

JTS Non-Invasive Extendible Distal Femoral Replacement

Surgical Technique: Rotating Hinged Polyethylene Tibia Option



973.264.5400

onkossurgical.com

Precision Orthopaedic Oncology

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



1 IMPLANT DESCRIPTION & INDICATIONS

1.1 Implant Description:

The Onkos Surgical JTS Extendible Distal Femoral Implant is manufactured in accordance with an approved prescription for a named patient. It **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:

The JTS Extendible Distal Femoral Implant is 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).

JTS Extendible Distal Femoral Implant and its 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

The Onkos Surgical JTS Extendible Implant is 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

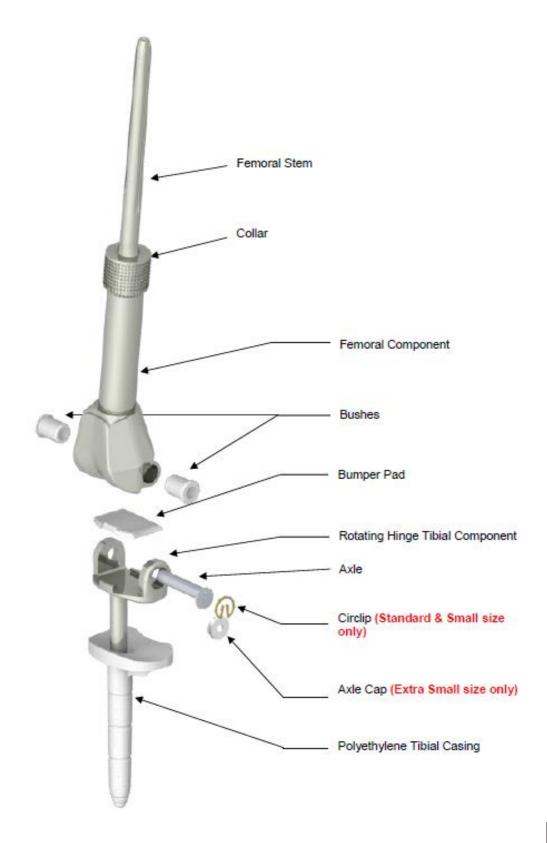
The JTS Extendible Distal Femoral Implant is available for use with the following tibial configuration options:

- · Passive rotating hinge tibia
- Passive fixed hinge tibia
- · Rotating hinge polyethylene tibia
- 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.

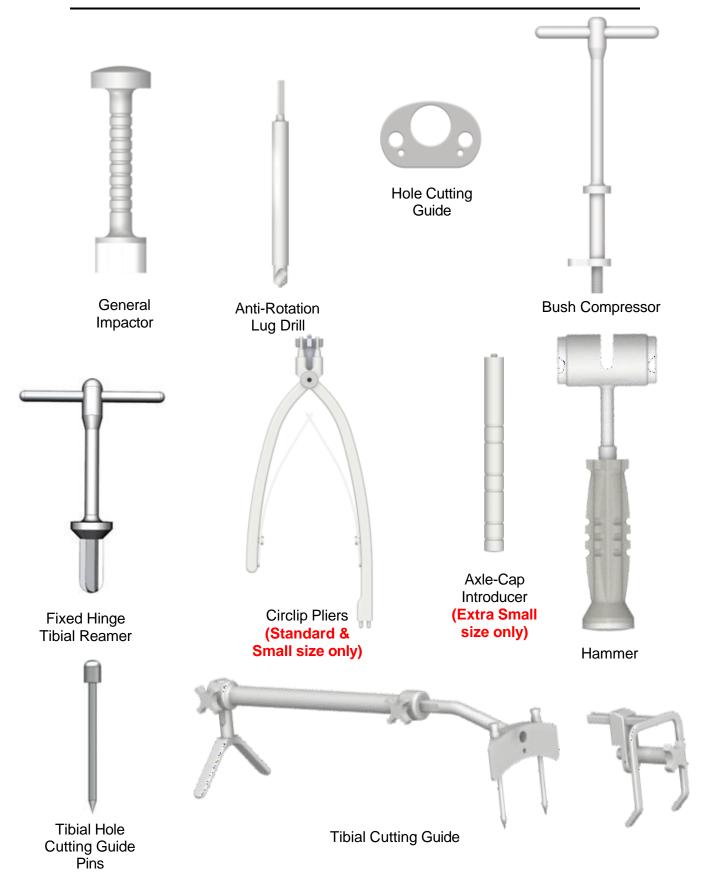


2 EXPLODED VIEW





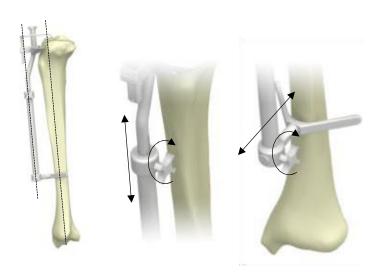
3 INSTRUMENTS



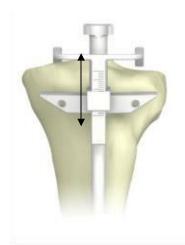
4 TIBIA PREPARATION



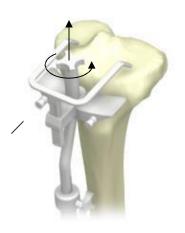
Adjust prongs of tibial cutting guide to sit on the plateau of the tibia



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



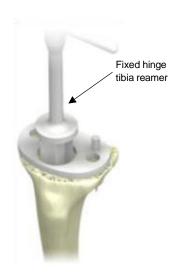
Trim tibial plateau in line with plane of reference cutting face



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



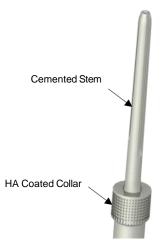
If not seated fully, use bush compressor to fully seat the bushes



Slide in from one side and screw in nut until bushes are fully seated

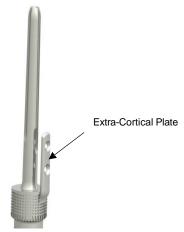


7 FEMORAL INSERTION



When cementing the femoral component into the femur, ensure that no cement adheres to the HA coated collar. Failure to do so may result in poor bony ingrowth.

It is advised to use a cement restrictor where possible.

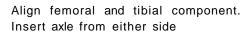


If the femoral component has an extracortical plate incorporated into the shaft, implant grade titanium alloy Ø4.5mm cortical bone screws of appropriate length (selected by the surgeon) may be used in the screw holes.



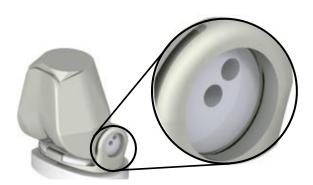
8 AXLE INSERTION







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 o 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

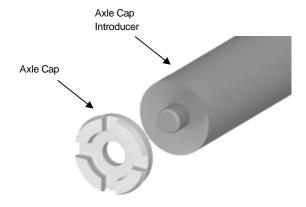
Straighten out to engage circlip into groove

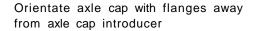
Once within groove, release grip on pliers to secure circlip within groove. Rotate circlip within groove, to ensure correct fit.



10 AXLECAP INSERTION

(Extra Small Size Only)







Press axle cap firmly onto introducer



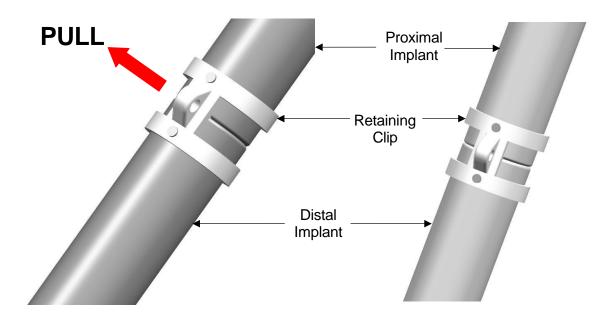
Insert axle cap by approaching from same side which axle was inserted



Press in fully until flanges click into groove on tibial component



11 GROWER CLIP REMOVAL



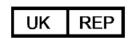
Once implant has been fully inserted and secured with axle and circlip, or axle and axle-cap, remove retaining clip located on the femoral component. Failure to do so will result in malfunction of the device and may lead to future surgical intervention.







Onkos Surgical 77 East Halsey Road Parsippany, NJ USA 07054 +1-973-264-5400 www.onkossurgical.com



IdealMed Ltd, Unit A2, Beech House Oaklands Office Park, Hooton Cheshire, CH66 7NZ, United Kingdom

All trademarks are the property of **ONKOS** SURGICAL unless otherwise indicated. A surgeon should rely exclusively on his or her own professional medical/clinical judgment when deciding which particular product to use when treating a patient. **ONKOS** SURGICAL does not prescribe medical advice and advocates that surgeons be trained in the use of any particular product before using it in surgery. A surgeon must always refer to the package insert, product label and/or instructions for use before using any **ONKOS** SURGICAL product.

JTS DFPRPT ST 06.13.23 v0

©2023 ONKOS SURGICAL. All rights reserved.

