Surgical Technique
INDICATIONS

The indications for use of the total hip replacement prostheses include:
• Noninflammatory degenerative joint disease, including osteoarthritis and avascular necrosis;
• Rheumatoid arthritis;
• Correction of functional deformity;
• Revision procedures where other treatments or devices have failed; and,
• Nonunions, femoral neck fractures, and trochanteric fractures of the proximal femur with head involvement that are unmanageable using other techniques.

Additional indication specific to use of ACCOLADE II Femoral Stems with compatible Howmedica Osteonics Constrained Liners:
• When the stem is to be used with compatible Howmedica Osteonics Constrained Liners, the device is intended for use in primary or revision patients at high risk of hip dislocation due to a history of prior dislocation, bone loss, soft tissue laxity, neuromuscular disease, or intra-operative instability.

ACCOLADE II Femoral Stems are intended for cementless use only and are intended for total and hemiarthroplasty procedures.

CONTRAINDICATIONS

• Active infection or suspected latent infection in or about the hip joint;
• Bone stock that is inadequate for support or fixation of the prosthesis;
• Skeletal immaturity;
• Any mental or neuromuscular disorder that would create an unacceptable risk of prosthesis instability, prosthesis fixation failure, or complications in post-operative care; and
• Obesity. An overweight or obese patient can produce loads on the device that can lead to failure of the fixation of the device or to failure of the device itself.

WARNINGS AND PRECAUTIONS

See package insert for warnings, precautions, adverse effects and other essential product information.

Before using ACCOLADE II instrumentation, verify:
• Instruments have been properly disassembled prior to cleaning and sterilization;
• Instruments have been properly assembled post-sterilization;
• Instruments have maintained design integrity; and,
• Proper size configurations are available.

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For cleaning and sterilization instructions of surgical instruments, refer to the instrument package insert.
The ACCOLADE II Femoral Hip System is a femoral stem that is intended for cementless, press-fit application. The proximal region of the stem is coated with PureFix HA over a commercially pure titanium plasma spray substrate. The Morphometric Wedge — an evolution of the tapered wedge — is characterized by its variable, size-specific medial curvature. The ACCOLADE II Femoral Hip System is suitable for various surgical approaches, including direct anterior.

The total system includes:
- 12 body sizes ranging from size 0 to size 11
- Two anatomic offset options for each size

The stem is designed for use with Stryker V40 femoral heads and their compatible acetabular components.

The ACCOLADE II Hip System includes a broach-only instrumentation platform that accommodates all surgical approaches and has been streamlined for surgical efficiency.

Surgical Technique

Stryker Orthopaedics wishes to thank the following orthopaedic surgeons for their expertise in the development of the ACCOLADE II System Surgical Technique:
- Dr. Richard Rothman
- Dr. Dermot Collopy
- Dr. David Jacofsky
- Dr. Frank Kolisek
- Dr. Art Malkani

This publication sets forth detailed recommended procedures for using Stryker Orthopaedics devices and instruments. It offers guidance that you should heed, but, as with any such technical guide, each surgeon must consider the particular needs of each patient and make appropriate adjustments when and as required.
Pre-operative planning aids in the determination of probable implant style and size. The pre-operative planning process should take qualitative and quantitative factors (including patient bone quality, density, and morphology) into consideration in order to evaluate and select the appropriate instrument/implant system for the patient.

Place an acetabulum template over the area on the x-ray (Figure 1). Be sure that the cup is well centered within the acetabulum and the size fills between the tear drop and the superior rim. After templating the acetabulum for size, mark the center of rotation (represented by the blue dot).

The ACCOLADE II has two offset options: the standard offset 132° neck angle and the high offset 127° neck angle. Choose the ACCOLADE II template for which the stem size achieves medio-lateral cortical engagement at the proximal two-thirds of the stem and recreate the desired leg length and offset (Figure 2). For both the 132° and 127° offset options, the template has markings that indicate the center of the femoral head for a range of head offset options.

The predicted change in leg length and offset is determined by the relative positioning of the center of rotation markings on the femoral and acetabular components. For example, if a given femoral component center of rotation marking is superior to the center of rotation marking of the acetabular component, leg lengthening is predicted. The desired change in leg length is determined by the radiographic leg length inequality that was previously determined. If 8mm of leg lengthening is required in order to equalize the leg lengths, the center of rotation marking of the femoral component should be positioned 8mm superior to the center of rotation marking of the acetabular component. The stem size and head offset that most closely meets this goal is chosen.

The predicted change in offset is also considered by comparing the relative medial/ lateral position of the center of rotation markings of the femoral and acetabular components. The templates should be used to estimate the final components that most closely restore the normal offset of the patient’s hip.

Once the final estimated stem size and position is determined, the neck resection level should be noted (Figure 3). This will be used as a reference during intra-operative neck resection.

**Tip**

Richard Rothman, M.D.
Templating is an important aspect of pre-operative planning, but it should only serve as a guide. Final decision making concerning fit, size, and soft-tissue tensioning occurs in the operating room.

**Tip**

David Jacofsky, M.D.
The templated stem size and neck resection level may have to be adjusted if leg length cannot be managed solely by changing the head offset or neck angle.

**Templates**

ACCOLADE II Acetate Templates Lit# LTEM105
Acetate templates include 20% magnification.
A proper neck resection level directly affects the final placement and fit of the femoral stem. By using the anatomic landmarks referenced during pre-operative x-ray templating, the pre-planned neck resection is made with an oscillating saw and with the aid of the Neck Resection Guide. The guide helps the surgeon to determine the correct stem orientation and placement. After careful pre-operative templating, the guide is placed on the anterior/posterior aspect of the exposed proximal femur and the planned femoral neck cut is marked using a marking instrument of choice. Care should be taken to align the body of the guide with the axis of the femoral canal (Figure 4).
To help ensure proper final orientation of the stem, lateral bias during implant preparation is preferred. Retraction of the gluteus medius and removal of the lateral cortical bone at the piriformis insertion will permit true axial introduction of the instruments and implant. The Modular Box Osteotome or a rongeur can be used to remove bone from this area (Figure 5).

**Tip**

Art Malkani, M.D.
Remove the lateral cortical bone at the piriformis fossa to obtain ideal proximal fit and to minimize the risk of undersizing and/or varus placement of the femoral component.
The ACCOLADE II Hip System is a broach only system. While use of an axial starter reamer is needed, use of cylindrical reamers is not necessary to prepare the femoral canal. The Axial Starter Reamer is used with the T-Handle to open the femoral canal and to aid in determining the orientation of the femoral axis. The tapered design allows for access to the canal and is graduated along the flutes, which helps provide a reference during insertion into the canal. Advance the Axial Starter Reamer into the femoral canal to a depth at which the first graduation mark is aligned with the medial aspect of the neck resection (Figure 6).

Slight lateral pressure on the reamer during operation will aid in preparing the femoral canal in the neutral orientation of the implant.

**Note**

Care should be taken not to sink the starter reamer below the first graduation mark to allow for proper press fit of the implant.
**STEP 4**

**BROACHING**

Broaching is then performed beginning with the size 0 broach. The broach should be oriented to the long axis of the oblong shape created by the neck resection (Figure 7).

**Instruments**

- **Offset Broach Handle**
  - 1020-1460
- **ACCOLADE II Broach**
  - Size 0-11
  - 1020-52xx
- **Mallet**
  - 1120-1000

**Tip**

*Art Malkani, M.D.*

During broaching, place lateral bias on the broach to fill the proximal femur and minimize undersizing the implant.

**Note**

ACCOLADE II broaches can be properly identified in several ways.
1. The broach size is engraved into a square pocket on both the anterior and posterior sides of the broach.
2. The size is marked on the top of the broach post.
3. The distal lateral tip of the broach is polished.
4. 1020-52xx part number

**Note**

Accolade and ACCOLADE II broaches cannot be used interchangeably.

**Figure 7**

Sequentially broach upward in size until the proper size is achieved. The surgeon’s clues to a firm fit and final size include 1.) changing pitch of sound that results from mallet blows to the broach handle; 2.) increased resistance to forward advancement; 3.) lack of further motion.

Two grooves on the anterior and posterior surfaces of the broach act as a point of reference to help the surgeon visualize the broach advancing into the femur (Figure 8).
Relying only on the neck resection height alone for final seating height may lead to improper sizing and inadequate component fixation.

The final broach should seat firmly against medial and lateral cortical bone (Figure 9).

**Tip**

*Frank Kolisek, M.D.*

If the broach and/or stem stops above or below the original neck resection, leg length and soft-tissue tensioning must be re-assessed during intra-operative trialing. Do not force the broach and/or stem to the resection level. This will increase the risk of proximal femur fracture.

Generally, if a broach sinks below the level of the neck resection, advance to the next larger broach. If the neck resection is deemed higher than desired, remove the broach and perform a new neck resection at a lower level.

**Figure 9**

For good fixation of the implant, it is important that axial alignment of the broach is maintained at all times in the canal. Generally, the broach rotation is self-aligning.

Upon reaching the final size and depth of the broach, detach the broach handle from the broach, leaving the broach fully seated in the femoral canal.

**Tip**

*Frank Kolisek, M.D.*

Pre-operative templating serves as a guide. Don’t keep hitting the broach harder just because you have not reached the size that was templated.

**Figure 10A**

Assemble the Calcar Planer to the reamer power adaptor. Guide the Calcar Planer over the broach post ensuring the Calcar Planer is axially aligned with the post and is stable. Initiate power prior to contacting the femur and slowly advance the Calcar Planer toward the broach using continuous power until the positive stop on the Calcar Planer contacts the broach face and the bone is removed (Figure 10A). Failure to operate the Calcar Planer in accordance with these instructions may result in damage to the femur.

**Figure 10B**

In the event that the Calcar Planer cannot fully engage the broach post (Figure 10B), remove the broach and perform a new neck resection at a lower level. Alternatively, a larger broach size should also be considered.
STEP 5
TRIAL REDUCTION

**TIP**
Richard Rothman, M.D.
Start with a standard offset (132°) neck trial. If the leg lengths are equal, but soft tissues are lax, move to a 127° degree neck trial.

Select a Neck Trial which has the same base neck length and angle as the planned implant size. This can be determined in two ways.

1. Match the color indicator located on top of the Neck Trial taper to the color indicator on top of the broach (Figure 11).

2. The table below indicates the correct neck length for each size stem and the corresponding color code. The size of the broach directly corresponds to the size of the implant.

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<td>0, 1</td>
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<td>10, 11</td>
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Figure 11

**Instruments**

132° Neck Trial 1020-32xx
127° Neck Trial 1020-27xx
ACCOLADE II Broach Size 0-11 1020-52xx
Assemble the Neck Trial onto the broach. Next, assemble a V40 Head Trial onto the Neck Trial (Figure 12). Femoral heads come in multiple offsets and are different for each femoral head implant material (see table at right.) For this reason, final head material should be chosen prior to trial reduction. Offsets add or subtract from the base neck length of the implant and help to achieve the desired leg length and offset.

Perform a trial reduction of the hip. Upon confirmation of the selected components, remove the trial head and trial neck, and reassemble the broach handle to the broach. Remove the broach from the femoral canal. The final broach size determines the correct implant size.

### Tip

**Richard Rothman, M.D.**
Remember, pre-operative templating serves as only a guide. Final sizing, leg length, and offset are determined intra-operatively.

### Tip

**Dermot Collopy, M.D.**
After trial reduction and prior to removing the broach, tap on the broach a few more times to see if it will advance. Due to the visco-elastic properties of the bone, the bone could relax during the trialing process. If the broach does advance, re-trial and make adjustments accordingly.

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Head Trials with an “R” suffix are made from a radiopaque material, making them visible on an x-ray.
STEP 6

IMPLANTING THE STEM

OPTION 1
Thread the Modular Threaded Stem Inserter into the drive hole on the proximal face of the stem (Figure 13). The inserter should be fully threaded and secured to the stem prior to impaction to prevent damage to the threads on the implant or the instrument. Using the inserter, the stem should be inserted into the femoral canal until it stops.

Tip
Dermot Collopy, M.D.
If the stem hangs up due to impingement of the Modular Threaded Stem Inserter against the overhanging tip of the greater trochanter, remove the threaded stem inserter leaving the stem in place and use the Modular Stem Impactor to fully seat the stem.

OPTION 2
The ACCOLADE II Femoral Stem can also be inserted using the Modular Offset Quick Connect Stem Inserter. Place tip of the inserter into the drive hole of the stem taking care to align the version tab on the inserter with the slot in the stem (Figure 14). The quick connect design provides the inserter with a stable spring connection, but it does not provide a mechanical lock. Therefore, this assembly should be handled with care, as excessive shaking or motion may result in the stem disassociating from the inserter.

Note: The Modular Offset Quick Connect Stem Inserter cannot be used with the Size 0 and Size 1 ACCOLADE II. Misuse could lead to instrument failure.

OPTION 3
The ACCOLADE II Femoral Stem can also be inserted by hand and then impacted into the bone using the Modular Stem Impactor. The Modular Stem Impactor has a spherical tip, which is placed onto the drive hole of the stem. This instrument allows for off-axis impaction of the stem. The Modular Stem Impactor does not connect to the stem, and, therefore, can only be used for final impaction of the stem.

Instruments

| Modular Threaded Stem Inserter 1020-1800 | Modular Offset Quick Connect Stem Inserter 1020-1860 | Modular Stem Impactor 1020-1870 |

Figure 13
A Mallet is then used to seat the stem into the canal (Figure 15).

The surgeon should NOT attempt to continue impacting the femoral component if visual and auditory clues indicate that the stem is firmly seated in the canal. These clues, rather than the broach seating level, should be used to determine the final seating height of the implant. Continued aggressive impaction could lead to femoral fracture. In the event that dense bone is encountered intra-operatively and compounding anatomical factors are present, the seating of the implant may not be consistent with the level of the broach due to the viscoelastic nature of the femoral bone.* If the final seating height is undesirable, the implant can be removed and additional broaching can be performed. If the stem inserter is contacting the greater trochanter during insertion, continued impaction could lead to a fracture.

Prior to final head assembly, neck length / head offset selection may be re-evaluated using a V40 Head Trial. Place the Head Trial onto the stem neck taper and reduce the hip. Leg length equality and proper soft tissue tension are evaluated. Remove the Head Trial and dry the implant trunnion with a laparatomy sponge or sterile towel.

Select the appropriate corresponding V40 Femoral Head (CoCr, Alumina Ceramic, BIOLOX delta Ceramic) or sleeve and place it onto the dry trunnion of the femoral stem with a slight twist. Impact the head with two moderate impactions using the Modular Head Impactor (Figure 16).

Verify the head is secure on the trunnion after head impaction by applying traction to the head and confirming stability on the trunnion. If necessary, the head can be removed utilizing the head disassembly instrument.*

Relocate the femoral head into the acetabular cup and re-check the hip biomechanics. The surgical site is then closed according to surgeon preference.

* If a ceramic head is placed on the trunnion and then removed, it must be replaced with a V40 cobalt chrome head or a V40 Titanium Adapter Sleeve (17-0000E) and a C-Taper ceramic head.

If the stem must be removed, utilize the Modular Threaded Stem Inserter.

Note

In no instance should any attempt be made to pre-assemble the adaptor sleeve inside the BIOLOX delta Universal Ceramic Head.

Intra-operatively assemble the BIOLOX delta Universal Taper Ceramic Head onto the sleeved femoral stem and set with two moderate strikes using the Head Impactor (6626-0-140). Care must be taken to avoid excessive impact forces when assembling the Ceramic Head to the sleeved femoral component.

### OPTIONAL STEP

When selecting a BIOLOX delta Universal Taper Ceramic Femoral Head (6519-1-0xx) for implantation, use of a Universal Adaptor Sleeve is necessary.

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After completing the trialing process, intra-operatively assemble the adaptor sleeve to the femoral stem manually. The Universal Adaptor Sleeve must be fully seated on the stem taper before the head is assembled.
### ACCOLADE II

#### IMPLANT INFORMATION

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ACCOLADE II SURGICAL TECHNIQUE

INSTRUMENTATION

1020-9001 Basic Tray
1020-9000 Single Tray Case

1020-2727
127°, 27mm Neck Trial

1020-3227
132°, 27mm Neck Trial

1101-2200
Orthonomic T-Handle

1020-2730
127°, 30mm Neck Trial

1020-3230
132°, 30mm Neck Trial

1020-2735
127°, 35mm Neck Trial

1020-3235
132°, 35mm Neck Trial

1020-1111
Calcar Planer - Standard

1020-1112
Calcar Planer - Large

1020-1100
Neck Resection Guide

1020-1460
2 x Offset Broach Handle

Optional
1440-1460
2 x Straight Broach Handle

1120-1000 Mallet
1020-2737 127°, 37mm Neck Trial
1020-2740 127°, 40mm Neck Trial
1020-3237 132°, 37mm Neck Trial
1020-3240 132°, 40mm Neck Trial

6264-8-028R 28mm -4mm V40 Head Trial
6264-8-928R 28mm -2.7mm V40 Head Trial
6264-8-128R 28mm +0mm V40 Head Trial
6264-8-228R 28mm +4mm V40 Head Trial
6264-8-328R 28mm +8mm V40 Head Trial
6264-8-428R 28mm +12mm V40 Head Trial
6264-8-032R 32mm -4mm V40 Head Trial
6264-8-632R 32mm -2.5mm V40 Head Trial
6264-8-132R 32mm +0mm V40 Head Trial
6264-8-232R 32mm +4mm V40 Head Trial
6264-8-332R 32mm +8mm V40 Head Trial
6264-8-432R 32mm +12mm V40 Head Trial
6264-8-036R 36mm -5mm V40 Head Trial
6264-8-436R 36mm -2.5mm V40 Head Trial
6264-8-136R 36mm +0mm V40 Head Trial
6264-8-236R 36mm +4mm V40 Head Trial
6264-8-336R 36mm +5mm V40 Head Trial
6264-8-736R 36mm +7.5mm V40 Head Trial
6264-8-338R 36mm +10mm V40 Head Trial

1601-1700 Modular Head Impactor
1601-1210 Modular Box Osteotome
1020-1460 Broach Handle
ACCOLADE II SURGICAL TECHNIQUE

INSTRUMENTATION

1020-9002 ACCOLADE II Broach Tray
1020-9000 Single Tray Case

1020-2900 Orthonomic Modular Handle
1020-1800 Modular Threaded Stem Inserter
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**Alternate Instrument Tray Configuration**

- **1020-9102** General Tray
- **1020-9000** Single Tray Case
- **1020-9101** Accolade II Tray
- **1020-9000** Single Tray Case

Refer to ACCII-B-1 for tray layouts
Stryker Orthopaedics has validated the following reusable instrument trays with Aesculap’s SterilContainer™ System and with CSR wrap. Refer to LSTPI-B (Instructions for Cleaning, Sterilization, Inspection, and Maintenance of Reusable Medical Devices).

6147-0-100 Universal Lid  
6147-1-101 Accolade II Basic Tray  
6147-1-102 Accolade II Broach Tray