

Distal Femoral Replacement

Surgical Technique: Fixed Hinge Tibia Option

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onkossurgical.com

Precision Orthopaedic Oncology

- ELEOS™ Limb Salvage Solutions
- My3D[®] Personalized Solutions
- GenVie® Regenerative Biologics



1 IMPLANT DESCRIPTION & INDICATIONS

1.1 Implant Description:

Onkos Surgical Specific Devices are manufactured in accordance with an approved prescription for a named patient. They **MUST NOT** be used for any other patient.

The patient name is detailed in the operation drawing supplied with the device/s.

The planned resection levels are indicated on the operation drawing supplied with the implant.

Before commencing surgery please refer to packaging insert for complete product information, including contraindications, warnings, precautions, and possible adverse effects/complications. Packaging Inserts are also available from Onkos Surgical.

1.2 Implant Intended Use:

Patient Specific Implants are indicated for limb sparing and skeletal restorative procedures where radical resection and replacement of the distal femur is required with the following conditions:

- Patients suffering from severe arthropathy of the distal femur that do not respond to any conservative therapy or better
 alternative surgical treatment
- Surgical intervention for severe trauma, revision arthroplasties, failed previous prostheses and/or oncology indications; and malignant diseases (e.g. osteogenic sarcoma).

Onkos Surgical Patient Specific Devices and their components are for single use only.

1.3 Indications:

- Primary bone tumors
- · Secondary tumors arising in bone
- · Non-neoplastic conditions affecting the shafts of long bone
- Failed joint replacements
- · Failed massive replacements

Onkos Surgical Specific Devices are indicated for cemented and cementless procedures where radical resection and replacement of the distal femur is required.

1.4 Contraindications:

- Absolute contra-indications include:
- Infection and sepsis.

Relative contra-indications include:

- Long delay between manufacturing and insertion of a patient specific implant may result in significant mismatch due to possible changes in bone geometry
- · Inadequate or incomplete soft tissue coverage
- · Uncooperative or unwilling patient or patient unable to follow instructions
- · Foreign body sensitivity. Where materials sensitivity occurs seek advice with respect to testing
- Obesity
- · Vascular disorder, neuromuscular disorders or muscular dystrophy.

1.5 Patient Selection:

Factors to be considered:

- · Resection of neoplastic or diseased bone
- At risk from pathological fracture
- · Pain relief and improved function
- · Ability of patient to willingly follow instructions and undergo rehabilitation

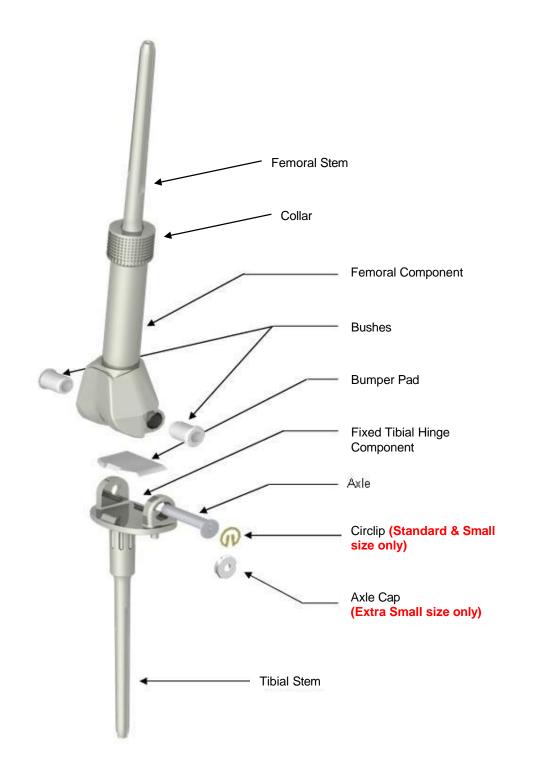
Onkos Surgical Specific Devices are available for use with the following tibial configuration options:

- Rotating hinge Passive tibia
- Passive fixed hinge tibia
- Rotating hinge polyethylene tibia
- Rotating hinge Metal cased tibia
- Fixed hinge tibia

The rotating hinge polyethylene tibial option is suitable for routine cases, the rotating hinge metal cased tibial option is suitable for extra-articular resection or difficult revisions and the fixed hinge tibial option is suitable for knees with marked instability.

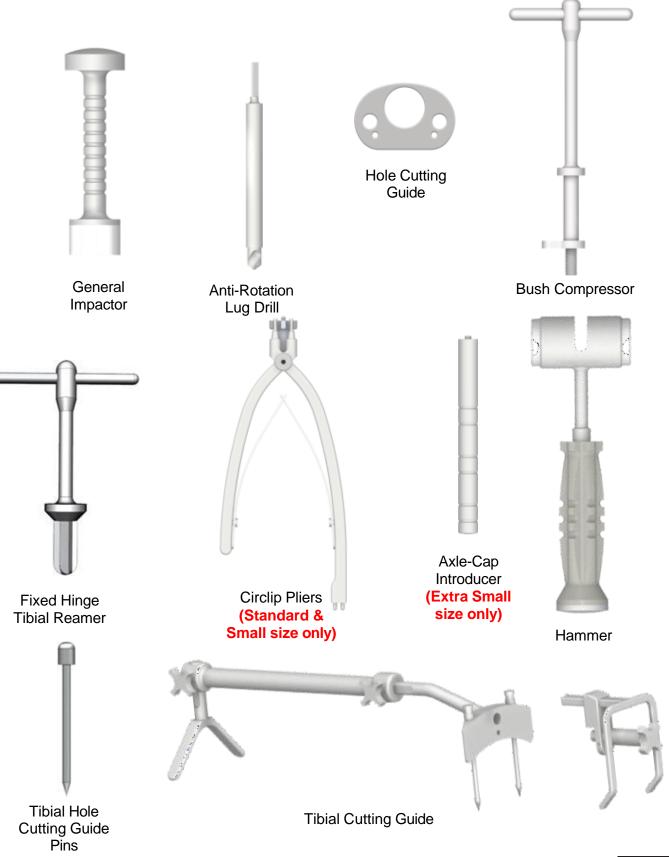


2 EXPLODED VIEW





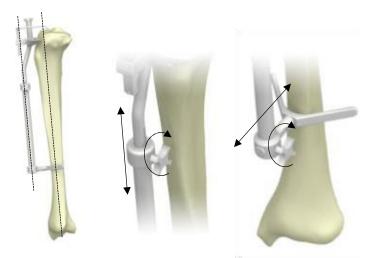
INSTRUMENTS



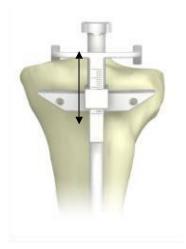


4 TIBIA PREPARATION





Adjust prongs of tibial cutting guide to sit on the plateau of the tibia Align cutting guide to be parallel with tibia and secure adjustments using locking screws



Adjust the reference cutting face to match the depth indicated on the operation drawing



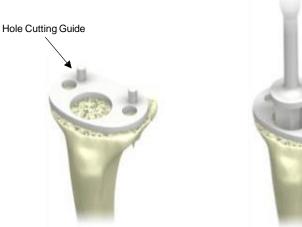
Secure the reference cutting face using bone pins and remove prong assembly



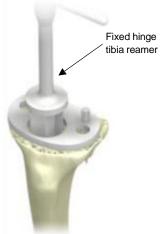
Trim tibial plateau in line with plane of reference cutting face



5 TIBIAL INSERTION



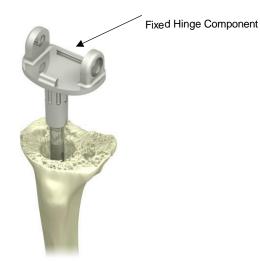
Secure cutting guide with pins, aligning straight edge with posterior of tibia



Ream tibial canal using fixed hinge tibia reamer



Use AR Lug drill to make two 10mm deep holes as indicated by cutting guide



Remove cutting guide and cement in fixed hinge tibia

Note: If optional plateau plate is to be used, it should be cemented with a thin layer of cement to the underside of the tibial component prior to impacting tibial component into tibia



6 BUMPER PAD & BUSHES



Insert bumper pad, aligning curve d anterior edge under anterior rail of tibial component





Use general impactor to clip posterior edge of bumper pad into place

On the femoral component, insert bushes into either end as far as possible

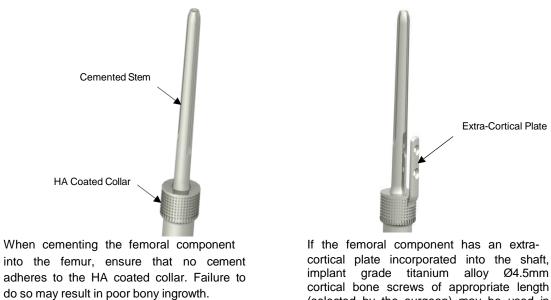




If not seated fully, use bush compressor to fully seat the bushes Slide in from one side and screw in nut until bushes are fully seated



7 FEMORAL INSERTION



It is advised to use a cement restrictor where possible.

cortical bone screws of appropriate length (selected by the surgeon) may be used in the screw holes.



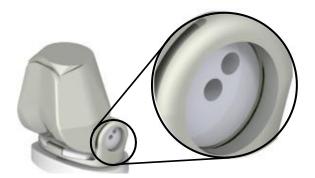
8 AXLE INSERTION



Align femoral and tibial component. Insert axle from either side



Using pointed implement, rotate axle to ensure offset head fits into recess within tibial component



Ensure offset head is seated within recess and not trapped in circlip groove



9 CIRCLIP INSERTION

(Standard & Small Size Only)



Align opening of circlip with prongs of circlip pliers Push circlip onto pliers, locating central locating pin into centre of circlip. Ensure prongs of pliers are oriented as shown



Engage circlip by approaching at an angle, from same side as axle insertion



Squeeze the circlip pliers to close the circlip. Insert circular

edge within groove in tibial component



Straighten out to engage circlip into groove

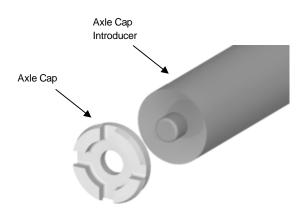


Once within groove, release grip on pliers to secure circlip within groove. Rotate circlip within groove, to ensure correct fit.



10 AXLECAP INSERTION

(Extra Small Size Only)



Orientate axle cap with flanges away from axle cap introducer



Press axle cap firmly onto introducer



Insert axle cap by approaching from same side which axle was inserted



Press in fully until flanges click into groove on tibial component







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