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Surgical management of hip osteoarthritis

Rajiv Gandhi MD MS, Anthony V. Perruccio PhD, Nizar N. Mahomed MD ScD

O steoarthritis is a leading cause of pain, disability and health care use among adults. The hip is the second most common large joint affected by osteoarthritis.¹⁻³ Although research has advanced our knowledge of osteoarthritis, no therapies currently exist that halt progression of the disease. In many cases, the disease progresses to damage and destruction of the joint. Consequently, orthopedic surgery has a critical role in the management of osteoarthritis.

More than 30 000 hospital admissions for hip replacement and revision surgery were reported across Canada in 2008/09, a 63% 10-year increase.⁴ Aging of the population; increased longevity, arthritis prevalence and rates of obesity; and expanding indications for hip surgery portend a continuing upward trend in demand for surgical management of hip osteoarthritis.

Although there have been many advances in surgical techniques and approaches for hip osteoarthritis, debate continues on the optimal management for the individual patient. In this review, we discuss indications for surgery, review surgical approaches and component materials, and suggest future directions. We reviewed randomized clinical trials, meta-analyses, and prognostic, observational and retrospective studies (Box 1).

When should patients be referred for surgical assessment?

Referral for surgical assessment should be considered for patients who experience hip symptoms (e.g., pain, restricted function and stiffness) that substantially affect quality of life and are unresponsive to pharmacologic and nonpharmacologic treatments.^{5–7} Evidence suggests that early referral, before extensive functional limitation and pain, and early intervention are associated with better patient-reported pain and function following surgery (Table 1).^{8–15}

Although patient-specific factors, including age, sex, obesity and comorbidities, may variably influence patient-reported outcomes after surgery (Table 2),^{16–18} there is no suggestion that these factors should be barriers to referral, and

these factors are not used for wait-list prioritization. Even if certain subgroups fare less well after joint replacement, this does not mean that, on average, they do not receive benefit.¹⁸ Further, there is no consensus on the use of scoring tools or algorithms by which referral is based on a specific threshold being reached.

Glycemic control is critical because diabetes confers an increased risk of deep infection (relative risk 2.11, 95% confidence interval [CI] 1.41 to 3.17) after surgery.¹⁹ Also, cessation of smoking for 6–8 weeks before intervention has been shown to decrease wound complications (absolute risk reduction of 26%).²⁰ Among patients taking newer-generation antiplatelet agents such as clopidogrel, it is recommended that these medications be stopped 7 days before surgery, particularly if spinal or epidural anesthesia is being considered.²¹ Finally, hip replacement should be delayed 1 year after cardiac stent placement.²² Individual risk–benefit evaluation is essential.^{21,22}

Competing interests:

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Correspondence to: Nizar N. Mahomed, nizar.mahomed@uhn.ca

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Box 1: Summary of literature review

We performed a literature search of PubMed (1980 to January 2013), Embase (1980 to January 2013) and MEDLINE (1950 to January 2013) databases. We used a combination of Medical Subject Headings, including "hip replacement," "hip arthroplasty," "hip resurfacing," "metal-on-metal," "ceramic hip," "minimally invasive hip surgery," "mini-incision hip surgery," "hip replacement outcomes," "metal-on-polyethylene hip replacement," "revision hip replacement" and "meta-analysis." Reference lists of selected articles were also reviewed for additional studies. Two of us abstracted and reviewed all data. We included the best evidence, including clinical trials, meta-analyses, prognostic studies, observational studies and retrospective studies, as available.

Key points

- Total hip replacement with a metal-on-polyethylene bearing surface remains the "gold standard" for the treatment of end-stage hip osteoarthritis, providing reliable improvement in pain and function with consistent implant longevity.
- Registry reports suggest an increased rate of revision for hip resurfacing compared with total hip replacement, particularly among female patients.
- A further registry report showed an increased failure rate of metal-onmetal bearing surfaces compared with ceramic and polyethylene at 7 years.
- The outcomes of revision total hip replacement are poorer than those of primary replacement, with patients reporting worse pain and poorer function at 5 years after revision surgery.

| Study | Design | Duration | Patients | Comparisons/ evaluations | Outcomes |
|---------------------------------|--------------------------------|---|---|--|---|
| Fortin et al. [®] | Observational; longitudinal | Preoperative to 6 mo after hip replacement | n = 116, undergoing total hip replacement for osteoarthritis | Compared 6-mo outcomes between those with "high" (better) and "low" (worse) function preoperatively | SF-36 physical function score, * WOMAC pain score, † WOMAC functional limitation score, ‡ hip range of motion: mean difference (high-low) (95% Cl) 16.4 (7.3 to 25.5), -1.9 (-3.0 to -0.7), -5.9 (-9.7 to -2.1), 12.8°, respectively (i.e., worse 6-mo status among those with worse function preoperatively). |
| Garbuz et al.° | Observational; longitudinal | From placement on waiting list to 1 yr after hip replacement | n = 147, undergoing total hip replacement for osteoarthritis | Examined probability of achieving better than expected outcome at 1 yr based on "long" (> 6 mo) v. "short" (\leq 6 mo) wait and by count of months on waiting list from decision to proceed with surgery to operation | Better than expected v. not better than expected WOMAC outcomes. 43% of patients with short waits v. 31% of those with long waits achieved better than expected functional outcome. Those wit long waits had 50% decreased odds for achieving a better than expected outcome compared with those with shor waits. Each additional month spent waiting was associated with an 8% decreased odds (adjusted OR 0.92, p = 0.05) of better than expected functional outcome. No evidence of negative effect of wait time found for WOMAC pain and stiffness domains. |
| Vergara et al. ¹⁰ | Observational; longitudinal | Preoperative to 6 mo after hip replacement | n = 527, undergoing total hip replacement for osteoarthritis | Examined predictors of change in WOMAC domains over 6 mo, and predictors of achieving minimal clinically important difference on WOMAC domains, including presurgery status and wait time | WOMAC pain score, WOMAC functional limitation score, WOMAC stiffness score. Change in function was poorer ($p = 0.025$) among those who waited > 6 mo for surgery. Progressive reduction in % of patients surpassing minimal clinically important difference with increasing watting < 3 mo, 3-6 mo and > 6 mo, respectively ($p < 0.001$). Likelihood of perceiving a gain greater than minimal clinically important difference was lower (OR 0.47, $p = 0.006$) with > 6 mo wait compared with < 3 mo wait. No effects on other WOMAC domains observed. |
| Hajat et al. ¹¹ | | | | Examined predictors of 12-mo Oxford Hip Score status, including presurgery status and wait time | Oxford Hip Score¶ (measure of severity of hip problems: pain, disability, loss of physical function). Trend of worse 12-mo status with worse presurgery status ($p < 0.001$), longer wait to first outpatient appointment ($p < 0.001$) and longer time on wait list ($p < 0.001$). |
| Fortin et al. ¹² | Observational; longitudinal | Preoperative to 2 yr after hip replacement | n = 84, undergoing total hip replacement for osteoarthritis (subset of sample from Fortin et al. ⁸ who completed 2-yr survey) | Compared 6-mo outcomes between those with "high" (better) and "low" (worse) function preoperatively | SF-36 physical function score,* WOMAC pain score,† WOMAC functional limitation score‡: mean difference (high–low) (95% Cl) 14.7 (2.6 to 26.8), -1.5 (-3.1 to 0.1), -6.6 (-11.8 to -1.4), respectively (i.e., worse 2-yr status among those with worse function preoperatively). |

Note: CI = confidence interval, OR = odds ratio, SF-36 = 36-Item Short-Form Health Survey, WOMAC = Western Ontario and McMaster Universities Arthritis Index. *The SF-36 physical function score ranges from 0 to 100: higher scores indicate better physical health, and a minimal clinically important difference is 20 points. †The WOMAC pain score ranges from 0 to 20 and is standardized to a range of 0 to 100: 0 represents the best health status and 100 the worst health status, and the minimal clinically important difference is 29 points.

*The WOMAC functional limitation score ranges from 0 to 68 and is standardized to a range of 0 to 100: 0 represents the best health status and 100 the worst health status, and the minimal clinically important difference is 26 points.

\$The WOMAC stiffness score ranges from 0 to 8 and is standardized to a range of 0 to 100: 0 represents the best health status and 100 the worst health status, and the minimal clinically important difference is 25 points.

¶The Oxford Hip Score ranges from 0 to 48: 0 represents maximum disability and 48 no disability, and the minimal clinically important difference is between 4 and 6 points.

What surgical options are there?

In the early stages of osteoarthritis, joint-preserving procedures such as pelvic osteotomy (for insufficient acetabular coverage of the femur) or hip arthroscopy (for femoroacetabular impingement) may be considered depending on the patient's underlying diagnosis. These procedures are generally not recommended for patients with advanced degenerative changes. Patients with

| Study; variable | Outcomes (generally assessed between 3 and 24 mo after surgery) |
|--------------------------------|---|
| Ethgen et al. ¹⁶ | |
| Age | Age was not a factor in pain outcomes. Results were mixed for physical function: either no effect or older age associated with somewhat worse scores. Reported change in pain and function were similar across age groups, but status was generally worse with older age. |
| Sex | Results were mixed: either no difference in change in pain and function, or men trended toward greater improvement in function and/or pain. Status at follow-up trended toward better among men. |
| Ethnicity | Black patients had less change in pain and function than white patients. |
| Obesity | Higher BMI was generally associated with greater change in pain and function, but worse status. |
| Education | Range of health-related quality-of-life outcomes: higher educational attainment was generally associated with greater improvement. |
| Comorbidity | Greater levels of comorbidity were generally associated with less improvement in pain and function. The influence appeared to be greater among older age groups. |
| Jones et al. ¹⁷ | |
| Age | Older age at surgery was associated with greater satisfaction. |
| Obesity | Results were inconsistent for health-related quality-of-life outcomes. Generally, no influence was identified; some patients reported worse postoperative pain and functional status. |
| Mental well-being | Preoperative psychological status explained some variation in postoperative pain and function. High levels of anxiety or depression were associated with worse outcomes. |
| Comorbidity | A greater number of comorbid conditions were associated with worse short-term pain and functional outcomes. The overall impact appeared to be relatively small. The influence of older age was believed to be mediated through number of conditions. |
| Santaguida et al. ¹ | 8 |
| Age | Results were inconsistent for revision surgery; younger patients were at somewhat greater risk at 2–20 yr. Older age was associated with greatest risk of death at 30–90 d. Older age was associated with poorer function, though not when assessed using WOMAC. Age was not associated with postoperative satisfaction, but older age was |
| Sex | associated with less satisfaction after revision surgery. Results were inconsistent for revision surgery; men were at somewhat greater risk, particularly younger men. Men were generally at greater risk of death at 30–90 d. Women generally had poorer function, showed less functional improvement and showed less postoperative pain. Sex was not associated with satisfaction following the primary procedure, but women reported less satisfaction following revision surgery. |
| Obesity | Higher BMI was associated with poorer postoperative function. Obesity status was not associated with postoperative satisfaction. |

advanced joint damage are best referred to a surgeon to consider the options of either hip resurfacing or total hip replacement.

In a traditional hip replacement, the femoral head and damaged acetabulum are both removed and replaced with metal, plastic or ceramic components (Figure 1). Cement may be used for fixation, but most hip replacements performed in Canada are now uncemented23 owing to longer implant survival.24-26 Although ethical concerns, and costs, preclude clinical trials for establishing the merits of hip replacement,27 many observational studies have shown this procedure to be highly effective (and cost-effective) in minimizing pain and restoring function.16,28-34 Because surgical techniques and the design of prostheses and materials have improved over several decades, the risks of complications and early revisions following replacement have diminished. Rates of complications occurring within 90 days after surgery were found to be 1.0% for mortality, 0.9% for pulmonary embolus, 0.2% for wound infection, 4.6% for hospital readmission and 3.1% for hip dislocation among the US Medicare population.³⁵ Ten-year revision rates after surgery can range from 5% to 20%, depending on age and fixation technique.36

Despite the overall success of hip replacement, studies have documented that 5% to 25% of patients who undergo this procedure report minimal improvement or dissatisfaction with their outcomes.^{14,16-18,37-43}

In hip resurfacing, the femoral head is left in place but trimmed and capped with a metal covering. The damaged acetabulum cartilage is removed and replaced with a metal shell, similar to a traditional hip replacement. The proposed advantages of resurfacing include bone conservation among patients likely to outlive a traditional replacement (i.e., younger patients), improved hip range of motion and the potential for allowing



Figure 1: Components of traditional hip replacement.

younger patients an increased level of activity. Reported disadvantages include increased risk of femoral neck fracture, more bone loss on the acetabular side, a more difficult operation requiring larger incisions, and increased risk of systemic exposure to metal ions resulting from wear of the metal-on-metal bearing surface.⁴⁴⁻⁴⁹

Hip resurfacing is predominantly considered for the young active patient with end-stage osteoarthritis.⁴⁷ Early and mid-term follow-up generally has shown comparable results to those of standard replacement.⁵⁰⁻⁵⁹ However, complications particular to this procedure have been identified (as stated previously), and emphasis is placed on patient selection, component selection and surgical technique to avoid poor and adverse outcomes and short-term failures.^{44,47,60-70} Many surgeons avoid resurfacing in postmenopausal women because of an increased risk of femoral neck fracture, and in those with known renal insufficiency owing to the potential for metal ion accumulation.

Earlier generations of resurfacing devices frequently failed, often because of problems with excessive wear of the bearing surface materials. Since then, improvements in surgical technique and design have led to a renewed interest and use of hip resurfacing.

Is hip replacement or is hip resurfacing the best approach?

Matched comparative studies and clinical trials have compared short- and long-term outcomes of hip resurfacing versus hip replacement (Table 3).^{48,56,58,70-73} One of the larger matched studies with 5-year follow-up compared patients who had undergone hip resurfacing with those who had undergone traditional hip replacement.⁵⁶ The authors found that the hip resurfacing group were more active in running (58.5% v. 13.7%, p < 0.001), sports (73.6% v. 33.3%, p < 0.001) and manual labour (60.4% v. 39.2%, p = 0.049).⁵⁶

The consensus from 3 clinical trials^{70,71,74} is that hip resurfacing has no added advantage with respect to gait speed, stride length, stair climbing,⁷¹ range of motion⁷⁴ or health-related quality of life.^{70,71,74} One study found that a substantially higher number of patients who had undergone hip resurfacing returned to moderate or high activity levels 1 year after surgery (77%) compared with those who had undergone hip replacement (39%).⁷⁵ No published trial appears to be adequately powered to compare the complication rates of resurfacing versus replacement, however.

Limited to short-term follow-up, current trial evidence does not adequately address the differ-

ence in implant lifespan between resurfacing and replacement. Norwegian⁷³ and Australian⁷² registry data showed increased rates of revision at 2 and 5 years, respectively, for resurfacing versus replacement. Longer-term follow-up periods for present resurfacing designs are needed. A recent UK-based registry study reported high revision rates among women who had undergone hip resurfacing (8.5% at 5 yr), and the authors advocated against resurfacing in female patients.⁴⁸

Given the added expense of implants used in hip resurfacing, a cost–benefit analysis should be considered in future trials comparing this procedure with hip replacement.

Is mini-incision hip surgery preferable to conventional hip surgery?

The traditional approaches to hip replacement are the direct lateral and posterior approaches. The traditional incision is typically 15 cm or greater in length. Differences have not been shown for dislocation, limp or function between the 2 approaches.76 Mini-incision hip surgery was developed with the goal of decreasing tissue injury and blood loss, and improving patient outcomes. Although there is no universally accepted definition of mini-incision, many define it as an incision less than 12 cm in length. Patients considered appropriate candidates should have a body mass index of 30 or less, or a thigh circumference of less than 50 cm.77,78 Clinical trial evidence comparing mini-incision with conventional approaches is limited (Table 477,79,80). For smaller incisions, findings suggest no difference for in-hospital morphine use (mean 42.9 \pm 97.4 mg v. 45.0 \pm 96.8 mg, p =0.89), and reduced intraoperative blood loss (mean 314 ± 162 mL v. 366 ± 190 mL, p = 0.03) but no difference in transfusion rates (mean 0.42 ± 0.95 units v. 0.30 ± 0.66 units, p = 0.27).^{77,79,80} Differences in patient function 3 months and 1 year after surgery were not found.80,81

| Study | Design | Final follow-up, yr | Sample size | Outcome | Findings at follow-up (total hip replacement v. hip resurfacing) |
|---------------------------------------|-------------------------------------|---------------------------------|----------------|---|--|
| Garbuz et al. ⁷⁰ | RCT | Mean 1.1 (range 0.8– 2.2) | 107 | WOMAC score, mean | 90.1 v. 90.4, <i>p</i> = 0.950 |
| | | | | SF-36 physical function score, mean | 51.2 v. 51.2, <i>p</i> = 0.979 |
| Lavigne et al. ⁷¹ | RCT | 1 | 48 | Gait speed, m/s | 1.46 ± 0.18 v. 1.44 ± 0.19, <i>p</i> > 0.05 |
| | | | | Step length, m | 0.68 ± 0.07 v. 0.67 ± 0.07, p > 0.05 |
| Pollard et al. ⁵⁶ | Retrospective, matched cohort | , 5–7 | 108 | UCLA activity score | 6.8 v. 8.4, <i>p</i> < 0.001 |
| | | | | EQ-5D score | 0.78 v. 0.9, <i>p</i> = 0.003 |
| Smith et al.48 | UK registry | 5 | > 400 000 | Implant failure | Total hip replacement: 2.8% (95% Cl 2.7% to 2.9%) |
| | | | | | Hip resurfacing: Men: 3.6% (95% Cl 3.3% to 3.9%) Women: 8.5% (95% Cl 7.8% to 9.2%) |
| Corten and MacDonald ⁷² | Australian registry | 5 | > 135 000 | Implant failure | Total hip replacement: 2.7% |
| | | | | | Hip resurfacing: 3.7%, <i>p</i> < 0.001 |
| Johanson et al. ⁷³ | Norwegian registry | 2 | > 170 000 | Implant failure, cumulative revision rate | Total hip replacement: 1.2% (95% CI% 1.2 to 1.3%) |
| | | | | | Hip resurfacing: 3.3% (95% Cl 2.2% to 4.3%), p < 0.001 |

Note: CI = confidence interval, EQ-5D = Euro-Qol 5-dimension, RCT = randomized controlled trial, SF-36 = 36-Item Short-Form Health Survey, UCLA = University of California Los Angeles, WOMAC = Western Ontario and McMaster Universities Arthritis Index.

What are the comparative benefits and risks of metal-on-metal versus other bearing surfaces?

Metal-on-metal and other "hard-on-hard" bearing surfaces such as ceramic have gained popularity in recent decades because of their potential for less wear and improved implant longevity over traditional polyethylene bearing surfaces. To date, trial evidence has not indicated any benefit of hard-on-hard bearing surfaces over polyethylene for implant survivorship (Table 5).^{82–88} A recent UK-based registry report showed higher revision rates for metal-on-metal bearing surfaces (6.57%, 95% CI 5.10% to 8.43%) compared with ceramic (3.00%, 95% CI 2.45% to 3.68%) and polyethylene (2.03%, 95% CI 1.69% to 2.44%) at 7-year follow-up.⁸⁹ The concern with metal-on-metal implants is the accumulation of cobalt and chromium ions in the body, with a potential for cardiac toxicity, local soft-tissue erosions (pseudotumours, estimated incidence of 1% at 5 yr⁹⁰) or neurologic complications.^{45,91-93} A case series that reported on early failures (within 2–3 yr of surgery) cited implant loosening in 56% of these failures, and the remainder as soft-tissue reactions, pseudotumours and persistent pain.⁹⁴ These

Table 4: Selected studies comparing outcomes of minimally invasive and standard-incision total hip

 replacement

| Study | Design | Final follow-up | Sample size | Primary outcome | Findings at follow-up (mini-incision v. standard incision) |
|----------------------------------|----------------------------------|--------------------|----------------|---|--|
| Dorr et al. ⁸⁰ | RCT | 6 mo | 60 | Total blood loss, mL, mean | 352.3 ± 145.5 v. 408.3 ± 158.3, <i>p</i> = 0.12 |
| | | | | Length of stay, h, mean | 63.2 ± 13.3 v. 73.6 ± 23.5, p = 0.04 |
| Ogonda et al. ⁷⁷ | RCT | 6 wk | 219 | 10-m walk time, s, mean | 54.4 ± 29.8 v. 54.5 ± 32.7, p = 0.97 |
| | | | | Stair climbing, s, mean | 19.31 ± 8.78 v. 19.58 ± 9.38 p = 0.83 |
| | | | | Hematocrit level on discharge, mean | 0.275 ± 0.04 v. 0.276 ± 0.04 p = 0.75 |
| | | | | 36-h VAS pain score, mean | 16.8 ± 20.3 v. 19.8 ± 21.2, p = 0.29 |
| Chimento et al. ⁷⁹ | Retrospective, matched cohort | 1 | 60 | Total blood loss, mL, mean | 378 ± 151 v. 504 ± 205, p < 0.009 |
| | | | | 6-wk limp | 21.4% v. 46.8%, <i>p</i> = 0.04 |
| | | | | 1-yr limp | None |
| | | | | Hospital length of stay, d, mean (range) | 5.8 (4–13) v. 5.5 (3–15), p = 0.6 |

Table 5: Selected studies comparing outcomes of metal-on-metal and metal-on-polyethylene hip replacement

| Study | Design | Final follow-up, yr | Sample size | Outcome | Mean difference (95% CI) (metal-on-metal v. metal-on- polyethylene) | |
|--|--------|------------------------|----------------|------------------|---|--|
| Dahlstrand et al. ⁸⁴ | RCT | 2 | 54 | Harris Hip score | 3.1 (–2.0 to 8.2) | |
| Engh et al. ⁸⁵ | RCT | 2 | 59 | Harris Hip score | 4.0 (–0.4 to 8.4) | |
| Lombardi et al. ⁸⁶ | RCT | 5.7 | 99 | Harris Hip score | 1.2 (–1.9 to 4.3) | |
| MacDonald et al. ⁸⁷ | RCT | 3.2 | 41 | Harris Hip score | 0.4 (–7.0 to 7.8) | |
| Zijlstra et al. ⁸⁸ | RCT | 10 | 200 | Harris Hip score | 1.0 (–2.9 to 4.9) | |
| Note: CI = confidence interval, RCT = randomized controlled trial. | | | | | | |

implants have recently garnered substantial media attention owing to some companies removing their products from the market over concerns of early failures, and owing to safety advisories from the US Food and Drug Administration and Health Canada.^{95,96} Metal-on-metal bearings should be avoided in women of childbearing age because of the risk of metal ion accumulation. Patients with metal-on-metal hip replacements without symptoms should have regular follow-up by their surgeon.

Ceramic implants have not shown improved longevity over polyethylene, and they have the potential risk of fracture.⁹⁷ Further, up to 10% of ceramic hips may have an audible "squeak," which is troubling enough that patients have required revision to change the bearing surface.⁹⁸ Clinical trial data and cost–benefit analyses, with a minimum 15–20 years of follow-up, are still needed to definitively address the issue of "best" bearing surface.

What management options exist when patients require revision?

Based on expert consensus, patients with implants should be followed biennially by an orthopedic surgeon, regardless of symptoms, for clinical and radiologic investigation of implant wear. Symptoms of implant wear or loosening may include groin or thigh pain, or symptoms of hip instability (i.e., a feeling of giving way).

Revision surgery, including multiple revisions, may be required for a diagnosis of aseptic loosening of 1 or both components, recurrent dislocation or infection. For recurrent dislocation, assuming adequate component alignment, the surgical options are revision to a larger femoral head or a hip-stabilizing (constrained) acetabular component. For aseptic loosening, revision options exist; however, bone grafting, cement or metallic augments may be needed to compensate for bony defects. For infection, the common procedure is a 2-stage revision, whereby the components are removed and a temporary implant is used to deliver local antibiotic therapy in addition to parenteral antibiotic therapy. Following infection eradication, new hip implants are inserted. Length of stay in acute care after revision is generally longer than for the primary operation (6.2 v. 4.0 d), and patients report worse pain and poorer function at 5 years following revision compared with the primary surgery.99,100

Future directions

More people are living longer and indications for hip surgery are expanding to include younger patients, which suggest a potentially greater future demand for revision surgery. To minimize this likelihood, continued improvement of design, materials and surgical techniques are key.¹⁰¹

As indicated, there are still gaps in our understanding of the benefits and disadvantages of hip resurfacing versus hip replacement, mini-incision versus conventional approaches, and optimal bearing surfaces. These areas require further research.

Efforts in tissue engineering and biologic therapies for osteoarthritis have focused on repair of cartilage defects. Methods have included bone marrow stimulation techniques, osteochondral grafting and chondrocyte implantation.^{102,103} Although many are investigating stem cell use in cartilage regeneration, limited evidence is currently available to support routine use of this technique.

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Affiliations: Division of Orthopedic Surgery, University of Toronto and University Health Network (Gandhi, Mahomed); and the Arthritis Program, University Health Network and the Institute of Health Policy, Management and Evaluation, University of Toronto (Perruccio), Toronto, Ont.

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