

Research Article

Subchondral Bone Changes in Early Versus Late Osteoarthritis

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Abstract

Background: Osteoarthritis (OA) is the most common joint disorder, characterized by focal loss of cartilage and increased subchondral bone remodeling at early OA stages of the disease.

Aim of the work: To detect subchondral bone changes by MRI and level of in early versus late osteoarthritis. **Methods:** 30 patients diagnosed as OA with age ranged between 27-70 years, the patients further subdivided into early and late OA according to Kellgren and Laurence grading, all patients subjected to through history taking and full examination, plain x ray knees PA view, MRI on the most severely affected knee joint. **Results:** there was significant difference between subgroups regarding BME, BME score, attrition and osteophytes ($P = 0.000$, $p < 0.0001$, $p = 0.000$ and $p = 0.01$ respectively). **Conclusion:** BME presence and BME score were more common in early than late OA but present in both while attrition, subchondral cyst and osteophytes were more common in late OA.

Keywords: Osteoarthritis – BME- MRI.

Introduction

Osteoarthritis (OA) is the most common joint disorder, characterized by focal loss of cartilage and increased subchondral bone remodeling at early OA stages of the disease⁽¹⁾. There are many trials demonstrated that the presence of bone marrow lesions (BMLs) are closely related for structural deterioration in knee OA. Roemer et al., attributed subchondral bone attrition to BMLs⁽²⁾. BMLs adjacent to the subchondral plate have been shown to have increased bone volume fraction and increased trabecular thickness, but reduced tissue mineral density, consistent with OA being associated with increased bone turnover. Both subchondral bone abnormalities are associated with cartilage loss as well⁽³⁾. Crema et al., also indicated that subchondral cysts arise at the same site as BMLs⁽⁴⁾. Clinical studies have reported BMLs in both patients with early asymptomatic OA and in those with severe late-stage OA^(5, 6).

Aim of the work

To detect subchondral bone changes by MRI and level of in early versus late osteoarthritis.

Patients and methods

30 patients (23 female, 7 males) who fulfilling Arthritis Rheum 1987 OA classification criteria, with permission from ACR, from Rheumatology outpatient clinic, Minia University Hospital in the period from April 2014 to February 2016 were included in the study. The mean of age was 50.00 ± 12.94 (range 27-70 years) and the mean of disease duration was 3.74 ± 3.32 years (range 0.20-12 years). OA patients were classified into 2 groups: I: early OA, II: late RA according to Kellgren and laurence scale. Excluded from the study other established diagnostic entities other than OA, osteoarthritis with co-morbid endocrine, metabolic or major organ impairment. The nature of the study was explained to all patients. Radiological procedures represent standard care and pose no ethical conflicts. A verbal consent was obtained from all patients.

All patients were subjected to detailed medical history and complete physical examination. X ray knees PA view were ordered in all the patients, MRI of knees was performed for patients using SiGNA

profiler 0.2 tesla GE medical system machine.

Statistical analysis

Analysis of data was done by personal computer using SPSS (Statistical program for social science) version 16. The data of all software patients and controls were fed into an IBM personal computer. Data were expressed as mean±SD for parametric variables and as number and percent for non-parametric variable. Comparison between groups for parametric data was done by independent samples t-test (unpaired t-test). Chi – square (X²) test was used to compare qualitative variables. The difference was expressed as probability of value (P value). The difference was considered significant if P < 0.05.

Results

Demographic data of the studied population:

Osteoarthritis patients' age ranged between 27 and 70 years, the mean of age was 00.00±12.94 years and disease duration ranged between 0.2-12 years, the mean of was 3.74±3.32 years.

Subchondral bone changes by MRI in OA patients:

Table 1 demonstrates MRI subchondral bone changes in knee joint in OA group where BME present in 53.3% of cases and BME score was 2.87±3.49 while attrition was present in 70%, osteophytes in 73.3% and subchondral cyst in 40% of cases.

Subchondral bone changes by MRI in both groups:

Table 2 demonstrates MRI findings in both groups where there was statistically significant difference regarding bone marrow edema, BME score, bone attrition, osteophytes (P = 0.040, p<0.0001, p=0.000 and p=0.01 respectively).

Table 1: Frequency of subchondral bone changes by MRI in OA group:

MRI findings	OA patients n=30
BM edema	16 (53.3%)
BME score	2.87±3.49
Bone attrition	21 (70%)
Osteophytes	22 (73.3%)
Subchondral cyst	12 (40%)

Table 2: Comparison of subchondral bone changes by MRI in both groups:

MRI findings	Group I (early OA)	Group II (late OA)	χ ²	P value
BM edema	11 (73.3%)	0 (33.3%)	3.394	0.040
BME score	0.13 7.40±4.42	0.4 1.13±1.09	3.096	<0.0001
Subchondral cyst	4 (26.7%)	8 (53.3%)	2.222	0.136
Bone attrition	7 (46.7%)	14 (93.3%)	7.778	0.000
Osteophytes	8 (53.3%)	14 (93.3%)	6.136	0.01

Discussion

Osteoarthritis (OA) is a low-grade inflammatory disease of synovial joints and the most common form of arthritis⁽¹⁾. There are well described changes that are observed in both articular cartilage and subchondral bone in OA⁽²⁾. Changes in the bone include sclerotic changes, thinning of articular cartilage, thickening of the subchondral plate and subchondral cortical thickness, osteophyte formation, advancement of tidemark associated with vascular invasion of the calcified cartilage and the development of bone marrow lesions and bone cysts in the subchondral compartment⁽³⁾.

Focal changes in the subchondral bone, termed bone marrow lesions (BMLs), are features detected by magnetic resonance imaging (MRI) that have been reported to be closely associated with the severity of symptoms of OA such as pain and osteochondral unit degeneration (e.g., loss of the overlying cartilage)^(4,5).

Study population was divided according to Kellgren and Laurence grading scale into 2 groups: group I: early OA; grade 1 and 2 K&L, group II: late RA; grade 3 and 4 K&L

Our study revealed that there was a statistically significant difference between both groups regarding MRI findings where there was statistically significant difference regarding bone marrow edema, BME score, bone attrition, osteophytes ($P=0.040$, $p<0.0001$, $p=0.000$ and $p=0.01$ respectively).

In a study done by Bassiouni et al., 2011, BME was present in 14 patients (47%) with OA, and this percentage is close to our percentage which is 53.3% of OA patients⁽⁶⁾.

There is no previous studies addressed the comparison between early and late OA regarding MRI subchondral bone changes.

Conclusion

This study suggests that BME and BME were higher in early than late OA while subchondral cyst, bone attrition and

osteophytes were common in late than early OA.

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References

1. Zamli, Z., Robson Brown, K., & Sharif, M. (2016). Subchondral bone plate changes more rapidly than trabecular bone in osteoarthritis. *International Journal of Molecular Sciences*, 17(9), 1496.
2. Roemer, F. W., Neogi, T., Nevitt, M. C., Felson, D. T., Zhu, Y., Zhang, Y., & Lewis, C. E. (2010). Subchondral bone marrow lesions are highly associated with, and predict subchondral bone attrition longitudinally: the MOST study. *Osteoarthritis and cartilage*, 18(1), 47-53.
3. Baranyay, F. J., Wang, Y., Wluka, A. E., English, D. R., Giles, G. G., Sullivan, R. O., & Cicuttini, F. M. (2007, October). Association of bone marrow lesions with knee structures and risk factors for bone marrow lesions in the knees of clinically healthy, community-based adults. In *Seminars in arthritis and rheumatism* (Vol. 37, No. 2, pp. 112-118). WB Saunders.
4. Crema, M.D., Roemer, F.W., Zhu, Y., Marra, M.D., Niu, J., Zhang, Y., et al. (2010). Subchondral Cyst Like Lesions Develop Longitudinally in Areas of Bone Marrow Edema-Like Lesions in Patients with or at Risk for Knee Osteoarthritis: Detection with MR Imaging—The MOST Study. *Radiology*, 256, 800-812.
5. Zubler, V., Mengiardi, B., Pfirrmann, C. W., Duc, S. R., Schmid, M. R., Hodler, J., & Zanetti, M. (2007). Bone marrow changes on STIR MR images of asymptomatic feet and ankles. *European radiology*, 17(12), 3077-3082.
6. Berenbaum, F. (2013). Osteoarthritis as an inflammatory disease (osteoarthritis is not osteoarthrosis!). *Osteoarthritis and Cartilage*, 21(1), 16-21.

5. Tanamas, S. K., Wluka, A. E., Pelletier, J. P., Martel-Pelletier, J., Abram, F., Wang, Y., & Cicuttini, F. M. (2010). The association between subchondral bone cysts and tibial cartilage volume and risk of joint replacement in people with knee osteoarthritis: a longitudinal study. *Arthritis research & therapy*, 12(2), R08.
6. Roman-Blas, J. A., & Herrero-Beaumont, G. (2014). Targeting subchondral bone in osteoporotic osteoarthritis. *Arthritis research & therapy*, 16(6), 494.
7. Davies-Tuck, M. L., Wluka, A. E., Wang, Y., English, D. R., Giles, G. G., & Cicuttini, F. (2009). The natural history of bone marrow lesions in community-based adults with no clinical knee osteoarthritis. *Annals of the rheumatic diseases*, 18(6), 904-908.
8. Wluka, A. E., Hanna, F., Davies-Tuck, M., Wang, Y., Bell, R. J., Davis, S. R., ... & Cicuttini, F. M. (2009). Bone marrow lesions predict increase in knee cartilage defects and loss of cartilage volume in middle-aged women without knee pain over 5 years. *Annals of the rheumatic diseases*, 18(6), 800-800.
9. Bassiouni, H., Zaki, K., Elshorbagi, M., Mustapha, A., Tantawi, R., Ali, H., Metyas, S., & Arkfeld, D. G. (2010). Relating bone marrow oedema to hs-CRP in knee osteoarthritis. *Indian Journal of Rheumatology*, 0(1), 11-10.