Radiographic Imaging of Hip Replacement Hardware

Catherine C. Roberts, MD,* and Felix S. Chew, MD, MBA†

Hip joint replacement surgery has been the most common type of elective and semi-elective orthopedic operation for nearly 25 years. As the science and engineering of implants have evolved, their appearances on imaging and the frequency of common complications have also evolved. Our objectives in writing this article are to explain the postoperative appearance of hip joint replacement hardware in terms of materials, design, and function, and to review the imaging appearance of common complications. Our emphasis will be on radiographic imaging because that modality remains the mainstay of follow-up and evaluation.

Indications for Hip Replacement

For most joints, the indication for replacement is articular disease involving that joint, most commonly osteoarthritis and rheumatoid arthritis. In these cases, the goal of joint replacement is twofold: first, the restoration of joint function, and second, the elimination of arthritic pain. Joint replacement has been proven to be highly efficacious in meeting these goals for most patients with arthritis.1-3 At the hip, additional common indications include femoral head disease, such as osteonecrosis, and femoral neck fractures, such as displaced subcapital fractures, where the risk of posttraumatic osteonecrosis is high.2-4 In the latter circumstance, hip replacement may not really be elective, as the morbidity and mortality from hip fractures in elderly patients who are not treated with hip replacement are both high. Patients with congenital hip disease can benefit from hip replacement.3 A hip replacement may also be one component of a larger reconstruction following tumor treatment.4

Materials Used in Hip Replacement

The principal material used in joint replacements is metal. The types of metal that are commonly used for implant components are pure titanium and titanium alloys (titanium–aluminum–vanadium), and cobalt–chromium alloys. Obviously, all metal components are radiopaque on radiographs, but titanium is less dense than cobalt–chromium, and both are less dense than lead. (The atomic numbers are 22 for titanium, 24 for chromium, 27 for cobalt, and 82 for lead.) The exact appearance on radiographs will depend on the radiographic technique. Titanium alloys are more commonly used in implants of recent vintage, but since the average life of a joint implant easily exceeds 10 years, there are many patients being followed who have cobalt–chromium implants. Stainless steel or titanium may be used for screws and washers.

Bone cement is commonly used to fix components to bone or to fill voids in bone to provide a closer fit with components. Bone cement is an acrylic plastic (polymethylmethacrylate) that is polymerized in situ at the time of implant. Because plastic is a hydrocarbon with the same primary atomic constituents as fat (carbon and hydrogen), it is radiolucent in radiographs. However, bone cement is typically mixed or premixed with barium powder to render it radiopaque, and thus it is radiopaque on radiographs, but less dense than metal. Bone cement can be used as both an adhesive as well as a space-filler.

Polyethylene is commonly used as a bearing surface, so that most total joint replacements have a convex metal component articulating with a concave polyethylene liner. The type of polyethylene used in joint replacements is not the common, pliable material used in kitchenware or other light consumer products, but it is an ultrahigh molecular weight material (molecular weight of 3 to 6 million Daltons) that is also used for bullet-proof vests and as the lining for skating rinks and ship docks. A high degree of crosslinking provides increased strength and resistance to surface abrasion.

Ceramics have become more common as prosthetic femoral heads and prosthetic acetabular bearing surfaces. The two materials in use are Zirconia,5 a strong material with an elastic modulus similar to metals that is more widely known for faux jewelry, and alumina, a hard and rigid material that is more widely known as an ingredient for antacids.

Fixation of Hip Replacements

Fixation of components to bone may occur through direct mechanical fixation, passive interference fit, bone cement,
and porous ingrowth. Methods of direct mechanical fixation such as screws are generally obvious on radiographs. Passive interference fit or press fit components are held in position by the shape of the components and the space into which they are tightly fitted. Bone cement may be used as an adhesive, literally gluing the component to bone, or it may be used to fill spaces and contribute to a closer interference fit. Porous ingrowth fixation is based on the principle that remodeling bone can attach itself directly to the component, holding it in place. The key to porous ingrowth is providing a surface that bone can grow into. Special surface treatments that have been used include layers of small metallic spheres, fine wire mesh, and hydroxyapatite crystals. On radiographs, it is difficult if not impossible in most cases to identify the specific type of surface treatment.

**Types of Hip Replacements**

There are two general types of hip replacement: hemiarthroplasty and total hip arthroplasty. The clinical indications for each type are different, and these general types can be identified from radiographs.

Hemiarthroplasty is an operation that replaces one surface of a joint. In the case of the hip, it is always the femoral side that is replaced, so that the resulting articulation is between a prosthetic femoral head and the native acetabular cartilage.

![Figure 1](image1.png) Austin–Moore unipolar hemiarthroplasty. Femoral head and neck are replaced with single-piece prosthesis that articulates with native acetabulum. This prosthesis is loose, as seen by lateral tilt of prosthesis tip (black arrow) and local lucency at prosthesis–bone interface along proximal, lateral aspect of remaining native femur (white arrowheads).

![Figure 2](image2.png) Bipolar hemiarthroplasty. Two-piece prosthesis replacing femoral head and neck articulates with native acetabulum. Unlike unipolar arthroplasty, bipolar prosthesis has an additional region of motion between prosthetic neck and prosthetic head. Similar to the first case, this prosthesis is loose, with lateral tilt of the prosthesis tip (black arrow) and lucency at cement–bone interface along the proximal, lateral aspect of the remaining native femur (white arrowheads).

![Figure 3](image3.png) Femoral head resurfacing arthroplasty. Damaged portions of femoral head have been removed and femoral head remodeled to accept a thin metal cup as a new articulating surface.
Hemiarthroplasty at the hip is generally performed for femoral head disease, not hip joint disease. Common clinical indications for hip hemiarthroplasty include osteonecrosis of the femoral head (before secondary osteoarthritis has developed), displaced subcapital fracture of the femoral neck, and resection for tumor. Hemiarthroplasty can be recognized from radiographs because the native acetabular cartilage will be present. One surgical strategy for hemiarthroplasty is replacement of the femoral head and neck by a one-piece (unipolar) prosthesis that is inserted into the medullary canal of the proximal femur (Fig. 1). In this operation, the femoral head and neck are resected and the prosthesis is secured to the proximal femur by cement or press fit. A more advanced type of hemiarthroplasty is called the bipolar hip replacement, in which a two-piece prosthesis is inserted into the medullary canal of the proximal femur (Fig. 2) following removal of the femoral head and neck. The bipolar prosthesis has a femoral stem and a separate head, so that two articulations result: an articulation between the prosthetic head and the native acetabulum, and an articulation between the two components of the prosthesis. Both the unipolar and the bipolar arthroplasty are commonly used today. Recent studies recommend the use of unipolar prostheses instead of bipolar prostheses in elderly patients with femoral neck fractures. A third type of hemiarthroplasty is one in which the articular surface of the femoral head is replaced by a metal cup (Fig. 3). This is referred to as a cup arthroplasty or a femoral head resurfacing arthroplasty. Arthroplasties such as these have been in use since the 1920s.

A total hip arthroplasty (THA) is an operation that replaces both surfaces of the hip joint. The clinical indication for total hip arthroplasty is typically hip joint disease such as osteoarthritis or rheumatoid arthritis. Patients with osteonecrosis are more commonly treated with hemiarthroplasty but total hip arthroplasty is also an effective treatment. Specifically, patients with Ficat grade III osteonecrosis have a better clinical outcome when treated with a total hip arthroplasty rather than with a hemiarthroplasty.
Figure 6  Hybrid total hip arthroplasty. Femoral component is cemented, while acetabular component is cementless.

Figure 7  Ceramic femoral head portion of THA. Ceramic head is more radiolucent than remainder of metallic femoral and acetabular components.

Figure 8  Metal-on-metal total hip arthroplasty. There is no polyethylene liner to separate femoral and acetabular components.

Figure 9  Surface replacement arthroplasty. Damaged portions of acetabulum and femoral head have been removed and lined with metal cups.
than a bipolar arthroplasty. Because both surfaces of the joint are replaced, the resulting articulation is always between the femoral and acetabular components of the prosthesis. Total hip replacements can be recognized from radiographs by the presence of both femoral and acetabular components. The most common type of total hip replacement is one in which the femoral head and neck are replaced

<table>
<thead>
<tr>
<th>Table 1 Potential Complications of Total Hip Arthroplasty</th>
</tr>
</thead>
<tbody>
<tr>
<td>Early</td>
</tr>
<tr>
<td>Nerve palsy: sciatic, femoral, peroneal</td>
</tr>
<tr>
<td>Hemarthrosis</td>
</tr>
<tr>
<td>Thromboembolism: deep venous thrombosis, pulmonary embolism</td>
</tr>
<tr>
<td>Vascular injuries</td>
</tr>
<tr>
<td>Hemorrhage and hematoma formation</td>
</tr>
<tr>
<td>Bladder infection</td>
</tr>
<tr>
<td>Limb length discrepancy</td>
</tr>
<tr>
<td>Fracture</td>
</tr>
<tr>
<td>Late</td>
</tr>
<tr>
<td>Loosening</td>
</tr>
<tr>
<td>Stem failure</td>
</tr>
<tr>
<td>Osteolysis (small particle disease)</td>
</tr>
<tr>
<td>Heterotopic ossification</td>
</tr>
<tr>
<td>Stress fractures: femur, pubis</td>
</tr>
<tr>
<td>Fracture of hardware</td>
</tr>
<tr>
<td>Implant sarcoma</td>
</tr>
<tr>
<td>Either</td>
</tr>
<tr>
<td>Infection of joint</td>
</tr>
<tr>
<td>Dislocation</td>
</tr>
<tr>
<td>Fractures: femur, acetabulum, pubic ramus</td>
</tr>
</tbody>
</table>

Figure 10 Acetabular reconstruction cage. Acetabulum has been revised with cage device to correct protrusio deformity, thus restoring anatomic positioning for femoral component of hip replacement.

Figure 11 Modular hip prosthesis. Custom length prosthesis replaces proximal femur, which was resected for osteosarcoma.

Figure 12 Periprosthetic fracture. Subtle longitudinal fracture line is seen extending distally beyond tip of femoral component (arrowheads). This fracture was noted at time of surgery and was stabilized with Cerclage wires.
by a prosthesis that fits inside the medullary canal of the proximal femur, and the acetabulum is replaced by a metal cup with a polyethylene liner. The articulation is thus metal on polyethylene. The components may be fixed to bone using cemented or cementless methods (Figs. 4 and 5). The combination of a cemented femoral component with a noncemented acetabular component is called a hybrid THA (Fig. 6). Some total hip replacements use ceramic-bearing surfaces (Fig. 7), and recently, the use of metal-on-metal hip prosthesis has become more widespread in the United States (Fig. 8). Total hip replacement by resurfacing both the acetabular and the femoral sides of the joint may also be done. When both the femoral and the acetabular articular surfaces are reshaped and lined by thin metal cups, the procedure is referred to as a surface replacement arthroplasty or a “double cup” arthroplasty (Fig. 9).

Specialized techniques and hardware may be used in circumstances where the native bone stock is insufficient for standard techniques. These circumstances include revisions of failed arthroplasty, developmental hip dysplasia, tumor resection, and severe protrusio acetabuli in rheumatoid arthritis. Autograft or allograft bone may be used to buttress the acetabulum or the proximal femur. Autograft bone is commonly derived from the femoral head and neck that have been resected during the course of hip arthroplasty. Allograft bone is typically obtained from a bone bank and is usually obtained from cadavers. Acetabular reconstruction cages are components that are used to reconstruct acetabular beds that are deficient in bone stock (Fig. 10). Patients with severe protrusio acetabuli from conditions such as rheumatoid arthritis or patients with marked bone loss from a failed acetabular component are typical candidates for acetabular reconstruction. Following the acetabular reconstruction, a standard total hip prosthesis may be implanted. When the proximal femoral shaft is resected along with the femoral head and neck, a modular prosthesis can be used for reconstruction. The proximal end of this prosthesis may be a unipolar, bipolar, or total hip arthroplasty, while the distal portion that replaces the missing segment of femoral shaft consists of modules customized for the individual patient. The distal portion has an intramedullary stem that fits into the medullary canal and is cemented into place (Fig. 11).

Complications

Complications related to hip replacements are mostly related to fractures, dislocations, component loosening, component failure, and infection. They may occur early during the hospital admission or years after the surgery (Table 1). Fractures of the proximal femoral shaft may occur as a complication of insertion of the stem of the femoral component during surgery, particularly if it is a press fit component. Most of these fractures are non-displaced and are often recognized at the time of surgery. Cerclage cables can be used to secure the fracture fragments (Fig. 12). In some cases a nutrient artery canal near the tip of a femoral stem can be confused with fracture. Comparison with preoperative imaging can help differentiate between an acute fracture and a vascular groove.
Periprosthetic fracture after surgery is most often secondary to prosthesis loosening, occurring at the tip of the femoral endoprosthesis. Fractures occurring after sports-related injuries are rare, likely due to the more sedentary nature of the population receiving the implants. Periprosthetic fractures often require revision of the hip arthroplasty.

Dislocations are more common in total hip replacement than in hemiarthroplasty and occur typically between the femoral component and the acetabular component. The proximal femur tends to rotate externally, and the head of the prosthesis dislocates anteriorly (Fig. 13). Recurrent dislocations often indicate malposition of components that require revision. Dislocation of the hip during transfer from the operating table to the gurney for transport to the recovery room may occur as an early complication. Postprocedure radiographs that are typically performed in the recovery room can exclude this possibility and document the immediate postsurgical appearance of the hip.

Loosening of components may occur from the cyclic mechanical loading that occurs during weight bearing or may result from biologic action. As implanted, joint replacement materials are biologically inert. However, repetitive mechanical stress on the surface of bearing surfaces causes the abraison of small particles. These particles are released from the artificial joint surfaces into the joint capsule, where they may incite a foreign body granuloma response. The particles that have been particularly implicated in this granulomatous response are particles of polyethylene. This granulomatous response results in the resorption of bone at its interface, and this resorption may be visible on radiographs as osteolysis if it is extensive enough (Fig. 14). Osteolysis may be evident as a region of lucency around a component at its interface with bone; this may be seen at the cement–bone interface in the case of a cemented prosthesis or at the metal–bone interface in the case of a noncemented prosthesis. Locations of osteolysis surrounding arthroplasty components can be described using the seven femoral zones described by Gruen and three acetabular zones (Fig. 15). As the zone of osteolysis extends along the interface between bone and component, the component loses its mechanical attachment. Once gross motion is possible, the zone of osteolysis may become progressively larger, and often a fine zone of sclerosis may form at the margins of the cavity. This process may occur slowly over several years. In general, a zone of osteolysis greater than 2 mm around a component is abnormal and suggests loosening from foreign body granulomatous reaction. Any gap be-

Figure 14 Osteolysis and polyethylene wear. Femoral component is eccentrically located in acetabular cup, indicating wear of polyethylene liner. Wide regions of lucency surround acetabular component and proximal femoral component likely secondary to a granulomatous response to polyethylene particles.

Figure 15 Periprosthetic zones used to describe location of osteolysis or complication.
between metal and cement is abnormal. The presence of osteolysis does not necessarily correlate with clinical symptoms, and clinical symptoms are the usual indication for revision, not radiographic appearance. Because polyethylene particles have been implicated in osteolysis, research has focused extensively on how to reduce the generation of these particles. The results of such research have included improved forms of polyethylene with greater resistance to abrasive wear as well as implants that do not use polyethylene. These include implants with ceramic-on-ceramic bearing surfaces as well as metal-on-metal bearing surfaces (Fig. 8).

Infections may occur in the immediate postoperative period or at a subsequent time. There are no specific radiographic features that indicate early infection, but dislocation may be associated with large effusions. In some cases, intra-articular foci of air, beyond the immediate postoperative state, suggest the presence of infection (Fig. 16). Periosteal reaction can also suggest infection. Imaging-guided arthrocentesis is the best radiological method for identifying infection. In most cases, this may be performed under fluoroscopy. Following informed consent, the patient is placed on the fluoroscopy table in the supine position. The hip capsule will extend from the native bone of the acetabulum to the native bone of the proximal femur, encompassing all of the exposed components in between (Fig. 17). The easiest target to hit is the prosthetic femoral neck, right in the middle. Following localization of this target under fluoroscopy and preparation of the skin with cleansers and local anesthetic, an 18- or 20-gauge spinal needle can be advanced through the soft tissues until it touches metal. At that point, fluid within the joint may be aspirated. If no fluid is aspirated, a small amount of contrast may be instilled into the joint to confirm location. Washing the joint with an injection of 10 mL of preservative-free saline followed by aspiration will provide material for microbiological analysis, but cultures of fluid from saline wash when the initial aspiration is dry are rarely positive. Because the majority of injected fluid will accumulate posteriorly, it is generally possible to re-aspirate only 1 to 2 mL. Flexing the hip while aspirating may improve the yield. The definitive method for identifying infection and the causative agents is considered to be surgical biopsy of the synovium, a procedure typically beyond the scope of the radiologist. It is sometimes possible to obtain a small piece of synovium using a 22-gauge Sure-Cut (Bauer Medical, Clearwater, FL) biopsy needle placed through the spinal needle. This small piece of tissue can be sent for culture and sensitivity.

Confirmed infections require surgical debridement and prolonged antibiotic therapy. Because it is difficult to achieve the requisite concentrations of antibiotics when they are...
given systemically, methylnmethacrylate beads that have been impregnated with antibiotics can be placed directly into the surgical bed for better results (Fig. 18). In most cases, the prosthesis itself must be removed (or retrieved as the surgeons refer to it). With the prosthesis gone, patients are not able to bear weight on the affected extremity. To prevent soft-tissue contractures and preserve the limb for possible reimplantation, spacers fabricated from polymethylnmethacrylate may be placed into the joint bed. The PROSTALAC® (Prosthesis of Antibiotic-Loaded Acrylic Cement) device is a temporary implant embedded with antibiotics tailored to the infecting agent (Fig. 19). This implant acts to help treat the infection and to maintain function of the hip between removal of the infected hardware and revision of the hip when the infection has resolved. Infections may require treatment for many months before definitive reconstruction can be performed. The ultimate outcome of infected hip replacements may be reconstruction with another prosthesis, surgical arthrodesis of the joint, or permanent pseudoarthrosis of the joint (Fig. 20).

Heterotopic bone may form in the surgical bed following total hip replacement. Although small amounts of heterotopic bone formation are unimportant, large amounts of heterotopic may interfere mechanically with motion at the hip and require resection (Fig. 21). Heterotopic bone formation appears to be more common in patients who have other evidence of excessive bone formation such as hypertrophic osteoarthritis or diffuse idiopathic skeletal hyperostosis. Therapeutic irradiation of the surgical bed following surgery may prevent recurrence of heterotopic ossification.

Figure 18 Antibiotic beads. Polymethylmethacrylate beads are impregnated with antibiotics and placed at site of infection, in this case along side plate with dynamic compression screw fixating subtrochanteric fracture.

Figure 19 PROSTALAC implant. This temporary antibiotic-impregnated implant is placed after removal of infected hardware.

Figure 20 Permanent pseudoarthrosis (Girdlestone procedure) of left hip joint following infection. Femoral head has been removed, without replacement.
Fracture of the greater trochanter may occur during surgery or as a delayed complication. Because these fractures displace widely, they are virtually impossible to reduce and fix, so they are generally left ununited (Fig. 22). The gluteus medius, one of the strong abductors of the hip, inserts on the greater trochanter. Its loss of action results in a permanent limp.

Approach to Radiographic Reporting

Radiographs in follow-up of hip replacements have two general functions, documentation of the procedure and the condition of the joint, and surveillance for complications. Joint reconstruction surgeons typically follow their patients several times in the first year following an implant, and then annually after that. At the time of the first perioperative set of radiographs, it is appropriate to identify the type of prosthesis and to search for immediate complications or problems. Because these radiographs are typically obtained using portable equipment in the recovery room, their quality may be substandard. At our institutions, AP pelvis and true lateral hip radiographs are obtained, allowing assessment of the placement of the components. On the AP radiograph, one can assess the orientation of the acetabular cup in the coronal plane of total hip replacements. The angle between a line drawn across the opening of the acetabular cup and a horizontal reference line drawn tangential to the ischial tuberosities is called the lateral opening angle and typically measures between 30 and 50° (Fig. 23). On the true lateral projection, the angle between a line drawn against the opening of the acetabular cup and a vertical reference line is called the degree of anteversion and typically measures between 10 and 15° (Fig. 24). The position of the center of rotation of the
prosthesis should be anatomic, and limb length should be symmetric with the other side. In some patients who have a THA subsequently revised, an extended trochanteric osteotomy\textsuperscript{28,29} may be necessary to remove the original femoral component. It is important not to confuse the osteotomy with a fracture (Fig. 25).

At the time of subsequent follow-ups, one should note the results of a comparison with the initial postoperative examination and search for evidence of complications. In the early follow-ups, one should look specifically for fractures; in subsequent follow-ups, it is important to search for the complications described above, including asymmetric polyethylene wear (Fig. 14), osteolysis, loosening, heterotopic ossification, subsidence (Fig. 26), textured coating disintegration (Fig. 27), and stress-related changes (Fig. 28). If there has been no change in appearance and there is no evidence of complications, it is sufficient to simply state that there has been no change in appearance and that there is no evidence of complications.

Table 2 summarizes the radiographic analysis of every total hip arthroplasty.

---

**Conclusion**

Hip replacement surgery has been one of the most common elective and semi-elective orthopedic operations for the past 25 years. Radiography remains the mainstay of imaging evaluation of hip replacements. The various types of hip replacements can be recognized on radiographs, and many complications have specific appearances that the radiologist should be familiar with.

**Acknowledgements**

We extend special thanks to William W. Daniel, MD, and F. Spencer Chivers, MD, for contributing cases.

Table 2 Radiographic Analysis of Total Hip Arthroplasty

<table>
<thead>
<tr>
<th>Question</th>
<th>Answer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Is the entire prosthesis imaged?</td>
<td></td>
</tr>
<tr>
<td>Check for acetabular version</td>
<td></td>
</tr>
<tr>
<td>Check for polyethylene wear. Is femoral head centrally located within acetabular prosthesis?</td>
<td></td>
</tr>
<tr>
<td>Check for loosening</td>
<td></td>
</tr>
<tr>
<td>Widening of the lucent zone at the cement-to-bone interface &gt;2 mm</td>
<td></td>
</tr>
<tr>
<td>Widening of the lucent zone at metal-to-bone interface &gt;2 mm</td>
<td></td>
</tr>
<tr>
<td>Migration of components from their original positions</td>
<td></td>
</tr>
<tr>
<td>Development of a lucent gap between metal and cement</td>
<td></td>
</tr>
<tr>
<td>Cement fracture</td>
<td></td>
</tr>
<tr>
<td>Periosteal reactive bone</td>
<td></td>
</tr>
<tr>
<td>Osteolysis</td>
<td></td>
</tr>
</tbody>
</table>

Figure 27 Textured coating of implant is disintegrating and free particles can be seen in joint space.

Figure 28 Stress fracture. Laterally located periosteal reaction along femoral shaft, between tip of THA and total knee arthroplasty, indicates region of stress fracture.