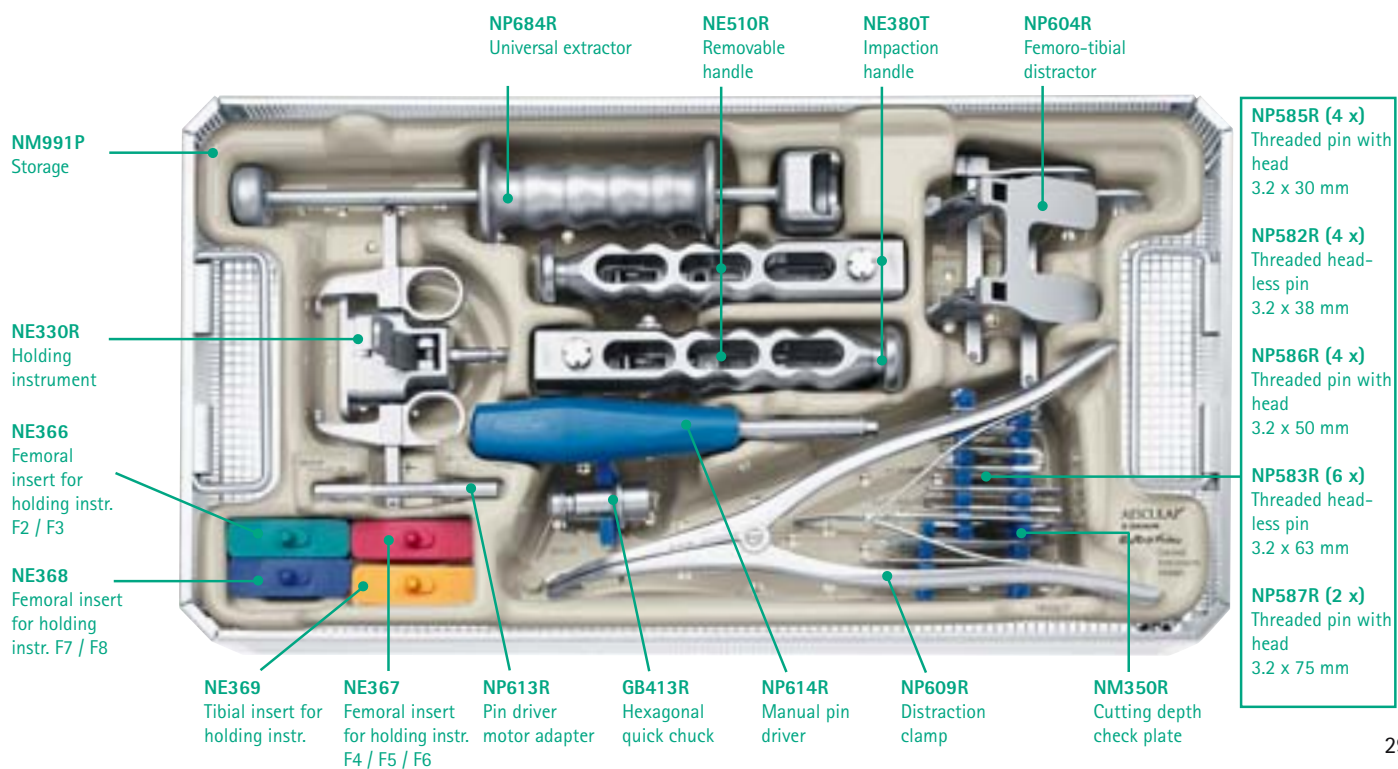
NM988 Trial Meniscal Components right



NM989 Manual Instruments



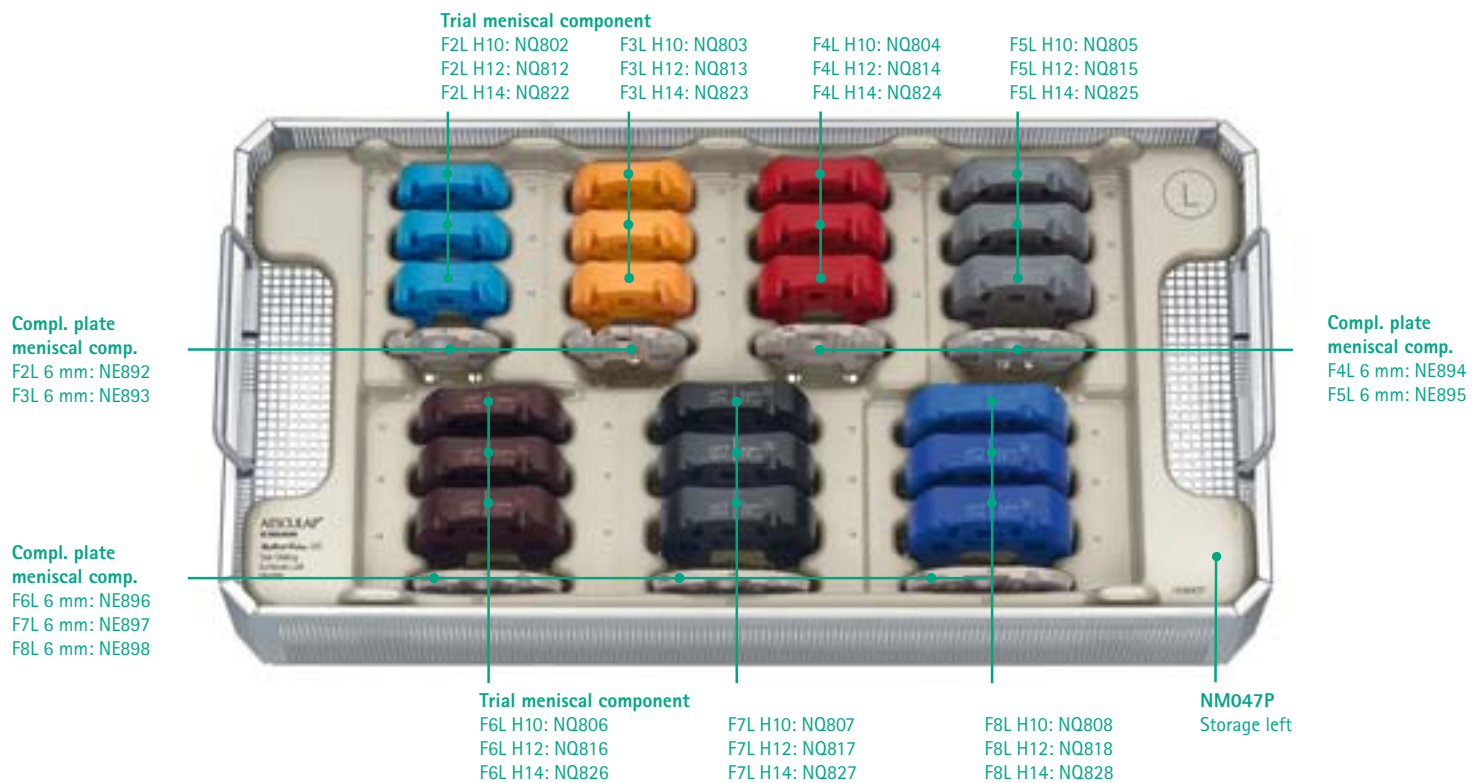
NM990 General Instruments



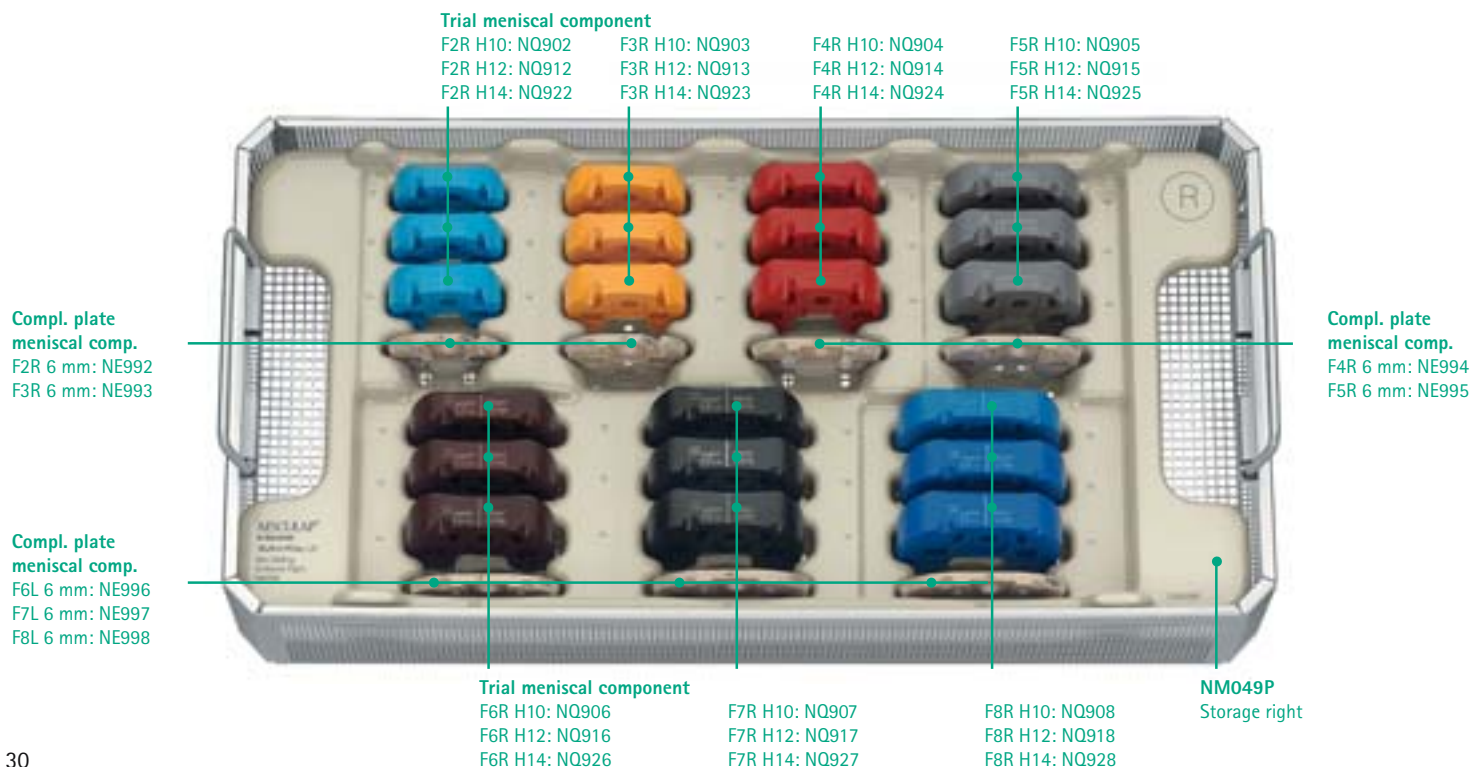
Instrumentarium

NM395 e.motion® UC

NM046 Trial Meniscal Components left

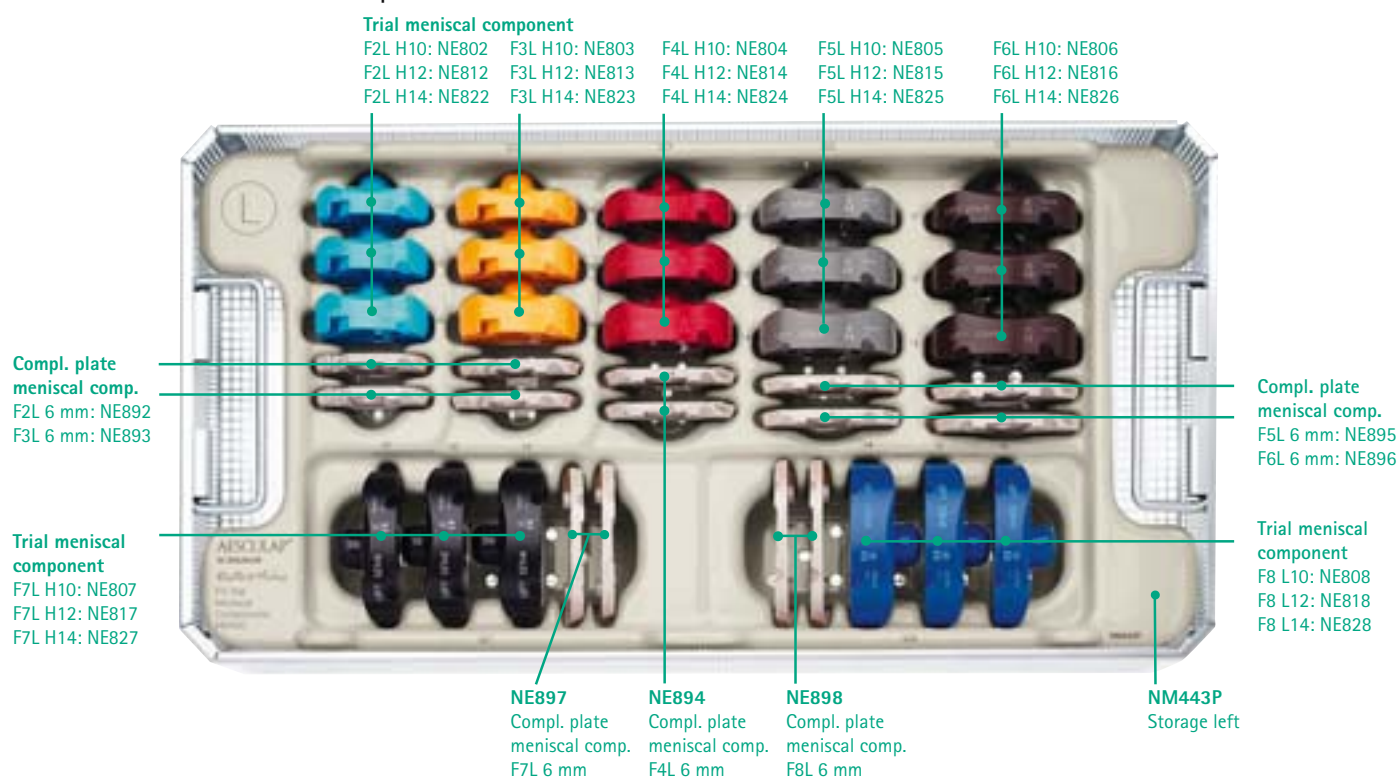


NM048 Trial Meniscal Components right

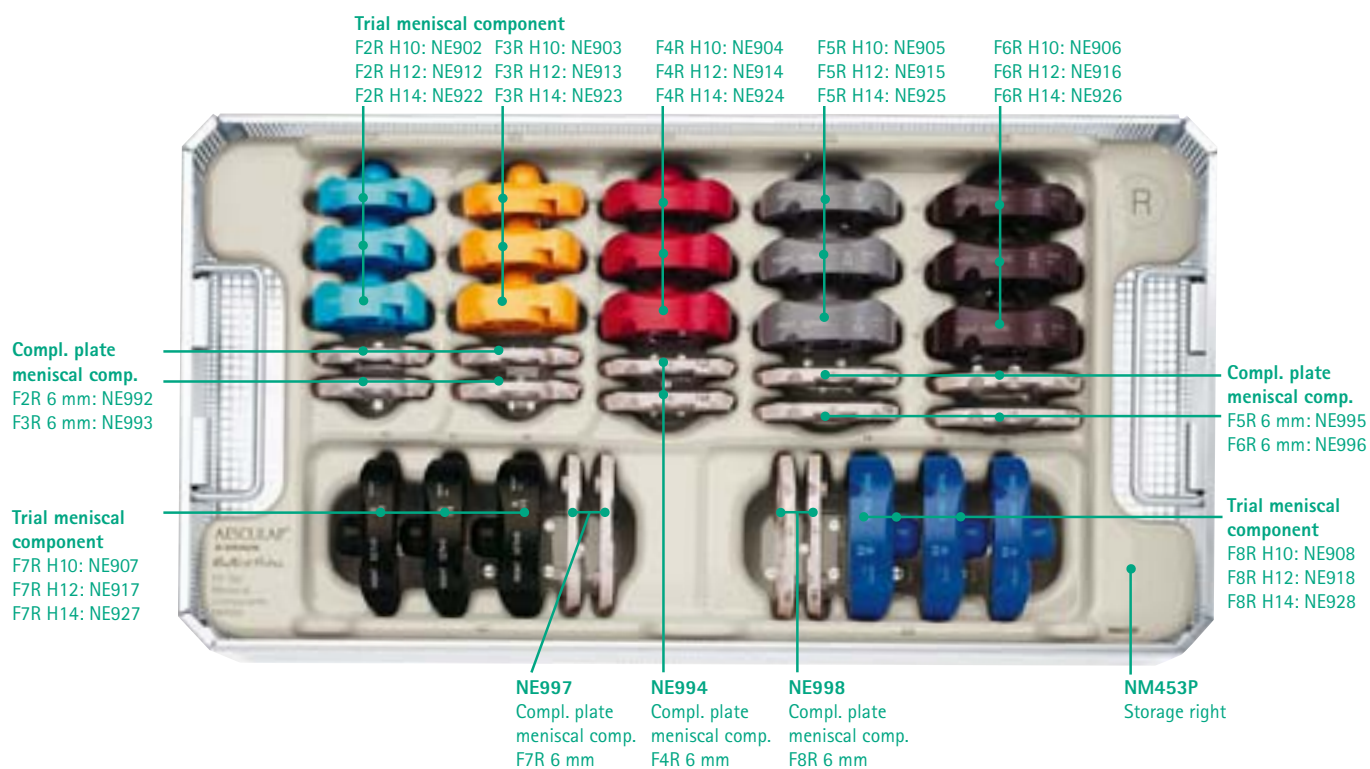


NM700 e.motion® PS

NM450 Trial Meniscal Components left

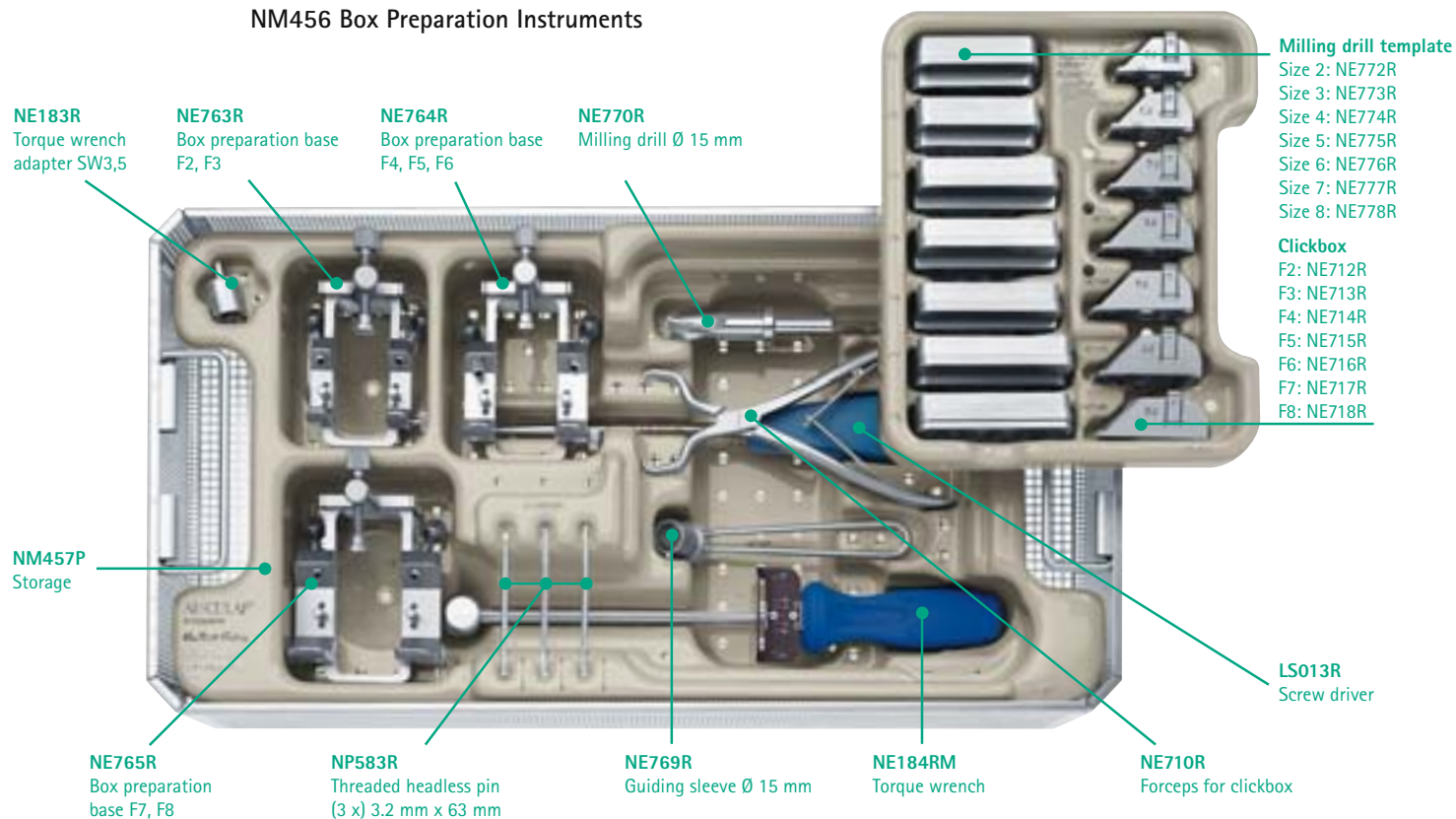


NM450 Trial Meniscal Components right



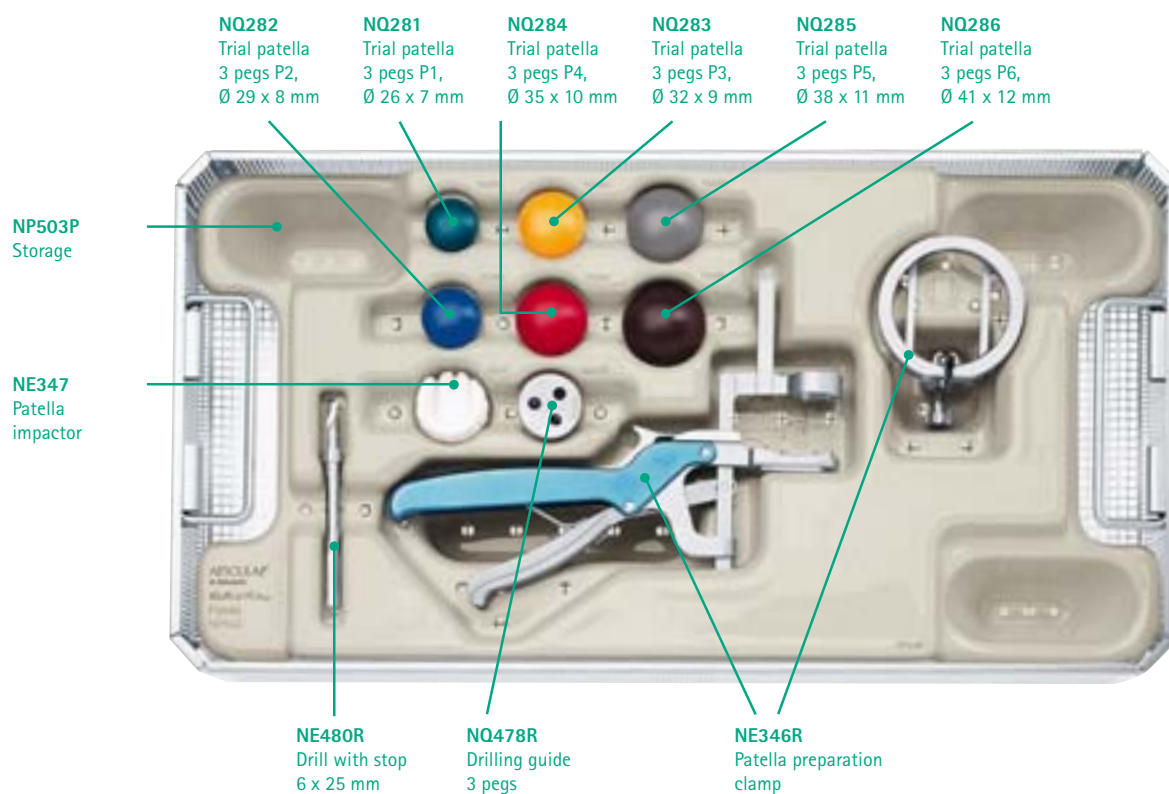
Instrumentarium

NM456 Box Preparation Instruments



NP502 e.motion® FP / PS / Revision

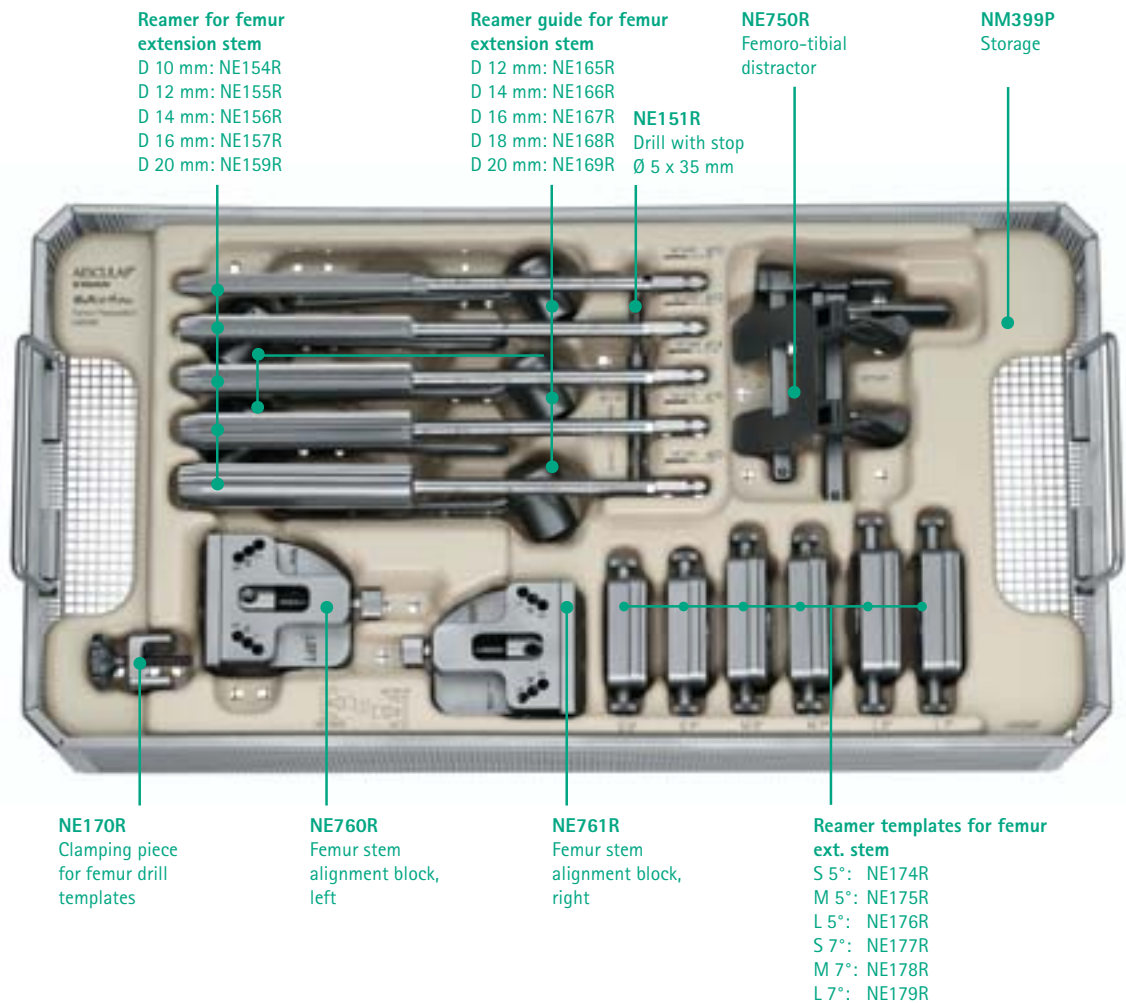
NP502 Patella Preparation Instruments



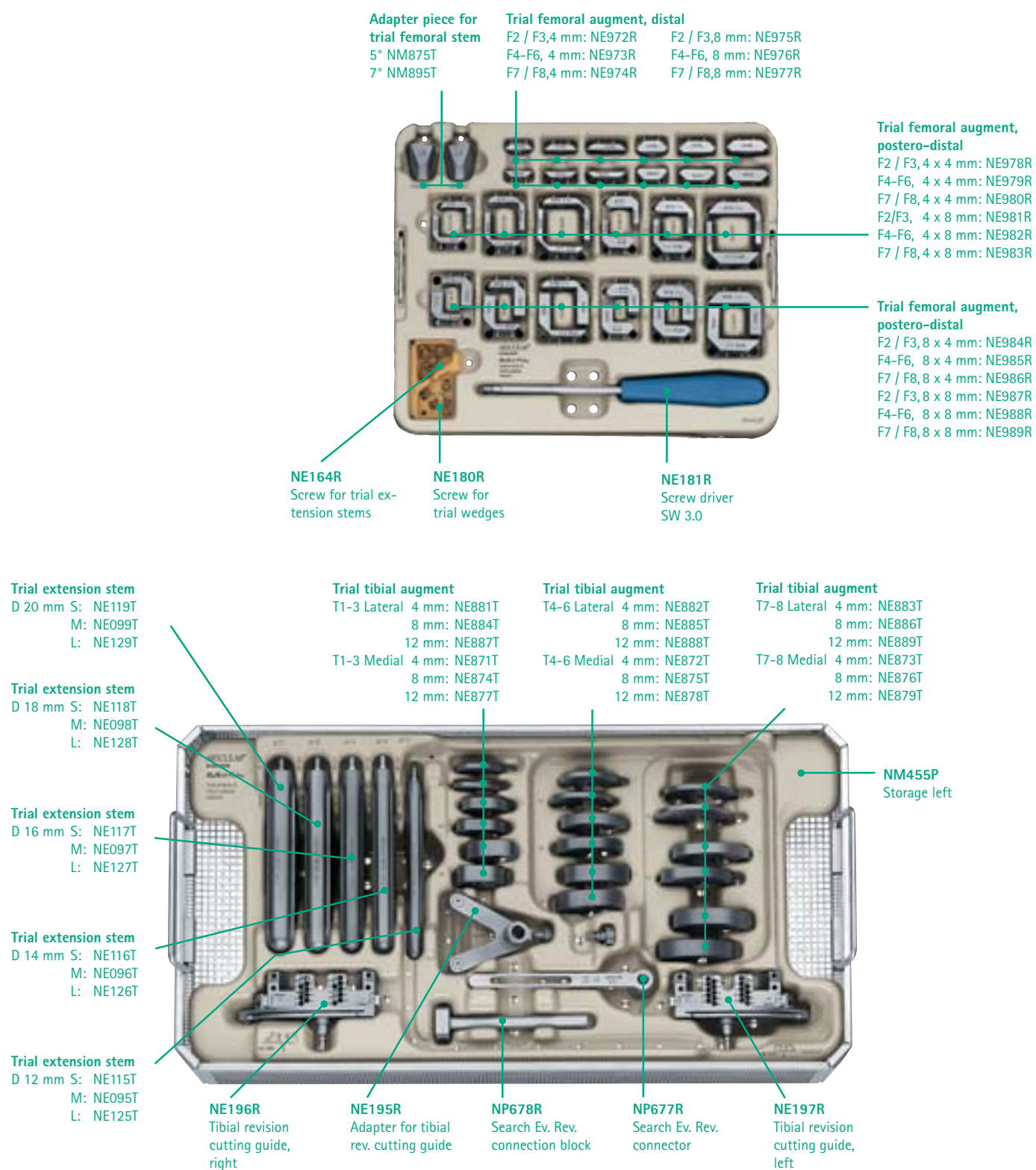
Instrumentarium

NM150 e.motion® PS / Revision

NM398 Femur Stem Preparation Instruments



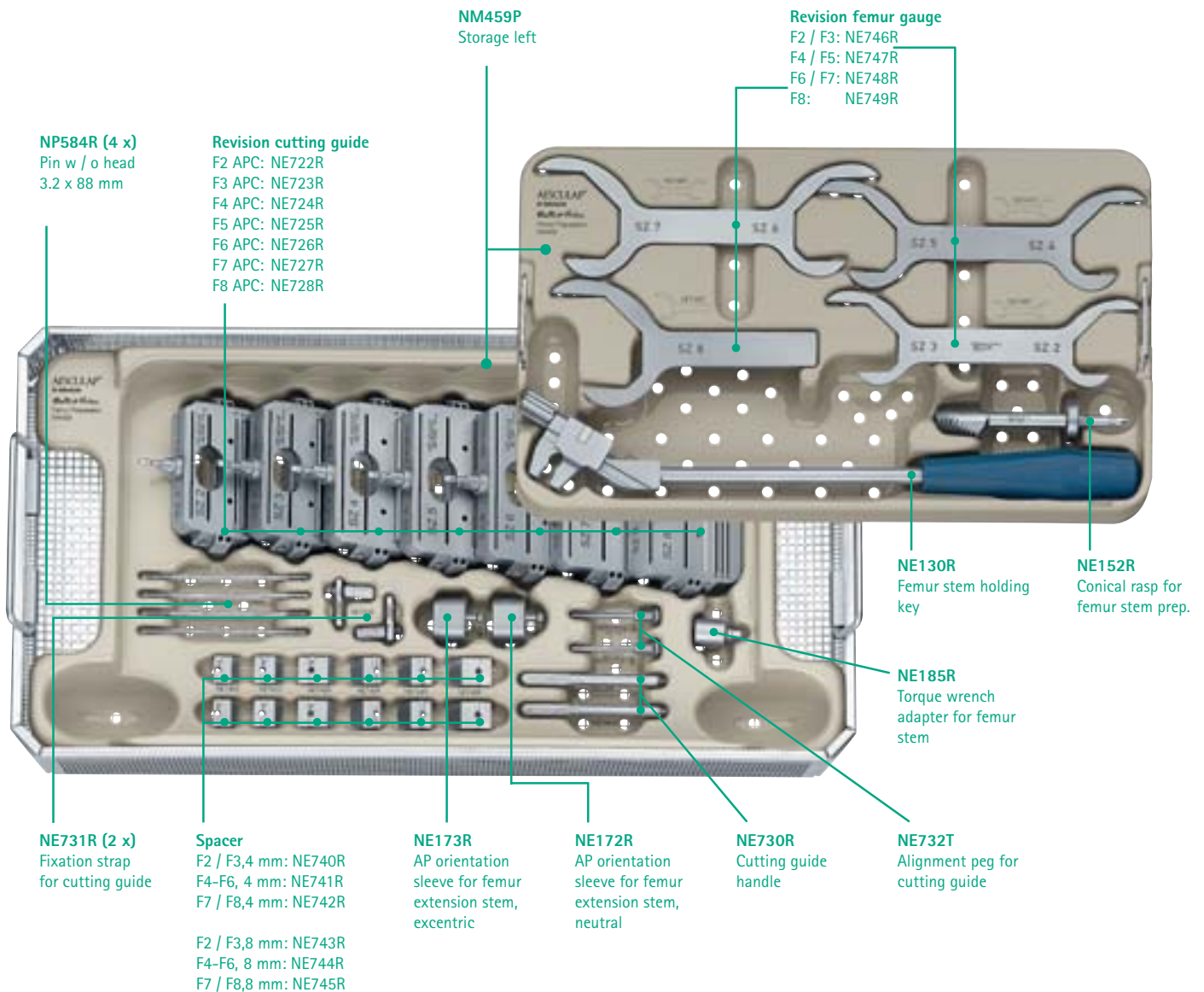
NM454 Instruments and Trial Implants



Instrumentarium

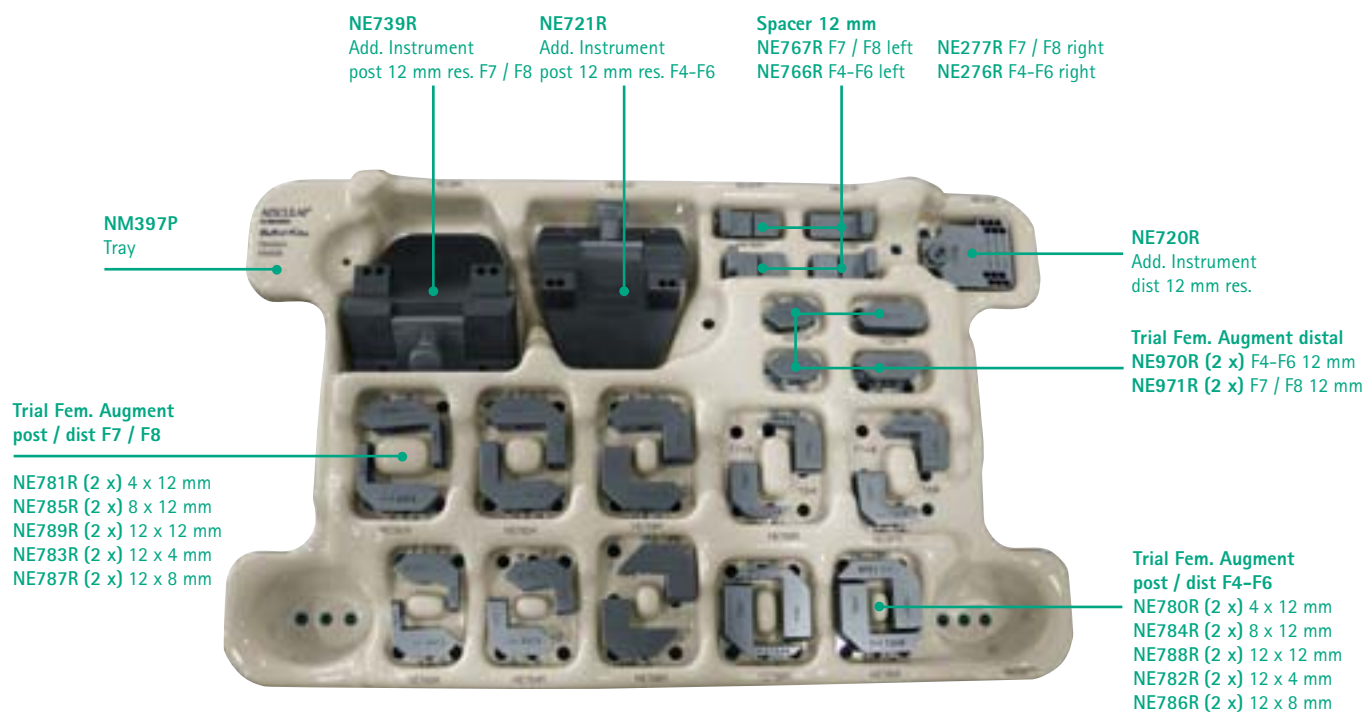
NM150 e.motion® PS / Revision

NM458 Femur Preparation Instruments



NM396 e.motion® PS / Revision

NM396 Trial Femur Augments 12 mm





Femur FP cemented

Types:	F2	F3	F4	F5	F6	F7	F8	Femur FP cementless				Nut for Femur Ext. Stem			
	Types:		F2	F3	F4	F5	F6	F7	F8					NB140K	
Left	N0502K	N0503K	N0504K	N0505K	N0506K	N0507K	N0508K	Left	N0582K	N0583K	N0584K	N0585K	N0586K	N0587K	N0588K
Right	N0602K	N0603K	N0604K	N0605K	N0606K	N0607K	N0608K	Right	N0682K	N0683K	N0684K	N0685K	N0686K	N0687K	N0688K

Femur PS cemented

Types:	F2	F3	F4	F5	F6	F7	F8	Femur PS cemented				Types:			
	Types:		F2	F3	F4	F5	F6	F7	F8					Short	
Left	NB702K	NB703K	NB704K	NB705K	NB706K	NB707K	NB708K	Left	NB702K	NB703K	NB704K	NB705K	NB706K	NB707K	NB708K
Right	NB752K	NB753K	NB754K	NB755K	NB756K	NB757K	NB758K	Right	NB752K	NB753K	NB754K	NB755K	NB756K	NB757K	NB758K

Gliding Surfaces

Types:	FP-left								UC-left								PS-left							
	F2	F3	F4	F5	F6	F7	F8	F2	F3	F4	F5	F6	F7	F8	F2	F3	F4	F5	F6	F7	F8			
10	N0542	N0543	N0544	N0545	N0546	N0547	N0548	NR802	NR803	NR804	NR805	NR806	NR807	NR808	NB802	NB803	NB804	NB805	NB806	NB807	NB808			
12	N0552	N0553	N0554	N0555	N0556	N0557	N0558	NR812	NR813	NR814	NR815	NR816	NR817	NR818	NB812	NB813	NB814	NB815	NB816	NB817	NB818			
14	N0562	N0563	N0564	N0565	N0566	N0567	N0568	NR822	NR823	NR824	NR825	NR826	NR827	NR828	NB822	NB823	NB824	NB825	NB826	NB827	NB828			
16	N0572	N0573	N0574	N0575	N0576	N0577	N0578	NR832	NR833	NR834	NR835	NR836	NR837	NR838	NB832	NB833	NB834	NB835	NB836	NB837	NB838			
18	N0592	N0593	N0594	N0595	N0596	N0597	N0598	NR842	NR843	NR844	NR845	NR846	NR847	NR848	NB842	NB843	NB844	NB845	NB846	NB847	NB848			
20	N0782	N0783	N0784	N0785	N0786	N0787	N0788	NR852	NR853	NR854	NR855	NR856	NR857	NR858	NB852	NB853	NB854	NB855	NB856	NB857	NB858			
22																	NB864	NB865	NB866	NB867	NB868			
24																	NB874	NB875	NB876	NB877	NB878			

Gliding Surfaces

Types:	FP-right								UC-right								PS-right							
	F2	F3	F4	F5	F6	F7	F8	F2	F3	F4	F5	F6	F7	F8	F2	F3	F4	F5	F6	F7	F8			
10	N0642	N0643	N0644	N0645	N0646	N0647	N0648	NR902	NR903	NR904	NR905	NR906	NR907	NR908	NB902	NB903	NB904	NB905	NB906	NB907	NB908			
12	N0652	N0653	N0654	N0655	N0656	N0657	N0658	NR912	NR913	NR914	NR915	NR916	NR917	NR918	NB912	NB913	NB914	NB915	NB916	NB917	NB918			
14	N0662	N0663	N0664	N0665	N0666	N0667	N0668	NR922	NR923	NR924	NR925	NR926	NR927	NR928	NB922	NB923	NB924	NB925	NB926	NB927	NB928			
16	N0672	N0673	N0674	N0675	N0676	N0677	N0678	NR932	NR933	NR934	NR935	NR936	NR937	NR938	NB932	NB933	NB934	NB935	NB936	NB937	NB938			
18	N0692	N0693	N0694	N0695	N0696	N0697	N0698	NR942	NR943	NR944	NR945	NR946	NR947	NR948	NB942	NB943	NB944	NB945	NB946	NB947	NB948			
20	N0792	N0793	N0794	N0795	N0796	N0797	N0798	NR952	NR953	NR954	NR955	NR956	NR957	NR958	NB952	NB953	NB954	NB955	NB956	NB957	NB958			
22																	NB964	NB965	NB966	NB967	NB968			
24																	NB974	NB975	NB976	NB977	NB978			

Distal Femur Wedges

Antero-Medial Femur Wedges								Postero-Distal Femur Wedges								
Types:	F2	F3	F4	F5	F6	F7	F8	Types:	F2	F3	F4	F5	F6	F7	F8	
4 mm	NB282K	NB283K	NB284K	NB285K	NB286K	NB287K	NB288K	P1	N0481	NB302K	NB303K	NB304K	NB305K	NB306K	NB307K	NB308K
8 mm	NB292K	NB293K	NB294K	NB295K	NB296K	NB297K	NB298K	P2	N0482	NB312K	NB313K	NB314K	NB315K	NB316K	NB317K	NB318K
12 mm			NB274K	NB275K	NB276K	NB277K	NB278K	P3	N0483			NB340K	NB341K	NB342K	NB343K	NB344K

Femur Extension Stems cementless

5°								7°							
Types:	Ø 14	Ø 16	Ø 18	Ø 20	Ø 14	Ø 16	Ø 18	Types:	Ø 14	Ø 16	Ø 18	Ø 20	Ø 14	Ø 16	Ø 18
Short	NB236K	NB237K	NB238K	NB239K	NB256K	NB257K	NB258K	Short	NB236K	NB237K	NB238K	NB239K	NB256K	NB257K	NB258K
Middle	NB241K	NB242K	NB243K	NB244K	NB261K	NB262K	NB263K	Middle	NB241K	NB242K	NB243K	NB244K	NB261K	NB262K	NB263K
Long	NB246K	NB247K	NB248K	NB249K	NB266K	NB267K	NB268K	Long	NB246K	NB247K	NB248K	NB249K	NB266K	NB267K	NB268K



e.motion® Implant Matrix-Tibial-Parts

Tibia FP Monobloc cemented



Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left	NB521K	NB522K	NB523K	NB524K	NB525K	NB526K	NB527K	NB528K
Right	NB621K	NB622K	NB623K	NB624K	NB625K	NB626K	NB627K	NB628K

Tibia PS Modular cemented



Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left	NB731K	NB732K	NB733K	NB734K	NB735K	NB736K	NB737K	NB738K
Right	NB781K	NB782K	NB783K	NB784K	NB785K	NB786K	NB787K	NB788K

Tibia FP Modular cemented

Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left	NB521K	NB522K	NB523K	NB524K	NB525K	NB526K	NB527K	NB528K
Right	NB621K	NB622K	NB623K	NB624K	NB625K	NB626K	NB627K	NB628K

Tibia FP Modular cementless

Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left	NB591K	NB592K	NB593K	NB594K	NB595K	NB596K	NB597K	NB598K
Right	NB691K	NB692K	NB693K	NB694K	NB695K	NB696K	NB697K	NB698K

Tibia PS Modular cementless

Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left	NB741K	NB742K	NB743K	NB744K	NB745K	NB746K	NB747K	NB748K
Right	NB791K	NB792K	NB793K	NB794K	NB795K	NB796K	NB797K	NB798K

Tibia Ext. Stems cemented



Types:	Ø 12	Ø 14	Ø 16
Short	NB213K	NB214K	NB215K
Middle	NB218K	NB219K	NB220K
Long	NB223K	NB224K	NB225K

Tibia Ext. Stems cementless



Types:	Ø 10	Ø 12	Ø 14	Ø 16
Short	NB114K	NB115K	NB116K	NB117K
Middle	NB094K	NB095K	NB096K	NB097K
Long	NB124K	NB125K	NB126K	NB127K

Tibia-Obturator



Types:	Ø 12	Ø 14	Ø 16
T1 / T2 / T3	NB105K	--	--
T4 / T5 / T6	--	NB106K	--
T7 / T8	--	--	NB107K

Tibial Wedges Medial



Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left + Right	NB401K	NB402K	NB403K	NB404K	NB405K	NB406K	NB407K	NB408K

Tibial Wedges Lateral



Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left + Right	NB411K	NB412K	NB413K	NB414K	NB415K	NB416K	NB417K	NB418K

8 mm

Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left	NB421K	NB422K	NB423K	NB424K	NB425K	NB426K	NB427K	NB428K
Right	NB431K	NB432K	NB433K	NB434K	NB435K	NB436K	NB437K	NB438K

8 mm

Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left	NB441K	NB442K	NB443K	NB444K	NB445K	NB446K	NB447K	NB448K
Right	NB451K	NB452K	NB453K	NB454K	NB455K	NB456K	NB457K	NB458K

12 mm

Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left	NB461K	NB462K	NB463K	NB464K	NB465K	NB466K	NB467K	NB468K
Right	NB471K	NB472K	NB473K	NB474K	NB475K	NB476K	NB477K	NB478K

12 mm

Types:	T1	T2	T3	T4	T5	T6	T7	T8
Left	NB461K	NB462K	NB463K	NB464K	NB465K	NB466K	NB467K	NB468K
Right	NB471K	NB472K	NB473K	NB474K	NB475K	NB476K	NB477K	NB478K

