Reproducibility and Diagnostic Accuracy of Kellgren-Lawrence Grading for Osteoarthritis using Radiographs and Dual Energy X-ray Absorptiometry (DXA) images

Running Title: Osteoarthritis grading using DXA
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Abstract

Introduction: Advances in image quality from modern Dual Energy X-ray absorptiometry (DXA) scanners now allow near radiograph-like quality images at a low radiation dose. This opens potential new applications for the use of DXA scanners to study other musculoskeletal conditions, such as osteoarthritis which is often investigated by visual assessment of radiographs. Together, osteoporosis and osteoarthritis are the two most common musculoskeletal conditions, both of which primarily affect older people. The aim of this study was to determine whether Kellgren-Lawrence grading of DXA images can be used to grade hip osteoarthritis as effectively as radiographs.

Methodology: People who had attended for recent pelvic radiographs underwent DXA images of hips (50 hips from 25 people) using a GE Healthcare iDXA scanner. Three observers assigned Kellgren-Lawrence grades to each image and grading was repeated at least one week apart. Intra-observer and inter-observer reliability for radiograph and DXA were calculated using quadratic weighted kappa (QWK). People were recalled 12 months later and the tests were repeated with both the radiograph and DXA scans taken within 2 weeks of each other.

Results: Hip DXA intra-observer reproducibility achieved a QWK range of 0.88-0.95 and inter-observer reproducibility of 0.85-0.88, similar to QWK from hip radiographs. Intra-observer reliability between subject-matched radiograph and iDXA images revealed QWK ranging between 0.80-0.88.

Conclusions: Reproducibility of hip osteoarthritis grading using DXA was comparable with that of radiographs in this study and similar to repeatability scores previously published in literature. Given the lower radiation dose and the opportunity to simultaneously investigate osteoporosis, DXA presents an attractive imaging option for osteoarthritis.

Keywords: DXA; Osteoarthritis; Osteoporosis; Bone Mineral Density; Arthritis; Kellgren-Lawrence
INTRODUCTION

Osteoporosis (OP) and osteoarthritis (OA) are the two most common musculoskeletal disorders in the developed world. Although an inverse relationship between OP and OA has been suggested (1), the diseases can coexist (2, 3) and it would be attractive to be able to use a single imaging modality to assess both in the same site (4). Modern DXA scanners have a lower radiation dose than radiographs (5-56 μSv for hip DXA (5, 6), 700 μSv for pelvic or hip (7, 8) radiographs) yet have decent image resolution, allowing assessment of vertebral fractures and aortic calcification (9).

Following the authors’ observation that typical features of hip OA, osteophytes, sclerosis and joint space narrowing were clearly visible on Dual Energy Absorptiometry (DXA) images acquired for diagnosing OP by measuring Bone Mineral Density (BMD), this study investigated whether Kellgren-Lawrence grading (KLG), a standard radiographic technique for assessing osteoarthritis severity using plain films (10), can be applied reliably to DXA images.

Figure 1: Comparison between radiograph and DXA images. (A) plain radiograph of the hip (B) iDXA image of the same hip. The scale and contrast of images have been adjusted for viewing purposes.
MATERIALS AND METHODS

Recruitment
Subjects for this study were identified from a larger, longitudinal study investigating osteoarthritis. Subjects for the parent study were recruited with differing degrees of hip OA identified from the local National Health Service (NHS) Radiology Information System (RIS). All patients over 30 with bilateral hip/pelvis radiographs taken within the previous year were identified via five computerised searches (April-October 2007). Based on the radiology reports (aged over 30 years, with a plain pelvic or antero-posterior radiographs of hips or knees taken on or after 1st February 2006 in any speciality except Accident and Emergency (A&E), invitation letters were sent to potential participants via their referring clinician. Radiographs were then examined for suitability. The following exclusion criteria were applied: prior surgical interventions such as total hip replacements (THR), known skeletal metastases, infective or inflammatory arthropathies, congenital/developmental dysplasia, avascular necrosis, fractures/dislocations, other bone disease (e.g. Paget’s disease), or absence of a formal radiology report.

For eligible subjects who gave informed consent, DXA scans of both hips were obtained posteroanteriorly, using an iDXA scanner (GE Healthcare, Madison, WI, USA), using standard DXA positioning protocols. As part of the longitudinal study, twelve months later, they were invited for a repeat DXA scan and non-weight-bearing antero-posterior radiographs of the pelvis were also obtained. Baseline images were used for initial comparison of KLG. Results were later confirmed using the 12-month images where DXA and radiographic images were taken within 1 week.

A subset of baseline radiographs (50 hips, 25 subjects) with subject-matched DXA scans, encompassing the full KLG range (0-4) was selected for this reproducibility study by JSG who was not involved in grading. Radiographs and DXA images were graded independently and in random order by 3 observers (KY, SG-S, DMR) from rheumatology or radiology backgrounds at consultant and trainee level. The images were graded again, randomly and independently, at least one week later without knowledge of the previous grades by the same observers. To enable off-site scoring radiographic images from both visits were digitized for DMR, whereas only the second set was digitised for the other observers. Radiographs were digitized using a Howtek MultiRAD 850 (Howtek, Hudson, New Hampshire) at 146 dpi and 8-bit depth. Observers could identify left and right hip images from the same patients; no other subject identifiable information was available.

DXA and radiographic images from the second visit were graded by KY (twice) and SG-S (once). Six subjects from the original 50 either had a THR or withdrew before this visit. These were replaced with subjects of similar age and baseline KLG to ensure QWK statistics were directly comparable.
Table 1: Distribution of baseline radiographic KLG in the study.

<table>
<thead>
<tr>
<th>Grade recorded</th>
<th>Grade 0</th>
<th>Grade 1</th>
<th>Grade 2</th>
<th>Grade 3</th>
<th>Grade 4</th>
<th>TOTAL</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pelvic X-ray</td>
<td>5</td>
<td>16</td>
<td>13</td>
<td>7</td>
<td>9</td>
<td>50</td>
</tr>
</tbody>
</table>

Note: Modes of all grades for each image by the three observers were used, unless more than one mode was possible, where the median was rounded to the nearest integer.

Figure 2: Intra-observer reproducibility for iDXA and radiograph of the Hip. Error bar represents 95% confidence interval (based on standard error, H_0≠0)

Osteoarthritis grading
Images were graded according to the Kellgren and Lawrence system using a reprint of the Atlas of standard radiographs of arthritis (11) and the Atlas of individual radiographic features, revised (12) (for KLG 0, since a “normal” image was not included in the original atlas). Observers were permitted to alter contrast and magnification of digital images using ImageJ.

Statistics
Reliability was calculated using Quadratic Weighted-Kappa (QWK) using MedCalc (v9.4.1, MedCalc, Mariakerke, Belgium) and WINPEPI (v9.3, PAIRSetc) and Intraclass Correlation Coefficients (ICC) using SPSS (v17, SPSS Inc, Chicago, USA) with two way random and absolute agreement.

A Kappa score of “1” indicates perfect agreement, “0” chance and “-1” perfect disagreement. Kappa is suitable for dichotomous or unordered categorical variables. When categories have a ranking or order, such as KLG, weighted Kappa is more appropriate (13). For comparison with previous studies, we also calculated the ICC which, while most appropriately used for continuous variables, is equivalent to QWK (14).
RESULTS
The study comprised 12 men and 13 women, average age 65.9 (±9.3) years. The average interval between the recruitment radiograph and baseline DXA was 225 (±104) days. All 12-month DXAs and radiographs were taken within 1 week. Figure 1 shows a typical osteoarthritic DXA and radiograph. Table 1 shows the baseline radiographic KLG distribution. There were no adverse events from performing the radiograph or DXA scan.

Intra-observer and inter-observer reproducibility
Good levels of intra-observer (Figure 2) and inter-observer (Table 2) reproducibility were achieved. All observers had similar intra-observer QWK values of 0.88-0.95 and 0.85-0.88 for DXA and radiographs respectively, and corresponding absolute agreements of 68-88% and 62-66%. Inter-observer agreement for DXA images was also similar (QWK 0.85-0.88, Table 2).

Calculation of ICC for all values confirmed QWK approximated to ICC (14) with a difference of no more than 0.01 for both two-way random and two-way mixed effect models.

Table 2: Inter-observer reproducibility scores for each pair of observers

<table>
<thead>
<tr>
<th></th>
<th>Absolute agreement %</th>
<th>QWK (SE)</th>
<th>ICC (CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Obs 2</td>
<td>Obs 3</td>
<td>Obs 2</td>
</tr>
<tr>
<td>Hip iDXA</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Obs 1</td>
<td>60%</td>
<td>58%</td>
<td>0.86 (0.037)</td>
</tr>
<tr>
<td>Obs 2</td>
<td>56%</td>
<td>56%</td>
<td>0.85 (0.035)</td>
</tr>
<tr>
<td>Hip radiograph</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Obs 1</td>
<td>54%</td>
<td>34%</td>
<td>0.81 (0.048)</td>
</tr>
<tr>
<td>Obs 2</td>
<td>52%</td>
<td>52%</td>
<td>0.73 (0.063)</td>
</tr>
</tbody>
</table>

*Abbr: CI: 95% confidence interval; ICC: Intraclass correlation coefficient; Obs, observer, QWK: quadratic weighted kappa. SE: Standard error (H₀≠0).*
Figure 3: Comparison of study results to literature. Comparison of results from the current study measuring intra-observer (3a) and inter-observer reliability (3b) from KLG of baseline DXA images (striped black) and radiographs (dotted grey) with published data from other studies (plain grey) that measured the reliability of OA grading of the hip using ICC or QWK (18, 19, 27). For Gossec et al. (18) Ingvarsson et al. (28), as only one ICC value was obtained, the width of the bar was widened to 0.01 for visual ease. For joint space narrowing (JSN) in Günther and Sun (27), the ICC range includes both superior and medial JSN. Joint space width (JSW) was categorised to measure JSW (JSWcat) (18).

DXA vs. radiograph
Intra-observer agreement between radiographs and DXAs achieved QWK values of 0.80-0.88 (Figure 2) with no mode or median grades differing by more than 1 KLG.

12 month time-point
Intra-observer reliability for KY was 0.99 (SE 0.005; 95% CI 0.99-1) for DXA and 0.88 (SE: 0.03; 95% CI 0.82-0.94) for radiographs; inter-observer reliability between KY and SGS was 0.87 (SE 0.024; 95% CI 0.83-0.92) for DXA and 0.85 (SE 0.037; 95% CI 0.78-0.93) for radiographs. The intra-
observer QWK values of 0.89 and 0.95 between DXA and radiograph images were similar to, but slightly higher than baseline grades for these observers.

To put our results in context we searched the literature for studies reporting radiographic OA reliability. Figure 3 shows ICC and QWK values from published studies looking at KLG, Croft, Osteoarthritis Research Society international (OARSI) grading and joint space width compared to the current study.

**DISCUSSION**

These results demonstrate that KLG can be applied to DXA images of the hip from iDXA scanners as reproducibly as standard radiographs and the same grade was assigned to the majority of subjects, regardless of the image source. All QWK scores lay in, or above the 0.61–0.80 range, referred to as ‘good’ (15) or ‘substantial’ (16) agreement.

A simple classification of ‘good’ agreement using a cut-off value can be considered to be somewhat arbitrary (16, 17) and does not fully evaluate the strength of DXA imaging for OA. However as Figure 3 demonstrates, DXA scoring was at least as good as radiographs and figures in the literature. Unfortunately, it is difficult to compare with published results in more depth as there is often a lack of detail of OA severity or prevalence, and of the statistical tests used, for example specifying the version of kappa (weighted/unweighted, Fleiss-Cohen/Cichetti-Allison weights), or ICC (McGraw, Shrout-Fleiss’s 6 types), or the statistical package.

This study suffers from some limitations. Although comparable to many (18, 19), the number of subjects is still relatively small, and only basic randomisation (the order of images as presented to graders) was achieved. In addition, the study did not include a consensus session to discuss images where there was disagreement. This should affect DXA and radiographs equally and not add bias but may have reduced inter-observer repeatability.

Intra-observer scores comparing radiographs and DXA were “good”, though unsurprisingly slightly lower than for each modality alone, probably because of differing contrast and resolution. Although slightly higher, 12-month repeatability was similar to baseline, indicating that the gap between radiograph and DXA acquisition at baseline (225 days vs. < 7 days), caused by using historical radiographs taken as part of the subject’s normal healthcare for recruitment, had little impact on KL repeatability.

DXA has some advantages compared with radiographs, including the low radiation dose, measurement of BMD and the use of positioning devices and strict protocols as standard practice (recent testing of our radiographers using 60 volunteers gave a precision error of 0.72% and least significant change 2.0%, less than half those recommended by the International Society for Clinical Densitometry (ISCD) (20)). Disadvantages of DXA include lower resolution and the inability to take weight-bearing images, so joint space measurements cannot currently be as precise as on radiographs.

Osteoarthritis is a complex disease where there is limited concordance between symptoms and radiographic features and links between them are not fully understood (21). Whilst individuals may often be diagnosed on symptoms alone (22), structural changes fundamentally underpin disease progression and are also critical for complete understanding of osteoarthritis, particularly for evaluation of therapeutic agents.
CONCLUSION
This study has shown hip KLG on iDXA images is at least as reproducible as on radiographs. Our results were comparable with published literature and it may be that the use of positioning devices in DXA minimises variability. The ability to use DXA images to assess radiographical OA will create further clinical and research opportunities, although testing would be recommended for each manufacturer and scanner model. The relative accessibility and lower radiation dose of DXA makes the technology an appealing modality. Furthermore, as the elderly population at risk of both osteoporosis and osteoarthritis expands and therapeutic agents effective in osteoporosis show promise for use in osteoarthritis (23-26), the potential for a one-stop scan to assess both diseases makes DXA an attractive modality to consider for use in standard clinical practice.

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