A Mid- to Long-Term Follow-Up Experience with a Specific Metal-on-Metal Total Hip Arthroplasty Design

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Abstract

Purpose Metal-on-metal (MoM) total hip arthroplasty (THA) has been a subject of recent discussion and concern due to the early failures caused by local and systemic adverse reactions related to specific designs. The aim of this study is to analyze the outcomes and survival rates of a single brand of MoM implants implanted in a consecutive series of patients at a single institution.

Methods Between 2007 and 2012, 116 (118 hips) patients were evaluated at a mean follow-up of 6.6 years after primary THA. The diagnosis leading to surgery was osteoarthritis (80 patients) and proximal femoral fracture (36 patients). A single design of THA was implanted. All patients were evaluated before surgery and postoperatively at 1, 3, 6, and 12 months by clinical scores and radiographic studies. The data analysis was made using Student’s t-test.

Results The minimum follow-up was of 4 years, with a mean follow-up of 6.6 years. Two aseptic loosening of the acetabular component were recorded (one per group), which were not associated with local or systemic complications related to metal ion release. Both were revised by an isolated acetabular cup substitution with metal-on-polyethylene couplings. Nonprogressive radiolucent lines < 2 mm in zone 2 were observed in other six patients around the acetabular component without clinical manifestation (four in the arthritis group and two in the fracture group). Postoperative Harris Hip Score and SF-36 (36-Item Short Form Survey) score improved in both groups.

Conclusion Despite several MoM implants showing early complications and failures, a specific MoM design may be associated with good clinical results at a mid- to long-term follow-up.

Level of Evidence This is a therapeutic case series, Level 4 study.

Keywords ► metal-on-metal total hip arthroplasty ► total hip arthroplasty ► metal ion release ► hip arthritis ► proximal femoral fractures

Introduction

Total hip arthroplasty (THA) represents the most successful orthopaedic procedure. Over the last decades, improvements in the surgical technique, prevention of perioperative complications, postoperative pain management, and dedicated rehabilitative protocols have produced better clinical outcomes as compared with the past. Today, THA is a fully reproducible operation with several indications not only limited to primary osteoarthritis (OA) but also to other hip alterations such as secondary OA, hip dysplasia, or fractures.1–4 Two of the most important fields of interest related to THA are the choice of the coupling and the properties of biomaterials.

“Hard” couplings such as ceramic-on-ceramic (COC) and metal-on-metal (MoM) have been enthusiastically introduced in the clinical practice for their extremely low rate of wear in terms of intra-articular particles release, as compared with the historical metal-on-polyethylene (MOP) coupling.5,6 Some concerns with MoM implants and
the potential effect of metal ions on local and systemic tissues arose but without substantial evidences.\textsuperscript{7,8} On the other hand, several biomechanical advantages were correlated with MoM implants: the use of larger heads improving hip stability and range of motion; the low percentage of aseptic loosening, given the less production of polyethylene debris; and the higher hardness compared with COC implants, in which some cases of brittleness were described.\textsuperscript{9}

However, the natural corrosion induced by the contact of metal bearings in the synovial fluid was thought to be correlated to an aspecific macrophage cell mediated local tissue reaction to the release of metal ions, leading to early aseptic loosening.\textsuperscript{10} Such condition was not well understood and considered until the recall of a specific implant (Articular Surface Replacement [ASR], DePuy Orthopaedics, Warsaw, Indiana, United States) in 2003. This risk was then potentially associated with all MoM implants. From that event until nowadays, there has been a perception (both from patients and surgeons) that the use of MoM implants should be discouraged because of the high probability of mechanical failure and also because of the theoretic harmful effect of metal ions in the periprosthetic tissues and systemic organs.\textsuperscript{11} Several models of the effect of the ions on joint tissues have been proposed (ALDVAI [aseptic lymphocyte-dominated vasculitis-associated lesion], LYDIA [lymphocyte-dominated immunological answer]).\textsuperscript{12,13} Moreover, cases of hypersensitivity to metals have been reported in patients with MoM implants either in stable or failed implants.\textsuperscript{7,14}

Simultaneously, the improvements in the manufacture of COC THAs, which are characterized by a superior hardness, a lower risk of brittleness, and advantages of an inert biomaterial, led to the increased use of such implants and a lesser consideration for MoM implants.\textsuperscript{15}

Even after these events, several researches on the good outcomes and long-term survivorship of other MoM implants have been published in recent years.\textsuperscript{16–19}

The aim of this study is to analyze the clinical outcomes and survival rates of a single model of MoM implants implanted in a consecutive series of patients at a single institution. The hypothesis of our study is that a single model of MoM implant used in our institute has led to good clinical outcomes and survival rates.

**Methods**

A total of 116 patients who consecutively underwent an MoM implant between January 2007 and March 2012 were prospectively studied. Of them, 74 were female patients and 42 male patients. The average age was 72 years (range: 63–89 years), and the mean body mass index was 25.8 (range: 23.0–28.5). In 59 cases, the right hip was involved, whereas in 55 cases, the left hip was involved. In two cases, a bilateral THA was performed. The diagnosis leading to surgery was OA in 80 cases and proximal femoral fracture in the remainder 36 cases. The exclusion criteria were patients not able to give consent to procedure and to follow-up, a different diagnosis from OA and femoral fracture, a contralateral implant with other bearings, and a referred hypersensitivity to metals.

The study based on the principles of the Helsinki declaration was approved by the Institutional Review Board, and all patients were informed regarding treatment and follow-up. All patients underwent a radiographic study (with a standing hip X-ray in cases of patients affected by OA) and a general evaluation by geriatricians (in case of femoral fracture). Harris Hip Score (HHS) and 36-Item Short Form Survey (SF-36) were evaluated preoperatively in all patients.\textsuperscript{20,21}

In 91 cases, a locoregional anesthesia was performed; in the remaining patients, a general anesthesia was performed. Two senior surgeons performed all surgeries by the same (direct lateral) approach and the same surgical technique. Two prosthetic designs belonging to the same brand (Wright Medical Technology, Memphis, Tennessee, United States) were used. In case of OA, Conserve with cementless cup and stem was implanted, whereas, in case of femoral fracture, Collegia with pressfit cup and cemented or cementless stem was used.

All patients underwent a deep venous thromboembolism (DVT) prophylaxis with low molecular weight heparin and a short-term antibiotic prophylaxis with cefazoline 2 g, as reported by the standard protocol of the authors’ institution. A postoperative mechanical intermittent compression of the lower legs was prescribed to all patients. After a mean period of 6.9 days (range: 5–10 days), all patients were discharged and sent to rehabilitative facilities to complete the functional recovery.

Follow-up visits were performed at 1, 3, 6, and 12 months, and then at a yearly interval, with standard X-rays, HHS, SF-36, and evaluation of the positioning of the components following the criteria of DeLee and Charnley for the cups and of Gruen et al for the stems.\textsuperscript{22,23} The acetabular inclination was calculated on X-rays following specific criteria,\textsuperscript{24} and the study of the periacetabular ossifications was conducted according to Brooker et al’s classification.\textsuperscript{25} A data analysis was performed to compare preoperative and follow-up data using paired Student’s t-test. Significance was set at $p < 0.05$.

**Results**

The mean follow-up was 6.6 years (range: 4–9 years). All patients were followed up for a period of at least 4 years. Eight (7%) patients were lost after the minimum follow-up.

No intraoperative complications were recorded. Early complications were recorded: six DVTs (four in OA patients, two in fractured patients), one superficial wound infection in an OA patient, and two cases of pneumonia (all in fractured patients). No cases of septic failure and no instability were recorded in the study population. Two aseptic loosening of cups (one in an OA patient, one in a fractured patient) were recorded 37 and 22 months postoperatively. Their inclination was 40 and 44 degrees, respectively. Both cups showed an abnormal rotation (excessive anteversion) and were thus revised by an isolated revision with jumbo cups fixed by screws and MOP couplings. In either case, no synovitis or bone alterations related to metal ions was found at histological analyses on intraoperative specimens (– Figs. 1 and 2).

The mean cup inclination in the non-failed implants was 42.9 degrees (range: 42–49 degrees). The mean preoperative
value of HHS was 41.1 (range: 24–52) in the OA group, with progressive postoperative improvements up to a mean score of 88.4 (range: 28–100) at the latest follow-up ($p<0.05$). In the fracture group, a mean preoperative HHS of 22 (range: 10–31) and a final mean score of 87.6 (range: 34–100) were recorded ($p<0.05$) (►Fig. 3).

Regarding the SF-36, in the OA group, the mean preoperative value was 23.7 (range: 15–40) for Physical Component Summary (PCS) and 34.5 for Mental Component Summary (MCS) (range: 23–50). At the latest follow-up, the mean value was 42.2 (range: 38–52) for PCS and 50.5 for MCS (range: 39–53) ($p<0.05$) (►Fig. 4). In the fracture group, the mean preoperative value was 12.7 (range: 9–18) for PCS and 33.1 for MCS (range: 30–39). At the latest follow-up, the mean value was 43.3 (range: 37–48) for PCS and 49.8 for MCS (range: 46–53) ($p<0.05$) (►Fig. 5).

Nonprogressive radiolucency lines < 2 mm in zone 2, according to DeLee and Charnley, were observed in four OA patients and two fractured patients. Furthermore, no osteolysis was found at follow-up, except in the two revised cups. No significant radiolucent lines or osteolysis in all stems were found following the criteria of Grue et al.

Finally, we found stage 2 ossifications, according to Brooker et al, in four OA patients and three fractured patients and stage 3 in other four patients (two of both groups), however, without complaints by the patients. All these patients were administered indometacin 25 mg three times a day for 3 weeks.

**Discussion**

THA is a highly successful surgical procedure. The introduction of more performing materials led to great expectations in terms of wear reduction and survivorship of hip implants. Hard bearings such as COC and MoM couplings seemed to fulfill such characteristics. However, first-generation implants failed due to aseptic loosening or other mechanical issues. Particularly, the first MoM implants dramatically failed because of brittleness. Improvements in the manufacturing of materials and new designs of the MoM components were made in the third generation, which were characterized by large head diameters, increased stability, and better clearance, compared with the past.

Metals in contact with biologic fluids tend to corrode, releasing metal ions locally and toward the blood flow. This
remains a hot point of debate because there is a risk of hypersensitivity and potential toxic effects on specific tissue in sensitive patients.\textsuperscript{14,39–41} Until 2003, no actual reason to suspect any of these effects was strongly considered. The large number of early failures of a specific design, the ASR (DePuy Orthopaedics), first used as hip resurfacing implant and later as THA, had a worldwide high resonance. In a high percentage of cases, local bone reactions, pseudotumors, and synovitis were associated with the failure of such implants, with rates reported up to 68%.\textsuperscript{39,42} The main mechanism of failure has been related to several factors: the poor positioning of the cup, the design of the implant, and the susceptibility of patients.\textsuperscript{43} Surely, the excessive wear due to an altered release of metal ions is the main reason for the failure of ASR cups. Moreover, while a moderate-to-mild release of ions induces a local osteolysis at the bone/implant interface,
a high production of metal particles may be considered the cause of potential systemic toxicity represented by apoptosis, tissue necrosis, and genomic alterations. Patients with ASR presented elevated blood and urinary levels of chrome (Cr) and cobalt (Co), while patients with other types of MoM implants usually have a slight elevation of such values. However, for the latter implants, no symptoms or early failures have been addressed, as the experience reported in this series. Recently, patients with M2a acetabular MoM components (Biomet Orthopaedics, Warsaw, Indiana, United States) have shown high rates of pseudotumor at a mid-term follow-up, leading to revision in a percentage ranging from 4 to 8% and loosening in 31%; 13.8% of the studied hips had a definite adverse reaction to metal debris (ARMD), and 53.8% of the studied hips had a definite, probable, or possible ARMD. Swelling, clicking, pain, and sensation of subluxation were also reported by patients with this implant.

A strong debate revolves also around the potential hypersensitivity in patients with MoM implants. Some interesting studies revealed that well-functioning MoM implants may produce a hypersensitivity to metals, whereas loose implants show a risk ranging from 50 to 60%. Patients with MoM implants have higher values of urinary and blood concentrations of CrCo than other couplings. As workers exposed to metals are prone to develop malignant tumors, it is reasonable to consider patients with MoM implants at high risk for cancer. However, to date, no study has highlighted such improved risk. According to the latest studies, the intrasynovial fluid concentration of CrCo in patients with MoM implants reflects blood concentrations of these ions. Nevertheless, despite contrasting opinions, there is no clear evidence that urinary or blood concentrations of CrCo ions are strictly related with the survivorship.

On the other hand, MoM implants showed excellent mid-to-long-term outcomes. Early revisions were substantially necessary only in ASR implants. Beyond the inner properties of MoM cups, other factors such as the proper positioning of the acetabular component, adequate components lubrication, and a good clearance are crucial for the correct functioning and long survivorship of the implant. As a matter of fact, acetabular cups inclination > 50 degrees have been associated with poor outcomes. In our experience, the mean inclination of approximately 43 degrees witnesses such a critical aspect in the final results.

This study has some limitations. The population is not numerically consistent. No blood or urinary samplings were harvested, and consequently no study has been performed on the hypothetic metal ion release. However, almost all patients were followed-up for a mid- to long term, and no one complained of symptoms to be related to metal hypersensitivity or other toxic effects. Thus, we did not have any need to perform laboratory assays. Furthermore, satisfactory outcomes and a midterm survivorship were reported.

In conclusion, modern MoM implants have theoretic advantages compared with other couplings. Unfortunately, some specific design has shown dramatic failures due to their abnormal metal ion release, associated with local bone alterations and theoretical systemic effects. Despite these events, most of the implanted MoM implants are well functioning up to now, however, mostly because of the refusal of this option expressed by patients.
Conflict of Interest
None declared.

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