Flexibility through Modularity
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Prostheses with Mobile Bearings

Innex Type CR

Mobile bearing (Cruciate Retaining)
For preservation of the posterior cruciate ligament

Suitable for:
- Osteoarthritis with sufficient posterior cruciate ligament
- Mild or passively correctable varus or valgus deformity
- Mild to moderate extension deficit
- Sufficient ligament stability

Innex Type UCOR

Mobile bearing (Ultra-Congruent Only Rotating)
For resection of the posterior cruciate ligament

Suitable for:
- Osteoarthritis with insufficient posterior cruciate ligament and chronic polyarthritis
- Partial contracture with moderate varus or valgus deformity
- Moderate to severe extension deficit
- Sufficient collateral ligament stability
Prostheses with Fixed Bearings

Innex Type FIXCR

Fixed bearing (Cruciate Retaining)
For preservation of the posterior cruciate ligament

Suitable for:
- Osteoarthritis with sufficient posterior cruciate ligament
- Mild or passively correctable varus or valgus deformity
- Mild to moderate extension deficit
- Sufficient ligament stability
- Increased dislocation risk of the mobile bearing due to residual ligament imbalance

Innex Type FIXUC

Fixed bearing (Ultra-Congruent)
For resection of the posterior cruciate ligament

Suitable for:
- Osteoarthritis with insufficient posterior cruciate ligament and chronic polyarthritis
- Partial contracture with moderate varus or valgus deformity
- Moderate to severe extension deficit
- Sufficient collateral ligament stability
- Increased dislocation risk of the mobile bearing due to residual ligament imbalance
Preoperative Planning

The preoperative planning comprises the indication, evaluation and preparation important for the success of the surgery. Preoperative planning also includes patient education, preoperative medical examinations, assessing the risk of the surgery and choice of anesthesia.

Preoperative X-rays are essential for surgical planning. A single-leg stance X-ray in the anterior/posterior (A/P) plane and a long-leg X-ray with weight on both legs are recommended. In addition, it may be useful to have a lateral X-ray of the knee joint in 90° flexion or in extension as well as a skyline view of the patella in 40° flexion.

Long-leg X-rays help to detect deviations of the axis and deformities in the diaphyseal area of the femur and the tibia. Long-leg X-rays help also to determine whether an intramedullary or an extramedullary alignment can be used.

Furthermore, the mechanical and anatomical axes of the leg can be plotted and the femoral angle \( \alpha \), representing the difference between the two, can be determined (see page 8). This angle is usually about 6° but varies depending on morphology and femur size. The distal femoral cut is determined by the femoral angle bushing. The bushing is chosen according to the femoral angle \( \alpha \) (see pages 20 and 22). Three femoral angle bushings are available with 4°, 6° and 8° valgus angle.

The entry point for the intramedullary alignment guide is determined by extending the line of the anatomical axis of the femur. Usually, the entry point is slightly medial to the center of the femoral condyles.

The extent of tibial resection can be determined on the long-leg X-ray. In this way, the magnitude of the medial and lateral bone resection can be evaluated (see page 8). This is especially important for extensive bone defects in order to avoid a too radical resection. It is preferable to treat defects in primary knee arthroplasty with autologous bone grafts.
Assessment of the posterior slope of the tibial plateau is made with a lateral X-ray. In general this slope ranges between 3° and 10°.

The status of the patella alta/baja is determined from the lateral X-ray. Any tilt/subluxation of the patella can be seen in the skyline view. This information can influence additional intraoperative decisions regarding the extensor mechanism.

It is not recommended to resurface the patella as a routine. In fact, resurfacing is doubtful in the case of a alta/baja patella position.

Where patella replacement is indicated, there are four sizes of cemented patella implants available.
Preoperative Planning with X-Ray Templates

When examining the A/P long-leg X-ray, it is recommended to proceed in the following manner:

1. Draw the anatomical axis of the femur (A) on the X-ray. If the femur is excessively curved, a line materializing the intramedullary alignment should be drawn instead of the line A.

2. Draw a line from the center of the femoral head to the center of the knee (mechanical axis D) on the X-ray.

3. The angle measured between the anatomical and the mechanical axis (α, the femoral valgus angle) determines the choice of the femoral angle bushing (see page 20). The valgus angle α is usually between 4° and 8° and should be determined for each patient.

4. Draw the axis of the tibia (B), and determine the tibial resection plane (E) perpendicular to B. Take care to prevent a too extensive resection in the case of any tibial defects.

5. Preoperative determination of component size and resection depth using the X-ray templates in the A/P and lateral planes.

6. After the resection, the mechanical axis of the leg (C) should match up with lines D and B.
**Tourniquet**

The surgery is carried out on patients under general or spinal anesthesia. Postoperative pain is significantly reduced without the use of a tourniquet. Furthermore, a tourniquet clamps the quadriceps muscle and interferes with its tension. Optimal muscle relaxation is necessary for surgery. In addition, it is safer to control bleeding throughout the procedure than only at the end of the surgery. If necessary to apply a tourniquet, it should be placed as high as possible and be inflated at maximum 100 mm Hg above systolic blood pressure.

**Patient Positioning**

Place the patient in the supine position. Flex the knee into a 90° position. Use a supporting roll on the table and a lateral support to facilitate extension and flexion of the leg. Alternatively, a knee positioner could be used as the position of the knee has to be changed frequently throughout the procedure in order to optimize access to the tibia and femur and better utilize the soft tissue window.
Surgical Instructions

Preparation of the Soft Tissues

The goal is to maintain a consistent rectangular gap during flexion and extension after bone resection. The gap should not be rhomboid.

The tibia first method is preferred by several surgeons who recommend performing necessary soft tissue releases before any bone cuts are made. They recommend using the following procedure:

1. Open the knee joint and make a preliminary balancing of the soft tissue structures appropriate to the situation. Remove osteophytes.

2. Perform bone cuts according to the preoperative plan.

3. Make a fine adjustment of soft tissues after checking the flexion and extension gaps with the gap gauge, or at the latest after inserting the trial prosthesis.

This prevents the release being too extensive, which would result in laxity of the ligaments after insertion of the final implant.

Based on their observations it is recommended undertaking a $\frac{1}{2}$- to $\frac{3}{4}$-resection of the Hoffa fat pad. This is independent of surgical approach. Fibrosis or a fat pad impingement can cause anterior knee pain.\(^1\)

In order to maintain the continuity of the joint capsule, the base of the meniscus should be left intact when resecting the meniscus or its remnants.

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Overview of the Cutting Steps
Tibia First

It is important to define the flexion gap first and then to adjust the extension gap to the flexion gap, and not vice versa. The correct ligament balance of the prosthesis can only be achieved if the flexion and extension gaps are identical.

1. Tibia cut
2. Anterior/posterior femur cut
3. Distal femur cut
4. Chamfer cuts and trochlea recess
5. Patella cut (optional). The patella cut can be done at the beginning of surgery in order to better expose the joint

Note
Only 1.3 mm saw blades should be used for all bone cuts!
Preparation of the Tibia

1. Tibia cut
Cut 1a and 1b for type CR and type FIXCR or cut 1b only for type UCOR and type FIXUC (see cutting steps on page 11).

Extramedullary variants
Preparation of the tibia alignment guide
The tibia alignment guide is assembled as follows:

1. Lock the drill guide tube (A) in the out-position. Due to its conical design, the tube is pressed on and therefore does not need a mechanical locking mechanism. (Fig. 1)
2. Place the tibia cutting block (B) on the tibia alignment guide (Fig. 2) by sliding it on to the two vertical guide rods. The Innex Knee System offers two asymmetric tibia cutting blocks for primary resection and two 2° tibia correction cutting blocks for correction resection.
3. Turn the height adjustment screw (C) upwards, as far as it will go (mechanical stop). The height adjustment screw can be moved over the threads more freely by pressing the locking button (D). (Fig. 2)
4. Use the locking screws for length (E) and slope (F) to temporarily set the tibia alignment guide for length and slope. (Fig. 2)
5. Set the distal locking screw (G) in the Medial/Lateral (M/L) plane to adjust for the varus/valgus slope of the proximal tibial cut. The neutral position is marked. (Fig. 2)

Positioning the tibia alignment guide
Fix the tibia alignment guide proximally in the center of the tibial plateau in the area of the tibial eminence using a headed guide pin. Hammer the pin through the guide tube, so that the tibia alignment guide can still be rotated. Set the definitive length on the tibia alignment guide with the locking screw (E). (Fig. 3)
Determination of rotation
The correct rotation positioning for the tibial alignment can be well defined, as a rule, with the following four orientation aids:

1. The reference point on the proximal tibia is an imaginary point, roughly 5 mm medial from the center of the tibial tuberosity. The tibia alignment guide should be aligned to this point.

2. Two distal holes in the tibia alignment guide (keep in mind right/left) allow the guide to be aligned in parallel to the malleolar axis. This is done with Steinmann pins.

3. Normally the tibia alignment guide points distally in the direction of metatarsal II.

4. In addition, there is also the possibility of aligning the tibia alignment guide in the middle of the ankle joint by releasing the stopping screw (G). (Fig. 4)

After correct alignment, the tibia alignment guide is fixed proximally in position with a second-headed guide pin. (Fig. 5)

Determination of slope
The angle for the posterior slope of the tibial cut is adjusted to the physiological inclination of the tibial plateau (in accordance with preoperative planning). A dorsally inclined tibial cut improves the ability of the knee to flex and prevents an overextension of the tendons during flexion.

The angle of slope is set by releasing the distal stopping screw (F) and shifting the distal part of the tibia alignment guide. (Fig. 5)

Note
The middle of the ankle joint is also suitable as a reference point. The tendon of the anterior tibial muscle can be easily palpated at the level of the ankle joint. This point corresponds approximately to the center of the top of the ankle joint. Alternatively, the lateral edge of the tendon can also be used as the reference point for the middle of the ankle joint. This edge is more distal, at the height of the top of the ankle joint.

Note
Setting the tibial alignment parallel to the axis of the tibial shaft corresponds to a posterior slope of 6°.

Note
The physiological inclination of the tibial plateau can be checked using the resection-guide-placed flush on the cutting block.

Note
If an anchoring stem is used, the surgeon must carefully choose the posterior tilt of the tibial cut. This is necessary to avoid the anchoring stem impinging on the anterior tibial cortex.
**Intramedullary option**

**Opening the medullary canal**
Position the knee in full flexion and subluxate the tibia forward with a Homann retractor.

Determine the insertion point for the intramedullary alignment rod based on the A/P and the lateral X-rays.

In general, the insertion point is situated mediolaterally in the middle of the tibial plane. In the A/P direction this point is directly in the area of insertion of the anterior cruciate ligament. (Fig. 6/7)

Open the medullary cavity with the $\varnothing$ 8 mm drill. Make sure that the drill enters the tibial metaphysis as close to the center as possible. (Fig. 8)

Clean the intramedullary canal by suction before inserting the intramedullary rod. This reduces the danger of fat particle embolisms.

Carefully insert the intramedullary alignment guide with turning movements and then remove the handle. (Fig. 9)

**Positioning the tibia alignment guide**
Slide the tibia alignment guide onto the intramedullary rod, and fix the rotation position with a headed pin. (Fig. 10)

The subsequent procedure corresponds to that of the extramedullary (beginning on page 15).
Determining the resection height

Screw and securely fasten the tibial stylus onto the tibia cutting block. (Fig. 11)

Adjust the resection height using the end of the tibial stylus. Lower the tibia cutting block with the height-adjustment screw (C) until the tip of the stylus is set at the highest point of the intact compartment. This position corresponds roughly to the physiological joint line. It is important to ensure that the height adjustment screw (C) is always in contact with the tibia cutting block (B). (Fig. 11)

Perform the tibial resection 10 mm below the highest point of the intact compartment (resection plane). The intact compartment can also have defects in cartilage. This must be considered when determining the height of the resection.

The resection plane can be lowered in 2 mm steps if required by turning the height adjustment screw (C) 360° clockwise.

After the height adjustment screw (C) has been set at the desired resection level, remove the tibial stylus so that the tibia cutting block (B) sits at the correct level.

Check the resection plane with the resection gauge before carrying out the final cut. This is especially important in the case of tibial bone defects. (Fig. 12)

Note
The tip of the tibial stylus (0 mm) lies at the same height as the cutting plane of the tibia cutting block. The setting of the desired resection plane is done manually by turning the height-adjustment screw (C).

The tibia cutting block is brought directly to the level of the recommended resection plane by using the tip of the tibial stylus (–10 mm).
Fixing the tibia cutting block
Release the drill bushings by turning and applying light pressure and at the same time pushing forward on the anterior edge of the tibia. This ensures an optimal control of the drill.

Drill the first hole (∅ 3.2 mm) and fix with a Steinmann pin. Prepare the second hole in the same manner. (Fig. 13)

A third Steinmann pin and a fourth oblique Steinmann pin can be used for additional fixation and improve stability.

Removal of the tibia alignment guide
Remove the tibia alignment guide with exception of the tibia cutting block. This is done as follows:

1. Move the height adjustment screw (C) to its lowest point (see page 12).
2. Loosen the stopping screw (E) to release the distal portion of the alignment guide.
3. The headed pin can be removed beforehand if need be. (Fig. 14)

Checking the alignment
Attach the modular handle to the tibia cutting block and insert the alignment rod through the handle. This is in order to check the correct alignment of the cutting block. (Fig. 15)

Correct the position of the tibia cutting block if the alignment guide shows a deviation from the middle of the ankle joint. A correction of 2° is done with the 2° tibia correction cutting block. (Fig. 16) The tibia cutting block must be repositioned for a deviation larger than 2°.

Note
It is recommended to drive in the Steinmann pin directly (without predrilling) if the patellar ligament is in the immediate vicinity of the bore hole. This avoids damaging the patellar ligament.
Tibia resection type CR and type FIXCR
Determine the approximate size of the cut out for the posterior cruciate ligament using the trial tibial plate. Use electrocautery for marking and protect the ligament with an osteotome.

Make two sagittal cuts (cut 1a, page 11) with a reciprocating saw. (Fig. 17)

The saw blade capture is attached to the tibia resection cutting block. Resect the tibia whilst protecting the collateral ligaments (cut 1b, page 11). (Fig. 18)

The saw cut should be completed in order to remove the resected bone in one piece. Later resection of posterior residues is difficult.

Tibia Resection type UCOR and type FIXUC
The posterior cruciate ligament is resected when implanting the UCOR and FIXUC types. Tibia resection is limited to the horizontal tibial cut (cut 1b, page 11) only.

There is no clearance for the posterior cruciate ligament in the final bearings (implants).

Note
The Steinmann pins should be replaced by the short hole markers. For the case of a later additional resection the Steinmann pins can be reinserted.

Note
Check the alignment again. Attach the modular handle to the trial tibial plate and insert the alignment rod through the handle. This is in order to check the correct alignment after the tibial cut.
Preparation of the Femur

Flex the knee 90° to access the femur from the front. Resect osteophytes from both femur and tibia (see handling of the soft tissues, page 10). Perform a preliminary mediolateral ligament balancing. This facilitates anterior femoral resection and prevents muscle violation.

2. A/P femoral cuts
   (Cuts 2a and 2b)
   Opening the medullary canal
   Osteophytes should have been removed prior to this step.

   However, if any osteophytes are still remaining, they should be removed now to facilitate orientation.

   Set the femoral intercondylar drill guide on the distal femur above the intercondylar fossa and drive it in. The alignment must correspond to the longitudinal axis of the femur in both planes. (Fig. 19–22)

   The intercondylar insertion point for the femoral intercondylar drill guide tends to lie slightly medial of the midline. This can be seen in the preoperative A/P X-ray of the femur (long-leg X-ray).

   Open the intramedullary space with the ∅ 8 mm drill. Align the drill with the anatomical axis of the femur. (Fig. 22)

   If necessary, the drill can be driven in deeper by removing the drill guide.
Rinse the intramedullary canal by suction before inserting the intramedullary alignment guide. This reduces the danger of fat particle embolisms. The alignment rod is then carefully inserted with turning movements and the handle removed. (Fig. 23)

**Sizing the femoral component**

Determine the size of the femoral component in both the A/P and the M/L directions.

Mount the femoral size gauge on the intramedullary alignment rod to determine the A/P dimension of femoral component. (Fig. 24)

Both feet of the femoral size gauge must have good contact with the posterior condyles. The point of the stylus should rest on the anterior femoral cortex.

**Place the point of the stylus on the deepest point of the anterior femoral cortex above the edge of the cartilage in order to obtain optimal measurement.**

The size of the femoral component is read on the vertical scale. In the case of intermediate sizes, choose the smaller size.

Check the dimension in the M/L direction with the femoral gauge. (Fig. 25) In cases of intermediate sizes, this measurement can facilitate the choice of the final component. The size determined in the A/P direction can be checked with the other side of the femoral gauge.

**Note**

The letters on the vertical scale indicate the size of the femoral component (S, S+, M, M+, L, L+). The A/P dimensions are identical for gender and standard implants.
A/P resection

Base the choice of A/P cutting guide on the preliminary determined size of the femoral component. Determine the femoral angle bushing according to the valgus angle. (Fig. 26) Insert the bushing into the A/P cutting guide. The A/P resection guide has a symmetrical design and the quick-release button (I) can be on the medial or lateral side. The arrow on the quick-release button should point to “open” when the femoral angle bushing is inserted.

The angle of the femoral angle bushing (4°, 6° or 8°) should correspond to the valgus angle α (see page 8). The valgus angle is usually around 6°.

If the angle α is between two bushing angles, select the next smaller angle for a genu varum or the next larger angle for the genu valgum.

Attach the anterior femoral stylus to the A/P cutting guide. (Fig. 26)

Slide the prepared A/P cutting guide onto the intramedullary alignment rod. In order to have an optimal measurement, position the point of the stylus on the deepest point of the anterior femoral cortex above the edge of the cartilage. Then fix the femoral stylus with a screw (H) on the correct size setting. (Fig. 27)

Femoral rotation is now roughly aligned to the epicondylar axis or the antero-posterior femoral axis. Lock the femoral angle bushing in the A/P plane with the quick-release button (I). The arrow on the quick-release button should point to “lock”. (Fig. 28) Now remove the femoral stylus.

Note

A 4–6° femoral angle is generally preferred in a varus knee. A 6–8° angle is preferred in a valgus knee. This makes the balancing of the ligaments easier and corresponds to the individual anatomical conditions of the knee joint.
Setting femoral rotation and ligament tension

Insert the balancer into the A/P cutting guide. For an easy assembly the balancer should be in the minimal position. Then stretch the balancer to the smallest overall tibial height of 10 mm which is the first stop. (Fig. 29) The desired external rotation of the A/P cutting guide (approximately 3°) is usually obtained automatically by equalizing tension of the medial and lateral soft tissues by cranking up the balancer.

Check the femoral rotation via the epicondylar axis and the Whiteside line with the aid of 2 Steinmann pins inserted laterally in the A/P cutting guide. (Fig. 29)

Repeat resection of the tibia if the gap is too small for the balancer or the ligament tension is too high when the balancer is inserted. Therefore the short note markers should be replaced by the Steinmann pins. The resection can be performed in 2 mm steps. (Fig. 30) If the ligament tension is insufficient, the balancer can be cranked up with the screwdriver to a thicker implant height until the desired ligament tension is achieved.
If the ligament tension is insufficient, the balancer can be cranked up with the screwdriver to a thicker implant height until the desired ligament tension is achieved.

The balancer has a snap mechanism which stops at each implant height (10/12.5/15/17.5 mm). To move to a thicker implant push the button and crank up the balancer.

To move forward to the next thickness, loosen the screw a quarter, push the button and move the paddles to the next thickness with the screwdriver at the same time. (Fig. 31)

Drill two holes at the “0” mark on the A/P cutting guide and fix it with Steinmann pins. Remove the balancer. Check the resection height with the resection gauge in order to prevent anterior notching. (Fig. 32) Anterior notching can weaken the anterior cortex.

Perform the anterior femoral cut (cut 2a, page 11). If the cut is too far anterior, it is still possible to lower the cutting plane (2 mm or 4 mm) by shifting the A/P cutting guide down. The intramedullary alignment rod must be removed in this case. Perform the posterior femoral cut (cut 2b, page 11). The attachable saw capture improves accuracy of the cut. (Fig. 33) Resect the femoral cuts whilst protecting the collateral ligaments.

Note
Do not move the femoral angle bushing. In certain circumstances, the tibial cut (cut 1b, page 11) must also be adjusted to the new situation. Ensure that the collateral and – when using CR/FIXCR – cruciate ligament structures are not damaged when making the posterior femoral cut.
3. Distal femoral cut

(Cut 3)

Remove posterior osteophytes on the femoral condyles and, if necessary, release the posterior capsule before making the distal femoral cut. This makes it easier to determine the depth of the distal femoral resection. This procedure is especially important in the case of a severe flexion contracture.

Assembling the distal cutting guide and the anterior platform

Insert the anterior outrigger into the distal cutting guide and slide it to the “primary” marking after pressing the button (F). *(Fig. 34)*

Attach the anterior outrigger and distal cutting guide to the A/P cutting guide. The distal cutting guide must lie flush on the anterior femoral resection plane. Drill with the ø 3.2 mm drill through the two guide holes and fix the distal cutting guide with Steinmann pins. *(Fig. 35)*

Remove the anterior outrigger by pressing the button on the distal cutting guide and by pulling the outrigger distally. Then remove the A/P cutting guide, the Steinmann pins and the intramedullary alignment rod. *(Fig. 36)*

Now assemble the second part (H) of the distal cutting guide on the “0” position. *(Fig. 37)*

**Note**

If there is a problem sliding the distal cutting guide onto the A/P cutting guide, open the quick-release button (G) on the A/P cutting guide. The Steinmann pin must be removed from the A/P cutting guide in order to do this.
Checking the alignment
Extend the leg to assess the alignment.

Assemble the alignment guide and the modular handle and attach it to the distal cutting guide. (Fig. 38)

The alignment guide should follow the mechanical axis of the leg, and be aligned with the center of the femoral head, about two finger widths medial to the anterior superior iliac spine.

Note
Correct the position of the femoral cutting block if the alignment guide shows a deviation from the middle of the upper ankle joint. A correction of 2° is possible by choosing a different fixation hole in the cutting block. (Fig. 39)

Checking the flexion gap
Put the femoral spacer plate and the tibial spacer plate together to form the spacer gauge. Check the flexion gap with the spacer gauge. (Fig. 40) The spacer gauge without the tibial inserts corresponds to the smallest overall gap height of 10 mm.
In case of insufficient ligament tension, add two tibial inserts of the same thickness to the spacer gauge until the desired ligament tension is achieved. (Fig. 41)

Repeat resection of the tibia if the gap is too narrow for the gap gauge or the ligament tension is too high. (Fig. 42) Therefore the short hole markers should be replaced by the Steinmann pins.

Remove the Steinmann pins from the proximal tibia if both flexion gaps are of identical height.

**Preliminary check of the extension gap**
Extend the knee joint and check the level of the distal femoral resection with the Balancer. Slide the femoral spacer plate toward the handle so that the spacer gauge can be inserted between the tibial resection surface and the distal cutting guide. This procedure allows an assessment of the extension gap height.

The balancer can be used to check the extension gap in the same manner as the flexion gap. (Fig. 43)

The extension and flexion gaps must be of identical height and the mediolateral ligament tension must be balanced (page 10).

**Note**
The distal cutting guide can be shifted proximally or distally by 1 mm increments if adjustments are necessary.
Attach the saw capture to the distal cutting guide and make the distal femoral cut (cut 3, page 11). \textit{(Fig. 44)}

Remove the femoral distal cutting guide.

\textbf{Final check of the extension and flexion gaps}

Insert the spacer gauge into the extension gap between the tibial and femoral resections with the leg in extension. The thickness of the spacer gauge in the extension gap must be the same as in the flexion gap. Check the balance and tension of the ligaments as well as the leg axes. \textit{(Fig. 45/46)}

If the spacer gauge cannot be inserted (extension gap too narrow), then resect the distal femur until the extension and flexion gaps are identical.

If the operative steps are correctly performed and checked, the vice-versa situation (extension gap larger than flexion gap) should not occur.
4. Chamfer cuts and trochlea recess
(Cuts 4a, 4b, 4c and femur peg holes)
Choose the chamfer cutting guide according to the predetermined femoral component size.

Assembly of the chamfer cutting guide with the holder
The holder for the chamfer cutting guide allows good positioning on the distal femur:

1. Locate the single peg in the bottom hole of the chamfer cutting guide and attach the holder. (Fig. 47)

2. Secure the holder to the chamfer cutting guide by turning the handle. (Fig. 48)

The chamfer cutting guide is now ready to be placed on the distal femur. (Fig. 49)

Mediolateral positioning is facilitated as the width of the chamfer cutting guide matches exactly the width of the femoral component. Both the A/P and M/L dimensions of the chamfer cutting guide correspond to the final implant size.
Fix the chamfer cutting guide with two Steinmann pins inserted through the anterior drill holes (Ø 3.2 mm). Remove the holder by turning the handle.

Drill both femur cog holes with the Ø 5 mm drill. (Fig. 50)

Use the saw blade slot in the chamfer cutting guide to perform the anterior (cut 4a, page 11) and posterior (cut 4b, page 11) chamfer cuts. (Fig. 51/52)
Prepare the trochlea recess of the femoral component using a jigsaw and corresponding osteotome (15 mm). *(Fig. 53)*

Bring the knee joint into maximum flexion and remove the remaining posterior osteophytes with a curved osteotome or oscillating saw (cut 4c). The posterior edge of the chamfer cutting guide serves as a reference. *(Fig. 54)* Avoid damaging the posterior femoral cortex. **Be careful not to harm the popliteal neurovascular structures.**

Take out the Steinmann pins and then remove the chamfer cutting guide with the extractor. *(Fig. 55)*

**Note**

The chamfer cutting guide protects the mediolateral soft tissues by capturing the saw blade inside. Therefore it could be necessary to finish the cuts after removing the chamfer cutting guide.
Preparation of the tibia
Choose the size of the tibial template that best covers the proximal tibia. Attach the appropriate trial insert (fixed) or trial mobile bearing CR (mobile) to the tibial template. The size of the trial insert (fixed) or trial mobile bearing CR (mobile) is based on the size of the selected femoral component (S, S+, M, M+, L, L+) and the height of the extension and flexion gaps (10.0/12.5/15.0 or 17.5 mm).

Mounting the trial insert (fixed) on the tibial template
1. Insert the adaptor for the tibial template into the trial insert (fixed).
2. Attach the trial insert (fixed) to the tibial template. (Fig. 56)
Place the trial insert (fixed) and the tibial template on the tibial resection surface. (Fig. 57)

Mounting the trial femur on the femoral clamp
1. Open both arms of the femoral clamp by pressing the levers (K) and position the trial femur in the center. (Fig. 58)
2. Release the two levers (K) gently to fit the arms in the notches on the trial femur.
3. Turn the handle clockwise to fix the trial femur with the femoral clamp. (Fig. 59)
The trial femur is now ready for placement on the distal femur.
Use the femoral clamp to mount the trial femur of the appropriate size on the femur. (Fig. 60) Uncouple the femoral clamp from the trial femur by turning the handle counterclockwise and pressing the levers together.

Use the impactor to drive in the trial femur until optimal bone contact is achieved. (Fig. 61)

**Setting tibial rotation**

The trial tibia is automatically rotated into the final position by repeated flexion and extension of the knee joint while maintaining optimum tibia coverage. (Fig. 62/63)

*Note*

A high anterior resistance can occur when driving in the trial femur. This resistance can be overcome by applying extension force. This leads to a neutral position of the trial femur and prevents misplacement in flexion.

When using the mobile bearing, the tibial template can be fixed for best coverage without having to preset rotation.

*Note*

The movement of the prosthesis during flexion and extension can be checked at the same time.
Extend the knee and fix the tibial template with two Steinmann pins. (Fig. 64)
It is recommended to mark the position of the trial tibia with a cautery. (Fig. 65)
These markings represent an additional aid for positioning the definitive tibia component. This is especially useful for a cemented tibia component.

Remove the trial insert (fixed) or trial mobile bearing (mobile) and the trial femur.

Place the tibia impactor guide on the tibial gauge, assemble it with the modular handle and drill the peg holes (Ø 5 mm). (Fig. 66)

Then drill the central tibial stem hole with a Ø 18.5 mm step drill until it can not go any further. (Fig. 66)

Prepare the tibial stem fins by compressing the cancellous bone with the tibial impactor. (Fig. 67)

Remove the Steinmann pins, the tibia impactor guide and the tibial template.

Note
If necessary the tibial template can be further stabilized with two additional Steinmann pins.

Note
Ensure that the locating pin on the tibial impactor is inserted into the posterior channel of the tibia impactor guide!
Preparation of the Anchoring Stem (optional)

Preparation of the anchoring stem
The following steps must be followed if the surgeon decides to use an anchoring stem.

Insert the tibial drill sleeve into the tibia reamer guide. (Fig. 68)

Choose the tapered tibial reamer according to the desired stem length.

Ream the tibial medullary cavity by pushing the reamer until it can go no further. (Fig. 69)

Note
Take the length of the anchoring stem into consideration when determining the posterior slope of the tibial cut to avoid that the tip of the anchoring stem contacts the anterior tibial cortex.

Note
Perform the reaming by hand. This allows for better control when reaming the tibial canal.
Positioning the Trial Components

**Trial assembly**

**Assembly of the trial tibia plateau with the tibial clamp**

A special tibial clamp is provided for simple and accurate positioning of the trial tibia plateau.

1. Position the tibial adaptor (labeled: “Tibial Trial”) in the hole of the trial tibia plateau and fix with the screw (L). ([Fig. 70])

2. Then slide the tibial clamp from the front onto the tibial adaptor. ([Fig. 71])

3. Fix the tibial clamp by tightening the screw. ([Fig. 72])

The trial tibia plateau is now ready for insertion.

Depending on the type of prosthesis, insert the mobile or fixed bearing in the following manner:

**Type CR**

Insert the corresponding guide pin into the trial meniscal bearing (mobile). Ensure that the guide pin is correctly aligned (ANT). Then insert both parts into the trial tibia plateau. ([Fig. 73])

**Type UCOR**

Insert the trial mobile bearing into the trial tibia plateau. ([Fig. 74])

**Type FIX**

Insert the adapter for the tibial gauge (S, M, L) into the trial insert (fixed). Insert the trial fixed bearing into the trial tibia plateau. ([Fig. 75])

**Note – Combination**

The combination matrix are displayed on page 44 (mobile) and 45 (fixed).
Mount the trial femur of the corresponding size on the femoral clamp. (Fig. 76)

Use the impactor to drive in the trial femur until optimal bone contact is achieved. (Fig. 77)

**Note**
A high anterior resistance can occur when driving in the trial femur. This resistance can be overcome by applying an extension force. This leads to a neutral position of the trial femur and prevents misplacement in flexion.

Checking flexion and extension
Check flexion, extension and stability of the knee joint with the trial components. (Fig. 78/79)
Preparation of the Patella (optional)

**Procedure when preserving the patella**

It is not necessary to replace the patella due to the anatomical nature of the trochlear recess of the femoral component.

If the surgeon decides not to replace the patella, it is recommended to perform a patellaplasty. The patellaplasty consists of the following steps:

- Perform circumferential denervation of the synovial edge of the patella with electrocautery
- Remove peripheral osteophytes to restore the patella to its normal shape and size.

Be careful not to damage tendon insertions on the bone. Measure the thickness of the patella with the caliper. Determine the amount of bone that should remain after resection by subtracting the implant thickness from the patella thickness.

**5. Procedure for patella replacement/universal saw guide technique**

Apply the universal patellar saw guide in line with the patellar tendon. Push the patella up between the jaws of the saw guide. Level the patella within the saw guide jaws and use the thumbscrew to tighten the guide.

Resect the bone across the top of the saw guide jaws. (Fig. 80) The amount of bone removed should be approximately the same on all sides. Check that the 10 mm gauge does not rotate beneath the anterior surface of the patella. (Fig. 80) If the gauge hits the anterior surface of the patella, this indicates that at least 10 mm of bone stock will remain after the resection. It is recommended that a minimum of 12 mm of bone remain to allow sufficient bone stock for the implant peg.

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Perform patella resection in one level resulting in a smooth surface. (Fig. 81)

**Drilling the anchoring hole**
Determine the size for the patella component with the trial patella. Place the trial patella on the resected patella surface. Place a Steinmann pin in the hole of the trial patella to mark the center of the resected patella. (Fig. 82)

Remove the Steinmann pin and the trial patella. Then drill a hole in the resected patella without pressure until the stop. (Fig. 83)

Check the tracking of the patella during flexion and extension with the trial components in situ.

Remove the trial components if the outcome is satisfactory.

Deflate the tourniquet and do careful hemostasis.
Assembly of the Anchoring Stem (optional)

Assembly of the tibial component and the anchoring stem
Use the extractor to remove the Zimmer® X-Ray PMMA plug from the tibial component. (Fig. 84)

It is mandatory to use the extractor provided in the set to remove the Zimmer X-Ray PMMA plug from the tibial component. This avoids damaging the surface of the taper.

Note
Position the tibial component in the holder of the support block before removing the Zimmer X-Ray PMMA plug.

Note
Parts made of Zimmer X-Ray PMMA can not be resterilized.

Place the tibial component upside down on the support block and attach the anchoring stem to it. (Fig. 85)

Insert carefully by hand the anchoring stem into the taper of the tibial component.
Mount the impactor on the end of the anchoring stem to fix the stem in the taper. The driving mechanism must be activated by pressing down on the rear part of the instrument. This procedure must be repeated 3 times. A clearly audible sound should be produced each time. (Fig. 86)
Insert the tibial component into the corresponding notch of the support block, then secure it with the fixation screw (M). (Fig. 87)

Insert the stem fixation screw with the adaptor (Fig. 87) and torque wrench into the tibial component. The fixation screw is included in the anchoring stem package.

Tighten the fixation screw with the torque wrench (appropriate torque 12 Nm). (Fig. 88)
Implantation

Final implantation
Use the same procedure for implanting the definitive prosthetic components as for the trial components (see page 34).

Different tibial adaptors are used depending on the type of bearing. (Fig. 89)

For the mobile tibial base plate, the tibial adaptors are labeled as follows:
- Without anchoring stem: “Tibial Impl. without stem”
- With anchoring stem: “Tibial Impl. with stem”

For the fixed tibial base plate, the tibial adaptors are labeled as follows:
- Without anchoring stem: “Tibial Impl. without stem – FIX”
- With anchoring stem: “Tibial Impl. with stem – FIX”

Note
Ensure that the correct rotational position is maintained while driving in the tibial base plate. The fins on the conical base stem and the anterior cautery markings can be used as references. This is achieved by lining up the fins on the stem with the anterior markings previously done with electrocautery in the tibia.
Depending on the type of prosthesis, insert the mobile or fixed bearing in the following manner:

**Type CR (Cruciate Retaining – mobile bearing)**
Insert the corresponding guide pin into the trial mobile bearing. Ensure that the guide pin is correctly aligned (ANT). Then insert both parts into the tibial base plate. *(Fig. 92)*

**Type UCOR (Ultra-Congruent-Only-Rotating – mobile bearing)**
Insert the guide pin (supplied with the UCOR PE) into the mobile tibial base plate. Then insert the mobile bearing onto the guide pin. *(Fig. 93)*

**Types FIXCR and FIXUC (Cruciate Retaining and Ultra-Congruent – fixed bearing)**
Drive the fixed bearing into the fixed tibial base plate using the impactors for PE inserts. Ensure that the fixed bearing engages correctly. Introduce the fixed bearing into the fixed tibial base plate posteriorly, and then fix anteriorly with the impactor. *(Fig. 94)*

**Note – UCOR**
When using a stem extension, no matter which size and thickness, the separately packaged guide pins must be used. The guide pins in the UCOR PE package can be disposed of.

**Note – Combination**
The combination matrix are displayed on page 44 (mobile) and 45 (fixed).
Ensure best possible prosthesis-bone contact with all resected surfaces when implanting the femoral component with the femoral clamp. (Fig. 95) Carefully remove the femoral clamp.

The femoral component must be fully seated in order to obtain a better press fit. This can be achieved by hitting the femoral impactor axially. (Fig. 96)

Cement the patella component into the central drill hole. Gently squeeze with the patella t-bar. Press until the cement has hardened and then remove the patella t-bar. (Fig. 97)

Notes on the cemented version
Apply cement in the cement pockets only under the mobile tibial base plate or under the fixed tibial base plate. Do not apply any cement in the area of the central tibial anchoring peg!

It is recommended not to implant the femoral component until the cement has hardened on the tibia.

However, if inserting both components in one cementing procedure, a very careful technique is important, because cement residues cannot be removed from the posterior recess. The advantage of inserting both components in one cementing procedure is that pressure is applied during polymerization of the cement.

Due to the fast setting of the cement it is recommended to insert the patella component as the first step or in a separate cementing procedure after having fixed tibial and femoral components.

Remove all excess cement and any loose cement particles to avoid damage to the surfaces of the prosthesis (third body wear).
CR (Cruciate Retaining) when retaining the posterior cruciate ligament
UCOR (Ultra-Congruent-Only-Rotating) when resecting the posterior cruciate ligament
### Combination Table of the Fixed Bearing System

**FIXCR (Cruciate Retaining) when retaining the posterior cruciate ligament**

**FIXUC (Ultra-Congruent) when resecting the posterior cruciate ligament**

#### Patellar component

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<th>Femoral component</th>
<th>Tibial insert</th>
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#### Femoral component

- **Femoral component**: left/right CR (1 cemented, 2 uncemented)
- **Tibial insert**: (FIXCR, FIXUC)

#### Tibial in var (FRCR/FIRC)

- **Tibial in var (FRCR, FIRC)**
- **Tibial met at back, fix cemented**

#### Patellar component

- **Patellar component**: (Gender Solutions™)
- **Femoral component**: left/right CR (1 cemented, 2 uncemented)

### Table

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<th>Tibial insert</th>
<th>Patellar component</th>
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### Diagrams

- **Patellar component**: (Gender Solutions™)
- **Femoral component**: left/right CR (1 cemented, 2 uncemented)
- **Tibial insert**: (FIXCR, FIXUC)

### Additional Information

- **Anchorage stem, conical**
- **Con:60 01.02001.470**
- **Con:60H 01.02001.471**
The Innex® Knee System is a CE marked product.
The Innex® Knee System is not available for sale in the USA.