

Correlation of Delayed Gadolinium-Enhanced MRI of Cartilage (dGEMRIC) Value With Hip Arthroscopy Intraoperative Findings and Midterm Periacetabular Osteotomy Outcomes

Jessica H. Lee,* MD, Darby A. Houck,* BA, Brandt A. Gruizinga,* BA, Tigran Garabekyan,† MD, Mary K. Jesse,‡ MD, Matthew J. Kraeutler,§ MD, and Omer Mei-Dan,*|| MD

Investigation performed at the Department of Orthopedics, School of Medicine, University of Colorado, Aurora, Colorado, USA

Background: Delayed gadolinium-enhanced magnetic resonance imaging of cartilage (dGEMRIC) is an advanced imaging technique that is purported to quantify cartilage damage in acute and chronic joint disease and predict periacetabular osteotomy (PAO) outcomes. There is a paucity of literature relating dGEMRIC values to arthroscopic findings before PAO and postoperative outcomes after PAO.

Purpose: To assess the utility and validity of dGEMRIC as a preoperative and prognostic assessment tool of cartilage status and integrity as it relates to intraoperative findings and midterm postoperative outcomes after PAO.

Study Design: Case series; Level of evidence, 4.

Methods: We analyzed a cohort of 58 patients (70 hips) with a median age of 30.1 years (range, 15-50) with hip dysplasia who underwent hip arthroscopy, followed by a PAO with preoperative dGEMRIC. The primary outcome measures were intraoperative assessment and correlation with cartilage damage (presence of cartilage flap, Outerbridge grade of the acetabulum and femoral head). Secondary outcome measures were postoperative patient-reported outcome (PRO) scores, including the International Hip Outcome Tool and Non-arthritic Hip Score. Correlation analyses were performed to determine the relationship between dGEMRIC values and (1) PROs and (2) intraoperative assessment of cartilage damage.

Results: There were significant negative linear relationships between dGEMRIC values and the primary outcome measures: presence of a cartilage flap (coronal, $P = .004$; sagittal, $P < .001$), Outerbridge grade of acetabular articular cartilage lesion (coronal, $P = .002$; sagittal, $P = .003$), and Outerbridge grade of femoral head articular cartilage lesion (coronal, $P = .001$; sagittal, $P < .001$). Despite significant overall improvement in all patients, there was no significant correlation between preoperative dGEMRIC values and improvement in PROs from presurgery to latest postoperative follow-up (median, 2.2 years; range, 1.0-5.0 years).

Conclusion: Although dGEMRIC values (sagittal and coronal) were significant predictors of the intraoperative presence of cartilage flaps and overall cartilage integrity, they were not associated with midterm outcomes after PAO.

Keywords: hip dysplasia; osteoarthritis; dGEMRIC; cartilage degeneration; CU PAO

Progressive cartilage damage of the hip joint commonly results from aberrant joint reactive forces associated with unfavorable anatomy, such as acetabular dysplasia and/or femoroacetabular impingement.^{15,19} Left unaddressed, chondral damage is thought to progress to end-stage arthritic change and is associated with poor clinical and

functional outcomes.¹⁵ Periacetabular osteotomy (PAO) is a surgical intervention that corrects acetabular orientation in dysplastic cases to improve hip joint stability and subsequently joint biomechanics, which delays the development of secondary osteoarthritis and enhances patient function and activity.^{25,26} Assessing articular cartilage damage before surgical intervention requires not only clinical information but also imaging for accurate identification, which is essential for selecting appropriate treatment strategies and surgical planning.²³ While preoperative

The Orthopaedic Journal of Sports Medicine, 10(9), 23259671221117606
DOI: 10.1177/23259671221117606
© The Author(s) 2022

This open-access article is published and distributed under the Creative Commons Attribution - NonCommercial - No Derivatives License (<https://creativecommons.org/licenses/by-nc-nd/4.0/>), which permits the noncommercial use, distribution, and reproduction of the article in any medium, provided the original author and source are credited. You may not alter, transform, or build upon this article without the permission of the Author(s). For article reuse guidelines, please visit SAGE's website at <http://www.sagepub.com/journals-permissions>.

plain radiographs are often used to analyze cartilage status via joint space measurements and osseous changes,¹⁹ magnetic resonance imaging (MRI) and arthrogram provide more direct cartilage assessment but lack information regarding the biological characteristics and health status of cartilage.^{25,26}

Gadolinium is an anionic contrast agent acting as a functional measure of cartilage status by binding avidly to damaged cartilage with less anionic glycosaminoglycans as compared with healthy cartilage.⁴³ It has been proposed that 3-dimensional delayed gadolinium-enhanced MRI of cartilage (dGEMRIC) improves specificity for identifying articular cartilage defects associated with hip pathologies, such as femoroacetabular impingement and dysplasia, via enhanced gadolinium binding and subsequent shortened T1 relaxation times.¹³

Although a few studies have associated low dGEMRIC scores (indicative of worse cartilage status) with poor PAO outcomes,^{13,18} there is a deficit of literature defining the relationship between dGEMRIC and both arthroscopic findings and clinical outcomes. This information would be useful to hip-preservation surgeons who may consider implementing the use of dGEMRIC into the standard preoperative imaging assessment, with the end goal of determining whether surgical success and/or failure can be predicted by dGEMRIC score.

The purpose of this study was to assess the utility and validity of dGEMRIC as a preoperative and prognostic assessment tool of cartilage status and integrity as it relates to intraoperative findings and postoperative outcomes after PAO.

METHODS

After institutional review board approval, a prospective longitudinal cohort study was performed involving 70 hips (58 patients) treated with the University of Colorado PAO²⁸ between November 2015 and June 2018. All patients were prospectively enrolled in an institutional review board-approved hip registry. All procedures were performed by the senior author (O.M.-D.). Inclusion criteria were (1) persistent hip pain refractory to nonoperative management lasting at least 6 months, (2) reproducible clinical examination findings suggestive of intra-articular pain and instability, (3) a joint-space width >3 mm on all radiographic views, and (4) radiographic findings consistent with frank or borderline hip dysplasia.²³

Patients with borderline dysplasia who exhibited substantial signs of instability²⁰ (Beighton Hypermobility Score⁴⁰ >6, excessive femoral and/or acetabular version) were typically advised that the risk of failure of hip arthroscopy alone was high, although this was presented as a valid first-line treatment option. When these patients elected to proceed with a PAO after a previously failed arthroscopic-only treatment, hip arthroscopy was again performed before the PAO to address any new chondrolabral pathology that may have developed since the previous arthroscopic surgery or residual cam pathology. Patients with concomitant excessive femoral ante/retrotorsion were offered a derotational femoral osteotomy in addition to hip arthroscopy and PAO.

Clinical diagnosis of acetabular frank or borderline dysplasia was determined according to accepted pathomorphologic signs and measurements. History of hip pain, positive findings on provocative hip tests indicating a labral tear, radiographic evidence of hip dysplasia (lateral center-edge angle <25°, sourcil angle >10°), excessive acetabular version and/or femoral antetorsion, and interruption of the Shenton line on the anteroposterior pelvic radiograph, as well as MRI findings of labral hypertrophy,^{16,21,32} articular cartilage thickening,¹ or a ligamentum teres tear,⁴¹ all aided in establishing a diagnosis of symptomatic hip instability. Measurements were performed by a hip-preservation fellow (J.H.L.) and verified by the senior author. Patients selected for surgery underwent preoperative computed tomography and MRI to assess acetabular version, femoral torsion, and femoral head sphericity as well as cartilage, labral, and subchondral bone integrity. Coronal and sagittal dGEMRIC indices of the hip joint were obtained during preoperative imaging.

Before the PAO operation, all patients underwent routine hip arthroscopy 3 to 10 days prior for direct visualization of cartilage status and to address intra-articular pathology. Hip arthroscopy was performed before PAO, rather than during the same anesthetic, for the following reasons: (1) to avoid prolonged anesthesia time, (2) to prevent fluid from hip arthroscopy in the surgical field during the PAO, (3) to reduce the rate of capsulolabral adhesions by enabling stationary bicycle use before PAO, and (4) to allow the surgeon and patient to discuss possibly opting out of the PAO if substantial articular cartilage damage is seen during arthroscopy.²⁸ Cartilage status during arthroscopy was assessed and subsequently graded using the Outerbridge grading scale.³⁰ Cartilage flaps were identified and confirmed with use of an arthroscopic probe. Acetabular

||Address correspondence to Omer Mei-Dan, MD, School of Medicine, University of Colorado, 12631 East 17th Ave, Mail Stop B202, Room L15-4505, Aurora, CO 80045, USA (email: omer.meidan@cuanschutz.edu).

*Department of Orthopedics, School of Medicine, University of Colorado, Aurora, Colorado, USA.

†Southern California Hip Institute, Los Angeles, California, USA.

‡Department of Radiology, School of Medicine, University of Colorado, Aurora, Colorado, USA.

§Department of Orthopedics & Sports Medicine, Houston Methodist Hospital, Houston, Texas, USA.

Final revision submitted March 21, 2022; accepted May 12, 2022.

One or more of the authors has declared the following potential conflict of interest or source of funding: T.G. has received consulting fees from Stryker. M.K.J. has received consulting fees and speaking fees from Medtronic USA and hospitality payments from Medicea USA. O.M.-D. has received consulting fees from Stryker. AOSSM checks author disclosures against the Open Payments Database (OPD). AOSSM has not conducted an independent investigation on the OPD and disclaims any liability or responsibility relating thereto.

Ethical approval for this study was obtained from the University of Colorado Denver (CRV006-1).

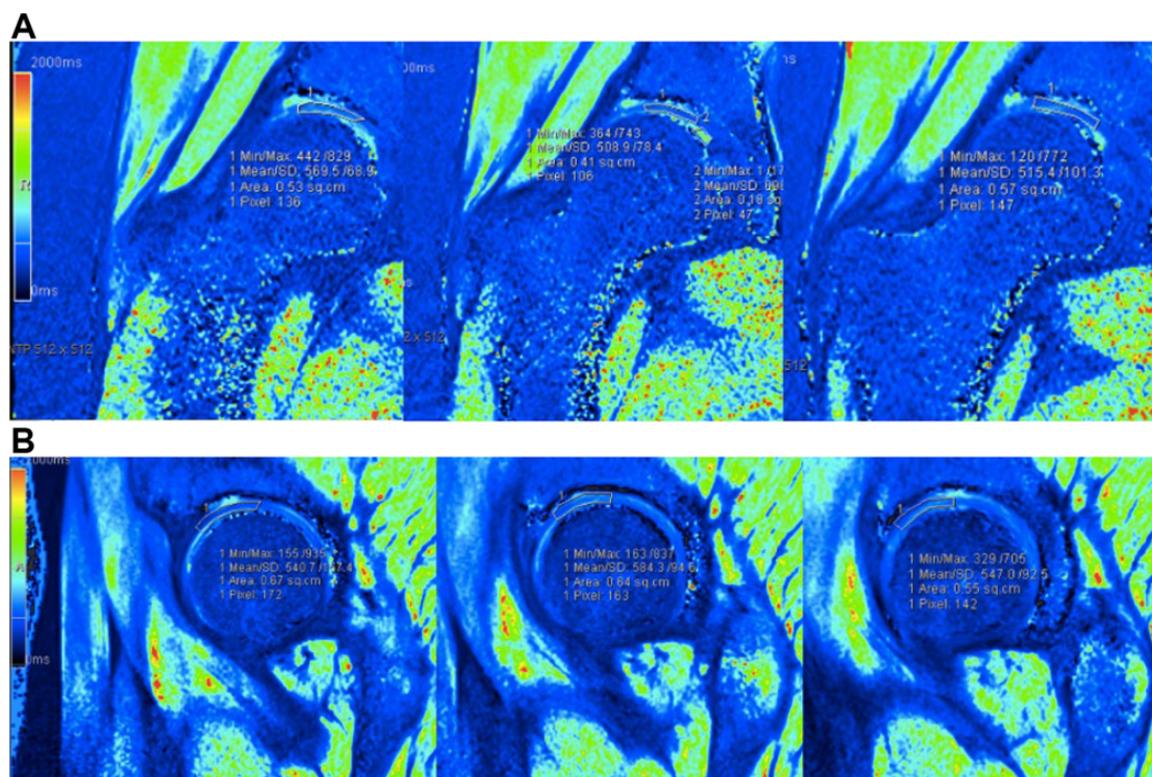


Figure 1. The bulk dGEMRIC values were the mean of 3 consecutive images obtained from the following regions of interest: (A) the weightbearing articular cartilage in the anterior, middle, and posterior coronal views (outlined in yellow) and (B) the anterolateral quadrant in the medial, central, and lateral sagittal views (outlined in yellow). All images were obtained from a patient who exhibited a cartilage flap intraoperatively. dGEMRIC, delayed gadolinium-enhanced magnetic resonance imaging of cartilage.

chondral flaps were often “inside-out” lesions with an intact chondrolabral junction.²² Any cartilage flaps identified at the time of arthroscopy were addressed with debridement, with or without microfracture, and subsequent limitations in weightbearing to allow viable fibrocartilage to develop into the former hyaline cartilage defect.

Imaging Protocol

The dGEMRIC protocol was based on established literature-supported parameters.^{2-4,8-10} Sequences of the hips were obtained on a single 1.5-T MRI scanner optimized for dGEMRIC. Coronal and sagittal dGEMRIC sequences of the hip were obtained 1 hour postinjection of a double weight-based dose (0.4 mL/kg) of gadobenate dimeglumine (Multihance). T1 map images were created by fitting saturation recovery curves to varying image intensity, and 3-dimensional variable flip angle images were then used to create the color dGEMRIC map on which regions of interest (ROIs) were drawn. Coronal ROI measurements were taken over the weightbearing femoroacetabular joint cartilage in 3 positions (anterior, middle, and posterior) (Figure 1A). Sagittal ROI measurements were taken across the anterior superior quadrant in 3 positions (medial, central, and lateral) (Figure 1B). Measurements were obtained by a single musculoskeletal radiologist (M.K.J.).

Outcome Measures

All patients enrolled in the study completed preoperative questionnaires and detailed demographic data sheets. Collected data included age, clinical diagnosis, sex, height, weight, body mass index, duration of pain at initial presentation, laxity using the Beighton Hypermobility Score, prior hip arthroscopy, radiographic parameters measured on anteroposterior pelvis radiographs,³⁹ and hip range of motion. The primary outcome measures were arthroscopic intraoperative assessment and correlation with cartilage damage. The secondary outcome measures were the Non-arthritic Hip Score (NAHS)¹² and the International Hip Outcome Tool (iHOT-12).¹⁷ Patient-reported outcome (PRO) scores were measured pre- and postoperatively at 6 and 12 weeks, 6 and 12 months, and 2 and 5 years.

Statistical Analysis

Patient demographics and outcome variables (baseline, intraoperative, PROs, and other outcomes) were summarized using descriptive summary measures expressed as means with standard deviations or medians with ranges for continuous variables, depending on the distribution, and as number (percentage) for categorical and ordinal variables. In patients with at least 12 months of follow-up, Wilcoxon tests were used to compare PROs (iHOT-12 and NAHS) between presurgery

TABLE 1
Baseline Patient Demographics (70 Hips in 58 Patients)^a

| Variable | Median [Range] or No. (%) |
|---|---------------------------|
| Age at periacetabular osteotomy, y | 30.1 [15-50] |
| Surgical hip laterality | |
| Left | 33 (47.1) |
| Right | 37 (52.9) |
| Sex: hips | |
| Female | 63 (90.0) |
| Male | 7 (10.0) |
| Sex: patients | |
| Female | 51 (87.9) |
| Male | 7 (12.1) |
| Body mass index | 22.9 [18.0-35.0] |
| Duration of pain at initial presentation, y | 3.00 [0.20-20.0] |
| Laxity: Beighton Hypermobility Score | 4.0 [0-9.0] |
| Prior hip arthroscopy | |
| No | 57 (81.4) |
| Yes | 13 (18.6) |

and latest postoperative follow-up. A Pearson correlation coefficient was computed to determine the relationship between the dGEMRIC values and the PAO outcome (defined as raw improvement in PROs from presurgery to latest postoperative follow-up). Additionally, biserial or polyserial correlations were computed for quantitative variables (dGEMRIC values) and a dichotomous variable (binary or ordinal; presence of cartilage flap, Outerbridge grade of the acetabulum, or Outerbridge grade of femoral head). A distribution-based method was used to calculate the minimal clinically important difference (MCID) of both PROs.²⁹ Linear regression analysis was performed using ordinary least squares regression to determine whether prior hip arthroscopy, lower preoperative scores, or duration of symptoms predicted latest postoperative PROs in patients with at least 12-month follow-up. All analyses were performed in R Statistical Software.³⁴ For all tests, statistical significance was set at $P < .05$.

RESULTS

Demographic Data

The demographic characteristics for the 70 hips (58 patients) are outlined in Table 1, and radiographic parameters are presented in Table 2. The mean dGEMRIC coronal value was 420 ± 101 , and the mean dGEMRIC sagittal value was 397 ± 108 .

Intraoperative Findings

Intraoperative findings are presented in Table 3. Direct visualization of cartilage status during hip arthroscopy before the PAO demonstrated an acetabular cartilage flap in 34 hips (Outerbridge grade 4; 48.6%), while 78.6% of hips had Outerbridge grade 0 of the femoral head. The majority of hips (62.9%) exhibited cartilage damage that extended

TABLE 2
Radiographic Parameters^a

| Variable | Value (milliseconds) |
|------------------------------------|----------------------|
| Tönnis grade | 0.214 \pm 0.45 |
| Mean \pm SD | |
| Median [range] | 0 [0 to 2.0] |
| Sharp angle, deg | |
| Mean \pm SD | 45.2 \pm 4.0 |
| Median [range] | 45.0 [36.0 to 54.0] |
| Lateral center-edge angle, deg | |
| Mean \pm SD | 18.4 \pm 7.4 |
| Median [range] | 18.0 [-4.0 to 37.0] |
| Tönnis angle, deg | |
| Mean \pm SD | 11.9 \pm 7.0 |
| Median [range] | 11.5 [-6.0 to 32.0] |
| Lateral joint space width, mm | |
| Mean \pm SD | 4.6 \pm 0.82 |
| Median [range] | 4.6 [3.1 to 6.6] |
| Medial joint space width, mm | |
| Mean \pm SD | 4.4 \pm 1.1 |
| Median [range] | 4.2 [2.7 to 8.0] |
| Acetabular equatorial version, deg | |
| Mean \pm SD | 22.5 \pm 5.3 |
| Median [range] | 23.0 [9.0 to 33.0] |
| Femoral torsion, deg | |
| Mean \pm SD | 16.9 \pm 9.4 |
| Median [range] | 16.0 [-3.0 to 37.0] |
| dGEMRIC: coronal, ms | |
| Mean \pm SD | 420 \pm 101 |
| Median [range] | 410 [231 to 643] |
| dGEMRIC: sagittal, ms | |
| Mean \pm SD | 397 \pm 108 |
| Median [range] | 404 [117 to 684] |
| COTAV | |
| Mean \pm SD | 39.4 \pm 11.1 |
| Median [range] | 38.5 [18.0 to 61.0] |
| Shenton line, ^b No. (%) | |
| Intact | 54 (77.1) |
| Interrupted | 13 (18.6) |
| Broken | 3 (4.3) |

^aCOTAV, combined index of femoral torsion and acetabular version; dGEMRIC, delayed gadolinium-enhanced magnetic resonance imaging of cartilage.

^bThe Shenton line was divided into interrupted and broken based on the degree of stepoff.

from 10% to 30% of the rim to the acetabular fossa distance (width of damage was measured according to hours of the clock).

In the 34 hips (48.6%) with a cartilage flap present, the mean dGEMRIC coronal value was 391.0 ± 99.1 , and the mean dGEMRIC sagittal value was 360.0 ± 110.0 (Table 4). In the 36 hips (51.4%) without a cartilage flap present, the mean dGEMRIC coronal value was 447.0 ± 96.8 , and the mean dGEMRIC sagittal value was 431.0 ± 96.2 .

Primary Outcome Measure

In the overall cohort of patients, there was a weak negative linear significant relationship between the dGEMRIC

TABLE 3
Intraoperative Findings (70 Hips)

| Variable | No. (%) |
|-----------------------------------|-----------|
| Presence of cartilage flap | |
| Yes | 34 (48.6) |
| No | 36 (51.4) |
| Outerbridge grade: acetabulum | |
| 0 | 11 (15.7) |
| 1 | 24 (34.3) |
| 2 | 0 (0) |
| 3 | 0 (0) |
| 4 | 35 (50.0) |
| Outerbridge grade: femoral head | |
| 0 | 55 (78.6) |
| 1 | 5 (7.1) |
| 2 | 4 (5.7) |
| 3 | 2 (2.9) |
| 4 | 4 (5.7) |
| Cartilage damage rim to fossa, % | |
| <10 | 16 (22.9) |
| 10-30 | 44 (62.9) |
| >30 | 10 (14.3) |
| Microfracture of acetabulum | |
| Yes | 35 (50.0) |
| No | 35 (50.0) |
| Bone grafting of acetabular cysts | |
| Yes | 7 (10.0) |
| No | 63 (90.0) |
| Labral reconstruction | |
| Yes | 4 (5.7) |
| No | 66 (94.3) |

TABLE 4
Intraoperative Cartilage Status: dGEMRIC Values^a

| | dGEMRIC Value, ms | |
|---------------------------------|-------------------|---------------|
| | Coronal | Sagittal |
| Presence of cartilage flap | | |
| Yes (n = 34) | 391.0 ± 99.1 | 360.0 ± 110.0 |
| No (n = 36) | 447.0 ± 96.8 | 431.0 ± 96.2 |
| Outerbridge grade: acetabulum | | |
| 0 (n = 11) | 476.0 ± 120.0 | 433.0 ± 103.0 |
| 1 (n = 24) | 436.0 ± 85.6 | 433.0 ± 95.4 |
| 2 (n = 0) | NA | NA |
| 3 (n = 0) | NA | NA |
| 4 (n = 35) | 391.0 ± 97.6 | 359.0 ± 109.0 |
| Outerbridge grade: femoral head | | |
| 0 (n = 55) | 431.0 ± 100.0 | 410.0 ± 103.0 |
| 1 (n = 5) | 410.0 ± 112.0 | 422.0 ± 87.0 |
| 2 (n = 4) | 391.0 ± 129.0 | 299.0 ± 146.0 |
| 3 (n = 2) | 392.0 ± 17.1 | 402.0 ± 86.8 |
| 4 (n = 4) | 315.0 ± 47.1 | 264.0 ± 111.0 |

^aData are reported as mean ± SD. dGEMRIC, delayed gadolinium-enhanced magnetic resonance imaging of cartilage; NA, not applicable.

coronal value and the presence of a cartilage flap during hip arthroscopy ($r = -0.34$; $P = .004$) (Table 5). A moderate negative linear significant relationship occurred between

TABLE 5
Correlation Between dGEMRIC Values and Parameters Related to the Arthroscopic Intraoperative Assessment of Cartilage Damage^a

| Parameter 1 | Parameter 2 | r | P Value |
|-------------------|---------------------------------|-------|---------|
| dGEMRIC: coronal | Presence of cartilage flap | -0.34 | .004 |
| dGEMRIC: sagittal | Presence of cartilage flap | -0.41 | <.001 |
| dGEMRIC: coronal | Outerbridge grade: acetabulum | -0.36 | .002 |
| dGEMRIC: sagittal | Outerbridge grade: acetabulum | -0.35 | .003 |
| dGEMRIC: coronal | Outerbridge grade: femoral head | -0.38 | .001 |
| dGEMRIC: sagittal | Outerbridge grade: femoral head | -0.45 | <.001 |

^aAll correlations, $P < .05$. dGEMRIC, delayed gadolinium-enhanced magnetic resonance imaging of cartilage.

the dGEMRIC sagittal value and the presence of a cartilage flap during hip arthroscopy ($r = -0.41$; $P < .001$).

In the overall cohort of patients, the dGEMRIC coronal value and the Outerbridge grade of acetabulum during hip arthroscopy demonstrated a weak negative linear significant relationship ($r = -0.36$; $P = .002$). There was a weak negative linear significant relationship between the dGEMRIC sagittal value and the Outerbridge grade of acetabulum during hip arthroscopy ($r = -0.35$; $P = .003$) and between the dGEMRIC coronal value and the Outerbridge grade of femoral head during hip arthroscopy ($r = -0.38$; $P = .001$). The dGEMRIC sagittal value and the Outerbridge grade of femoral head during hip arthroscopy had a moderate negative linear significant relationship ($r = -0.45$; $P < .001$).

PRO Scores

Fourteen hips were lost to follow-up. In the remaining 56 hips with a minimum follow-up of 1 year (median, 2.2; range, 1.0-5.0), significant improvements in the iHOT-12 ($P < .001$) and NAHS ($P < .001$) were achieved from pre-surgery to latest postoperative follow-up (Table 6). Using the distribution-based method, the MCID was 9.7 for the iHOT-12 and 9.0 for the NAHS. At the latest postoperative follow-up, 85.7% (n = 48) of patients reached the MCID for the iHOT-12, and 67.9% (n = 38) of patients reached the MCID for the NAHS. There was no relationship between dGEMRIC values and improvement in PROs from pre-surgery to the latest postoperative follow-up ($P > .05$ for all). Additionally, linear regression models demonstrated that prior hip arthroscopy, lower preoperative scores, and duration of symptoms did not predict latest postoperative PROs.

DISCUSSION

Our results were demonstrative of dGEMRIC value being a significant predictor of cartilage damage in a population of

TABLE 6
Patient-Reported Outcome Scores (n = 56)^a

| Outcome Measure | Preoperative | Postoperative | P Value |
|-----------------|------------------|------------------|---------|
| iHOT-12 | 37.8 [4.17-91.1] | 87.8 [28.3-99.3] | <.001 |
| NAHS | 58.8 [2.50-100] | 91.3 [41.3-100] | <.001 |

^aData are reported as median [range]. Each pre- and postoperative comparison, $P < .05$. iHOT-12, International Hip Outcome Tool; NAHS, Non-arthritis Hip Score.

patients that required PAO surgery for bony instability. Decreasing dGEMRIC scores on both views (coronal and sagittal) were related to the presence of a cartilage flap. Additionally, dGEMRIC values negatively correlated with Outerbridge score of the femoral head and the acetabulum graded at the time of hip arthroscopy. Although surgical intervention alone resulted in significant improvements in PROs (iHOT-12 and NAHS) from presurgery to latest postoperative follow-up, there was no relationship between preoperative dGEMRIC values and improvement in PROs.

Because there was no significant relationship between the dGEMRIC value and either the iHOT-12 or the NAHS at follow-up, this calls into question the utility of the test as a predictor of failure after PAO. Therefore, dGEMRIC may not be a useful diagnostic adjuvant in determining whether a patient with dysplasia is a candidate for hip preservation surgery vs adult reconstruction. Although a lower dGEMRIC score was correlated with the presence of cartilage flaps, it was not correlated with clinical outcomes in our cohort. The presence of chondral injury can be assessed on preoperative MRI without the use of dGEMRIC. Thus, our study does not support the routine use of dGEMRIC in the preoperative assessment of patients undergoing PAO.

Prior studies utilizing dGEMRIC value thresholds have likewise demonstrated mixed results in diagnostic and predictive utility. The standard dGEMRIC value of healthy cartilage in patients (mean age, 37 years) has been reported to be 570 ± 90 ms, while values <390 ms are associated with osteoarthritis as well as increased risk of failure after pelvic osteotomy.¹⁸ Similarly, Cunningham et al¹³ found a mean dGEMRIC index of 370 ± 88 ms to be the greatest predictor of PAO failure at a follow-up, ranging from 2 years to 3 years 10 months. By contrast, Chandrasekaran et al¹¹ reported significantly greater PROs at 2 years in nondysplastic cases of hip arthroscopy with a dGEMRIC >323 ms, but they were unable to establish an actual correlation between dGEMRIC and PROs. The inconsistencies in dGEMRIC thresholds and subsequent variability in postoperative outcomes in these studies reinforce the concept that value thresholds are population-based and therefore limited in their predictive utility outside the population included in each study. The mean coronal and sagittal dGEMRIC values for our cohort were greater for patients without a cartilage flap than with a cartilage flap (447.0 and 431.0 ms vs 391.0 and 360.0 ms, respectively). Although our mean sagittal dGEMRIC value of 360 ms in patients with a cartilage flap falls below the 390 ms,¹⁸ suggestive of increased risk of PAO failure, this was not borne

out clinically in our cohort. This further suggests that employing a dGEMRIC threshold may have a more limited role than previously demonstrated.

It is possible that the use of dGEMRIC bulk values as detailed in the protocol may be slightly confounded in a dysplastic population, which may have clinical implications. A recent study evaluated the use of dGEMRIC in assessing human cartilage adaptation to exercise. It called into question the validity of dGEMRIC bulk values, which may be misleadingly elevated in individuals with inherently thicker cartilage (ie, elite athletes with cartilage hypertrophy) and subsequently higher precontrast T1 values from which the bulk values are primarily derived.³⁶ In a dysplastic population in which cartilage thickness is significantly greater¹ than a nondysplastic cohort as measured by lateral center-edge angle, there exists potential for similar confounding in the dGEMRIC values. The implications of a falsely elevated dGEMRIC secondary to increased cartilage thickness as it relates to postoperative outcomes suggest that bulk values may be misleadingly high in a dysplastic population regardless of cartilage viability.

While there may be a small role for dGEMRIC in localizing isolated acetabular chondral lesions in preoperative patients,^{24,31} arthroscopic surgical intervention in and of itself for appropriate pathologies necessitating hip-preservation procedures seems to be a better predictor, albeit an uncontrolled one, of prognosis than dGEMRIC score, as demonstrated by our study and others.^{31,35} Schmaranzer et al³⁵ found significant decreases in preoperative vs minimum 1-year postoperative dGEMRIC scores in operative and nonoperative groups. Despite a more pronounced negative difference (decrease) in operative group dGEMRIC score after surgery, all PROs (Western Ontario and McMaster Universities Osteoarthritis Index, Hip disability and Osteoarthritis Outcome Score, modified Harris Hip Score) improved significantly in the operative group, and only 1 (modified Harris Hip Score) improved in the nonoperative group. This begets further skepticism of the predictive capabilities of dGEMRIC with respect to clinical outcome.

In our dysplastic cohort, the relationship between dGEMRIC score and cartilage flap was marginally stronger for sagittal ($r = -0.41$) over coronal ($r = -0.34$) sequences. This is consistent with our initial expectations, as cartilage flaps are better identified and characterized on sagittal sequences because of the primarily anterolateral orientation of cartilage flaps in a dysplastic cohort. Conversely, the finding that the relationship between dGEMRIC and acetabular Outerbridge grade was marginally stronger for coronal ($r = -0.36$) over sagittal ($r = -0.35$) views was less consistent with expectations, especially given that all patients with acetabular grade 4 cartilage damage had a cartilage flap identified at the time of arthroscopy. It is possible that hyaline cartilage flaps, which represent the thickest area of cartilage and are best visualized on coronal sequence, may occasionally present with higher dGEMRIC scores by virtue of thickness alone, as discussed earlier,³⁶ thus confounding the relative relationships observed. Even so, this study differentiates itself by utilizing coronal and sagittal views. By contrast, Palmer et al³¹ utilized only

sagittal imaging, while Lerch et al²⁴ utilized radial imaging volumetrically reconstituted for their studies.

Studies have touted dGEMRIC as a quantitative molecular assessment of cartilage integrity via preferential binding of glycosaminoglycans,^{3,37} but it adds significant burden and risk to patients, including time, pain, and unstandardized billing ranging from US\$2300 to US\$24,000.²⁷ Patients must adhere to specific timed protocols to obtain requisite imaging for optimal dGEMRIC values.³³ Additionally, the use of gadolinium contrast agents, specifically those belonging to the group I subtype, is reported to have significant risk of systemic fibrosis in patients with renal disease^{38,42} and may cause nonrenal acute reactions, such as anaphylaxis and local necrosis at the site of injection.⁶ Therefore, dGEMRIC must demonstrate clinical benefits that outweigh these potential harms before this advanced imaging modality can be adopted by hip preservation surgeons as a routine imaging supplement.

There exist multiple noncontrasted alternate advanced imaging modalities that demonstrate noninferior and superior diagnostic and/or predictive capabilities as compared with dGEMRIC that do not expose patients to the aforementioned factors.⁵ Beaulé et al⁵ identified strongly significant correlations between T1 ρ and T2 ρ MRI mapping (noncontrasted studies) and dGEMRIC values in a cohort of patients with developmental dysplasia of the hip. Furthermore, Bittersohl et al⁷ found dGEMRIC to have weak correlation with intraoperative findings of cartilage damage among those undergoing surgical intervention for femoroacetabular impingement, while standard MRI evaluation alone had a moderate correlation. Additional quantitative magnetic resonance techniques, such as T1 rho imaging, quantitative T2 mapping, and sodium MRI, may provide similar diagnostic capabilities without the use of contrast.³³

Limitations

The limitations of this study should be noted. First, the sample size was low and limited our ability to quantify dGEMRIC values based on each Outerbridge grade of the acetabulum and femoral head. In addition, there is a lack of a clinically significant threshold dGEMRIC value, which could provide a guideline. We did not include this grouping because a significant relationship was demonstrated but still not predictive of outcome, thus lending little clinical utility to such a value. All cartilage flaps were included in the same category, though the size and location of these defects may affect outcomes. In addition, microfracture and debridement of chondral flaps at the time of hip arthroscopy may have affected follow-up PROs differently from prior research¹³ in which intra-articular procedures were not performed before PAO. All procedures and intraoperative diagnoses were performed by a single surgeon, which may introduce some observer bias. The median follow-up time for PROs was 2.2 years with a range of 1 to 5 years in our study cohort. Finally, because distribution-based MCIDs are typically sample dependent,¹⁴ application of this MCID may be limited to populations with similar characteristics.

We did not stratify the data based on specific patient demographics or characteristics to determine MCIDs for these subpopulations, as this was not the purpose of the present study.

Future Directions

Further study is warranted to determine the utility of dGEMRIC in predicting long-term outcomes after PAO. Additional investigation is necessary to determine if various noncontrasted methods of imaging are noninferior to dGEMRIC with regard to diagnostic and prognostic value. Finally, future studies may seek to determine if cartilage thickness based on MRI or plain radiographs (in terms of joint space width) correlates with dGEMRIC values.

CONCLUSION

Although dGEMRIC values (sagittal and coronal) were significant predictors of the intraoperative presence of cartilage flaps and overall cartilage integrity, these findings were not associated with midterm outcomes after PAO.

REFERENCES

1. Ashwell ZR, Flug J, Chadayammuri V, Pascual-Garrido C, Garabekyan T, Mei-Dan O. Lateral acetabular coverage as a predictor of femoroacetabular cartilage thickness. *J Hip Preserv Surg*. 2016;3(4):262-269.
2. Bashir A, Gray ML, Boutin RD, Burstein D. Glycosaminoglycan in articular cartilage: in vivo assessment with delayed Gd(DTPA)(2-)-enhanced MR imaging. *Radiology*. 1997;205(2):551-558.
3. Bashir A, Gray ML, Burstein D. Gd-DTPA2- as a measure of cartilage degradation. *Magn Reson Med*. 1996;36(5):665-673.
4. Bashir A, Gray ML, Hartke J, Burstein D. Nondestructive imaging of human cartilage glycosaminoglycan concentration by MRI. *Magn Reson Med*. 1999;41(5):857-865.
5. Beaulé P, Melkus G, Rakhra K, Wilkin G. Quantitative hip cartilage MRI of patients with hip dysplasia: evaluation of dGEMRIC, T1 ρ and T2 mapping. *Orthop Proc*. 2020;102B:25.
6. Bellin MF, Van Der Molen AJ. Extracellular gadolinium-based contrast media: an overview. *Eur J Radiol*. 2008;66(2):160-167.
7. Bittersohl B, Hosalkar HS, Apprich S, Werlen SA, Siebenrock KA, Mamisch TC. Comparison of pre-operative dGEMRIC imaging with intra-operative findings in femoroacetabular impingement: preliminary findings. *Skeletal Radiol*. 2011;40(5):553-561.
8. Burstein D, Bashir A, Gray ML. MRI techniques in early stages of cartilage disease. *Invest Radiol*. 2000;35(10):622-638.
9. Burstein D, Gray M. New MRI techniques for imaging cartilage. *J Bone Joint Surg Am*. 2003;85(suppl 2):70-77.
10. Burstein D, Velyvis J, Scott KT, et al. Protocol issues for delayed Gd(DTPA)(2-)-enhanced MRI (dGEMRIC) for clinical evaluation of articular cartilage. *Magn Reson Med*. 2001;45(1):36-41.
11. Chandrasekaran S, Vemula SP, Lindner D, Lodhia P, Suarez-Ahedo C, Domb BG. Preoperative delayed gadolinium-enhanced magnetic resonance imaging of cartilage (dGEMRIC) for patients undergoing hip arthroscopy: indices are predictive of magnitude of improvement in two-year patient-reported outcomes. *J Bone Joint Surg Am*. 2015;97(12):1305-1315.
12. Christensen CP, Althausen PL, Mittleman MA, Lee JA, McCarthy JC. The nonarthritic hip score: reliable and validated. *Clin Orthop Relat Res*. 2003;406:75-83.
13. Cunningham T, Jessel R, Zurakowski D, Millis MB, Kim YJ. Delayed gadolinium-enhanced magnetic resonance imaging of cartilage to

- predict early failure of Bernese periacetabular osteotomy for hip dysplasia. *J Bone Joint Surg Am.* 2006;88(7):1540-1548.
14. de Vet HC, Terwee CB, Ostelo RW, Beckerman H, Knol DL, Bouter LM. Minimal changes in health status questionnaires: distinction between minimally detectable change and minimally important change. *Health Qual Life Outcomes.* 2006;4:54.
 15. Ganz R, Leunig M, Leunig-Ganz K, Harris WH. The etiology of osteoarthritis of the hip: an integrated mechanical concept. *Clin Orthop Relat Res.* 2008;466(2):264-272.
 16. Garabekyan T, Ashwell Z, Chadayammuri V, et al. Lateral acetabular coverage predicts the size of the hip labrum. *Am J Sports Med.* 2016;44(6):1582-1589.
 17. Griffin DR, Parsons N, Mohtadi NG, Safran MR, Multicenter Arthroscopy of the Hip Outcomes Research Network. A short version of the International Hip Outcome Tool (iHOT-12) for use in routine clinical practice. *Arthroscopy.* 2012;28(5):611-616.
 18. Kim YJ, Jaramillo D, Millis MB, Gray ML, Burstein D. Assessment of early osteoarthritis in hip dysplasia with delayed gadolinium-enhanced magnetic resonance imaging of cartilage. *J Bone Joint Surg Am.* 2003;85(10):1987-1992.
 19. Kraeutler MJ, Garabekyan T, Goodrich JA, Fioravanti MJ, Chadayammuri V, Mei-Dan O. Standardizing the prearthritic hip joint space width: an analysis of 994 hips. *Arthroscopy.* 2018;34(7):2114-2120.
 20. Kraeutler MJ, Garabekyan T, Pascual-Garrido C, Mei-Dan O. Hip instability: a review of hip dysplasia and other contributing factors. *Muscles Ligaments Tendons J.* 2016;6(3):343-353.
 21. Kraeutler MJ, Goodrich JA, Ashwell ZR, Garabekyan T, Jesse MK, Mei-Dan O. Combined lateral osseolabral coverage is normal in hips with acetabular dysplasia. *Arthroscopy.* 2019;35(3):800-806.
 22. Kraeutler MJ, Goodrich JA, Fioravanti MJ, Garabekyan T, Mei-Dan O. The "outside-in" lesion of hip impingement and the "inside-out" lesion of hip dysplasia: two distinct patterns of acetabular chondral injury. *Am J Sports Med.* 2019;47(12):2978-2984.
 23. Kraeutler MJ, Safran MR, Scillia AJ, Ayeni OR, Garabekyan T, Mei-Dan O. A contemporary look at the evaluation and treatment of adult borderline and frank hip dysplasia. *Am J Sports Med.* 2020;48(9):2314-2323.
 24. Lerch TD, Ambuhl D, Schmaranzer F, et al. Biochemical MRI with dGEMRIC corresponds to 3D-CT based impingement location for detection of acetabular cartilage damage in FAI patients. *Orthop J Sports Med.* 2021;9(3):2325967120988175.
 25. Lerch TD, Steppacher SD, Liechti EF, Tannast M, Siebenrock KA. One-third of hips after periacetabular osteotomy survive 30 years with good clinical results, no progression of arthritis, or conversion to THA. *Clin Orthop Relat Res.* 2017;475(4):1154-1168.
 26. Leunig M, Siebenrock KA, Ganz R. Rationale of periacetabular osteotomy and background work. *Instr Course Lect.* 2001;50:229-238.
 27. Marshall KW, Braithwaite K, Safdar N, et al. The rise and fall of dGEMRIC evaluation in the adolescent hip. *Orthop J Sports Med.* 2019;7(3)(suppl):2325967119S00016.
 28. Mei-Dan O, Welton KL, Kraeutler MJ, Young DA, Raju S, Garabekyan T. The CU PAO: a minimally invasive, 2-incision, interlocking periacetabular osteotomy: technique and early results. *J Bone Joint Surg Am.* 2019;101(16):1495-1504.
 29. Norman GR, Sloan JA, Wywich KW. Interpretation of changes in health-related quality of life: the remarkable universality of half a standard deviation. *Med Care.* 2003;41(5):582-592.
 30. Outerbridge RE. The etiology of chondromalacia patellae. *J Bone Joint Surg Br.* 1961;43:752-757.
 31. Palmer A, Fernquest S, Rombach I, et al. Diagnostic and prognostic value of delayed gadolinium enhanced magnetic resonance imaging of cartilage (dGEMRIC) in early osteoarthritis of the hip. *Osteoarthritis Cartilage.* 2017;25(9):1468-1477.
 32. Pareek A, Carey JL, Reardon PJ, Peterson L, Stuart MJ, Krych AJ. Long-term outcomes after autologous chondrocyte implantation: a systematic review at mean follow-up of 11.4 years. *Cartilage.* 2016;7(4):298-308.
 33. Potter HG, Koff MF. MR imaging tools to assess cartilage and joint structures. *HSS J.* 2012;8(1):29-32.
 34. *R Core Team. R: a Language and Environment for Statistical Computing.* Version 1.1.456. R Foundation for Statistical Computing; 2016.
 35. Schmaranzer F, Haefeli PC, Hanke MS, et al. How does the dGEMRIC index change after surgical treatment for FAI? A prospective controlled study: preliminary results. *Clin Orthop Relat Res.* 2017;475(4):1080-1099.
 36. Tiderius CJ, Hawezi ZK, Olsson LE, Dahlberg LE. Pre-contrast T1 and cartilage thickness as confounding factors in dGEMRIC when evaluating human cartilage adaptation to physical activity. *BMC Med Imaging.* 2019;20(1):1.
 37. Trattng S, Huber M, Breitenheiser MJ, et al. Imaging articular cartilage defects with 3D fat-suppressed echo planar imaging: comparison with conventional 3D fat-suppressed gradient echo sequence and correlation with histology. *J Comput Assist Tomogr.* 1998;22(1):8-14.
 38. Weinreb JC, Rodby RA, Yee J, et al. Use of intravenous gadolinium-based contrast media in patients with kidney disease: consensus statements from the American College of Radiology and the National Kidney Foundation. *Radiology.* 2021;298(1):28-35.
 39. Welton KL, Jesse MK, Kraeutler MJ, Garabekyan T, Mei-Dan O. The anteroposterior pelvic radiograph: acetabular and femoral measurements and relation to hip pathologies. *J Bone Joint Surg Am.* 2018;100(1):76-85.
 40. Wolf JM, Cameron KL, Owens BD. Impact of joint laxity and hypermobility on the musculoskeletal system. *J Am Acad Orthop Surg.* 2011;19(8):463-471.
 41. Woodward RM, Vesey RM, Bacon CJ, White SG, Brick MJ, Blankenbaker DG. Microinstability of the hip: a systematic review of the imaging findings. *Skeletal Radiol.* 2020;49(12):1903-1919.
 42. Woolen SA, Shankar PR, Gagnier JJ, MacEachern MP, Singer L, Davenport MS. Risk of nephrogenic systemic fibrosis in patients with stage 4 or 5 chronic kidney disease receiving a group II gadolinium-based contrast agent: a systematic review and meta-analysis. *JAMA Intern Med.* 2020;180(2):223-230.
 43. Zilkens C, Jager M, Bittersohl B, et al. Delayed gadolinium enhanced MRI of cartilage (dGEMRIC): molecular MRI of hip joint cartilage. Article in German. *Orthopade.* 2009;38(7):591-599.