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The Relationship Between Bone Mineral Density and Lumbar Disc Herniation in Postmenopausal Women

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Abstract

Introduction: In previous studies, the relationship between BMD (bone mineral density) and LDH (lumbar disc herniation) has been investigated in young people, except for postmenopausal women. The aim of this study was to evaluate this association in postmenopausal women.

Methods: A cross-sectional analysis of 545 consecutive postmenopausal women was performed at a single center. The study included patients aged 45 to 85 with low back pain. Age, weight, height, L1-L4 BMD, L1-L4 T-score, L1-L4 Z-score, femoral neck BMD, femoral neck T-score, and femoral neck Z-score of patients were collected. MRI scans were assessed for the diagnosis of LDH. To explore the impact of the number of herniated segments, patients with LDH were further divided into single-level and multi-level LDH groups.

Results: Five hundred and thirteen postmenopausal women were included in the final analysis. The mean age of the patients was 61.3 ± 8.6 years in the LDH group and 58.4 ± 7.8 years in the non-LDH group (p=0.001). The LDH group had higher lumbar (p<0.001) and femoral neck (p<0.05) BMD, T, and Z-scores than the non-LDH group. In binary logistic regression analysis, age, lumbar, and femoral neck BMD were significantly associated with LDH (p<0.001, p=0.03, and p=0.040, respectively). Patients with multi-level herniation had significantly higher rates of obesity (BMI \geq 30) compared to patients with single-level herniation (58.0% vs. 47.0%; p=0.031). However, in terms of obesity rates, the LDH group and the non-LDH group were statistically similar (53.9% vs. 54.2%; p=0.961). There was no association between the single and multi-level herniation groups in terms of L1-4 and femoral neck BMD (p=0.760 and 0.435, respectively).

Conclusion: Higher lumbar bone mineral density and higher femoral neck bone mineral densities were found to be associated with lumber disc herniation in postmenopausal women with low back pain. These results suggest that bone mineral density assessment may be useful in clinical practice to determine which patients are at higher risk of lumbar disc herniation.

Categories: Orthopedics

Keywords: cross sectional study, postmenopausal women, low back pain, disc herniation, bone mineral density

Introduction

Lumbar disc herniation (LDH) is one of the most frequent diseases that has a serious impact on people's quality of life and puts a heavy burden on families and society [1]. It is also the most frequent cause of sciatica and affects 1% to 5% of the population every year [2,3]. Therefore, it is important to elucidate the risk factors and the pathophysiological mechanisms involved in the development of LDH. Clarifying these mechanisms may enable prevention or early diagnosis of the disease. Previously reported associated factors include smoking, genetic factors, occupational factors, and obesity [4-6]. In addition, a recent study reported that symptomatic LDH increases with age, and the incidence is higher in women than in men [7].

Since bone mineral density (BMD) levels affect not only the structure of bones but also nonosseous tissues, osteopenia and osteoporosis disrupt the morphology of lumbar vertebral bones and intervertebral discs [8,9]. On the other hand, it has also been shown that stronger bones may be associated with more disc degeneration [9]. Nevertheless, although lumbar vertebral bones and lumbar intervertebral discs are close to each other both biomechanically and anatomically, to our knowledge, the association between BMD and LDH has not been studied in postmenopausal women before. Two studies were detected evaluating this relationship in young healthy people living in an urban area and young patients, with the exclusion of postmenopausal women, and the studies found no significant relationship [10,11].

Close contact between the vertebral bone marrow and the vertebral endplate is important for adequate nutrition of the nucleus pulposus of the disc [12]. High BMD causes increased end-plate resistance and affects the vascularization of the disc by disrupting diffusion [13]. Furthermore, high BMD may cause higher vertebral stiffness, and by this mechanism, mechanical stress on the disc may cause repetitive tears in the

annulus fibrosus and result in compression and herniation of the disc [14]. As the accumulation of small tears in the disc through these mechanical and vascular pathophysiological pathways occurs over many years, and the structure of the disc is stronger in young adults, high BMD may not result in LDH in young people through these mechanisms. Thus, we conducted this study to evaluate the relationship between BMD and LDH in postmenopausal women.

Materials And Methods

Study design

This study was a retrospective cross-sectional analysis based on an electronic database from a single center.

Patient selection

An analysis of 545 consecutive patients at a state hospital between January 2021 and February 2022 was performed. Inclusion criteria were a diagnosis of low back pain with or without radiculopathy that persisted for at least one-month, postmenopausal women, patients aged 45 to 85 years, and patients who had undergone both a BMD and a lumbar MRI within one month. Patients with malignancy, inflammatory joint disease, vertebral fractures, a history of spinal surgery for any reason, and those using medications that could affect bone metabolism were excluded from the study.

Outcome measures and confounders

Age, weight, height, L1-L4 BMD, L1-L4 T-score, L1-L4 Z-score, femoral neck BMD, femoral neck T-score, and femoral neck Z-score of the patients were collected from the electronic database of the hospital. Patients' weights and heights were measured before BMD measurements using a Soehnle Professional 7830 electronic scale (100g precision) and a standard stadiometer (1mm precision), respectively. Body mass index (BMI) was calculated by dividing the patient's body weight by the square of their height and was expressed as kg/m². BMI was stratified as: 18.5 to 24.9 kg/m², normal; 25.0 to 29.9 kg/m², overweight; 30.0 to 39.9 kg/m², obese; and >40 kg/m², morbidly obese, according to the World Health Organization (WHO) definitions.

According to WHO guidelines, a T-score between -1 and -2.5 was diagnosed as osteopenia, and a T-score of -2.5 or less as osteoperosis. Patients with a T-score of -1 or above were included in the normal BMD group. BMD measurements were performed using the dual energy X-ray absorptiometry (DXA) method using the LUNAR DPX densitometer (Madison Corporation, USA). MRI scans were obtained using a 1.5 Tesla machine (Philips, Suzhou, China). LDH was defined as "localized or focal displacement of disc material beyond the limits of the intervertebral disc space," according to the latest recommendations of the North American Spine Society [15]. The term "localized" or "focal" was defined as the extension of the disc material to less than 25% of the circumference of the disc as viewed in the axial plane [11]. The MRI scans (L1 to S1) were assessed by an independent reader who was blinded to the patients' details. This study was approved by the Institutional Review Board (Approval Number: 2022-1/23). A written informed consent was obtained from all individual participants included in the study.

Statistical analysis

Data were included in a database created by the Excel 2007 program by Microsoft (Microsoft Corporation, Redmond, Washington, USA). Statistical analysis was performed using PASW statistics for Windows (version 18, USA). The Shapiro-Wilk test was used to determine whether the continuous variables were normally distributed. Continuous variables that provided the parametric test assumption in comparison between groups were tested with Student's t-test. Categorical variables were compared using the Chi-Square test. A binary logistic regression analysis was performed to adjust for the effects of covariates, including age and BMI. To explore the impact of the number of herniated segments, patients with LDH were further divided into single-level and multi-level LDH groups. An ANOVA test was performed to analyze the relationship between herniated segment numbers and BMD. A p-value of <0.05 was considered significant.

Results

Characteristics of patients

Five hundred and forty-five postmenopausal women were initially enrolled in the study. After the exclusion of six patients with malignancy, 12 patients with vertebral fractures, two patients with rheumatoid arthritis, nine patients with a history of spinal surgery, one patient using glucocorticoids, and two patients using antiosteoporotic drugs, 513 patients were included in the final analysis. Four hundred and six patients had at least one level of LDH, and 107 patients had normal MRI findings. The mean age of the patients was 61.3 ± 8.6 years in the LDH group and 58.4 ± 7.8 years in the non-LDH group (p=0.001). The two groups were comparable in terms of height, body weight, and BMI (p=0.392, 0.324, and 0.154, respectively) (Table 1).

Characteristic	LDH	Non-LDH	P-value
Sample size	406 (79.1%)	107 (20.9%)	-
Age, years	61.3±8.6	58.4±7.8	0.001
Height, cm	157.7±6.5	158.2±5.2	0.392
Weight, kg	75.2±12.3	73.8±13.2	0.324
BMI, kg/m²	30.3±5.0	29.5±5.1	0.154
L1-4 BMD	1.1±0.2	1.0±0.1	<0.001
L1-4 T score	-0.6±1.3	-1.2±1.2	<0.001
L1-4 Z score	0.1±1.2	-0.6±1.0	<0.001
Femur neck BMD	0.9±0.1	0.8±0.1	0.004
Femur neck T score	-0.8±1.1	-1.0±0.9	0.023
Femur neck Z score	0.1±0.9	-0.3±0.7	<0.001

TABLE 1: Characteristics of the patients.

Notes: Variables are expressed as the mean and standard deviation. A student-T test was used to compare the LDH and non-LDH groups.

LDH: Lumbar disc herniation, BMD: Bone mineral density.

Lumbar disc herniation: bone mineral density relationship

The LDH group had higher lumbar and femoral neck BMD, T, and Z scores compared to the non-LDH group (p<0.05) (Table 1). In binary logistic regression analysis, the covariate-adjusted model showed that higher age, higher lumbar L1-L4 BMD, and higher femoral neck BMD were significantly associated with an increased risk of LDH (p<0.05) (Table 2).

Variable	Odds ratio (95% CI)	P-value
Age, years (per 1 year increase)	1.08 (1.05-1.12)	<0.001
BMI, kg/m² (per 1 kg/ m² increase)	0.98 (0.93-1.02)	0.332
L1-4 BMD (per 0.1 g/cm² increase)	1.35 (1.11-1.64)	0.003
Femur neck BMD (per 0.1 g/cm² increase)	1.32 (1.01-1.72)	0.040

TABLE 2: Factors associated with lumbar disc herniation and binary logistic regression analysis results (the covariate-adjusted model).

Notes: n=513. Covariates included in the model for statistical analysis are: age, BMI, L1-4 BMD, and Femur neck BMD.

BMI: Body mass index, BMD: Bone mineral density.

The relationship between lumbar disc herniation and T-scores

The proportion of patients with a normal L1-L4 T-score was 58.4% in the LDH group and 41.1% in the non-LDH group (p=0.001, OR 95%CI: 2.01 [1.30-3.10]). The proportions of patients with a normal femoral neck T-score were 59.6% and 48.6% in the LDH group and non-LDH group, respectively (p=0.041, OR 95% CI: 1.56 [1.02-2.39]) (Table 3).

	LDH group (n=406)	Non-LDH group (n=107)	Odds ratio (95% CI)	P-value
Normal (≥1) L1-L4 T-score	58.4%	41.1%	2.01 (1.30-3.10)	0.001
Normal (≥1) femur neck T-score	59.6%	48.6%	1.56 (1.02-2.39)	0.041

TABLE 3: Relationship between LDH and T-scores.

Notes: The Chi-Square test was used to compare the LDH and non-LDH groups.

LDH: lumbar disc herniation.

Analysis of patients according to the number of herniated segments

There was no association between the single and multi-level herniation groups in terms of L1-4 and femoral neck BMD (p=0.760 and 0.435, respectively).

Patients with multi-level herniation had significantly higher rates of obesity (BMI \geq 30) compared with single-level herniation (58.0% vs. 47.0%; p=0.031). However, in terms of obesity rates, the LDH group and the non-LDH group were statistically similar (53.9% vs. 54.2%; p=0.961).

Discussion

While the effect of BMD on disc degeneration and osteoarthritis has been widely observed in previous studies [16-20], the LDH-BMD relationship has been understudied. Previous studies were conducted in a young, healthy adult population collected from another study or in young patients with a small sample size [10,11]. We evaluated postmenopausal women who suffer from low back pain, and our results indicated that, after adjusting for covariates, higher lumbar BMD and higher femoral neck BMD were significantly associated with an increased risk of LDH (p<0.05).

Lumbar end plates are the main anatomical structures that carry loads from bone to disc. Stronger bones mean more load from end plates to intervertebral discs [9]. These anatomical regions also contain small vessels for the supply of the disc. Therefore, the morphology, content, and density of these plates are crucial for the integrity of the discs [21]. Structural changes in the end plates can cause higher pressures on the disc and, thus, herniation of the disc.

Several studies have reported that osteophytes and end plate sclerosis may cause a false increase in DXA measurements [22-24]. Therefore, some recent studies have used quantitative computerized tomography instead of DXA and measured BMD from the lumbar trabecular bone region [10,25]. In the present study, we evaluated both L1-L4 and femoral neck BMD using the DXA technique to better interpret the relationship between LDH and the systemic BMD. Both L1-L4 and femoral neck BMD were significantly associated with LDH (p<0.05).

The incidence of LDH has been shown to increase with age [7,10,11]. In the present study, consistent with previous literature, there was a positive correlation between the age of the patients and LDH (p=0.001).

In a previous study, BMI was significantly higher in patients with \geq 2 herniated segments than in healthy adults [10]. In the present study, patients with multi-level herniation had significantly higher rates of obesity (BMI \geq 30) compared with single-level herniation (58.0% vs. 47.0%; p=0.031). However, in terms of obesity rates, the LDH group and the non-LDH group were statistically similar (53.9% vs. 54.2%; p=0.961). Despite the findings of the present study suggesting that obesity may be associated with multi-level LDH, we must emphasize that an association is not always causal, and patients may develop obesity because of reduced activity due to their low back pain.

Consistent with previous literature [10], the number of herniated segments was not significantly associated with BMD in our study. Although we did not find any significant differences between subgroups, the observation needs to be further investigated by longitudinal studies. The reason for the lack of statistically significant differences between multi-level LDH and single-level LDH subgroups may be related to the initiation of treatment after the diagnosis of the first LDH in many patients. This may prevent the development of multi-level LDH in these patients. This issue can be better evaluated with longitudinal studies.

LDH is one of the most common causes of low back pain [26]. LDH was present in 79.1% of patients with low back pain in the present study. In a previous study, 66.4% of patients with low back pain had LDH [27]. However, in their study, the population was younger, and the patients included were between 20 and 64

years of age [27].

Differently from previous studies regarding LDH, we selected our study group from postmenopausal women. With the rapid decrease in estrogen levels, menopause both causes rapid loss of bone mass as well as accelerated lumbar disc degeneration [28,29]. As we only included postmenopausal patients, we minimized the effect of estrogen level differences on outcomes. Moreover, we evaluated the femoral neck BMD in addition to the L1-L4 BMD. This prevented a possible misinterpretation of BMD levels due to lumbar osteophytes and end plate sclerosis. However, this study has several limitations. The first limitation was the retrospective cross-sectional evaluation. The second limitation was that some variables were not available, including smoking status and occupational factors. Third, although both the femoral neck and L1-L4 DXA were assessed, quantitative computerized tomography was not used in this study to eliminate possible false increases in the L1-L4 BMD levels.

Conclusions

Higher lumbar bone mineral density and higher femoral neck bone mineral density were found to be associated with lumber disc herniation in postmenopausal women with low back pain. These results suggest that bone mineral density assessment may be useful in clinical practice to determine which patients are at higher risk of lumbar disc herniation. According to this assessment, preventive methods such as activity modification, exercise programs, and frequent radiological follow-up can be recommended to patients before lumbar disc herniation develops or progresses. Considering the retrospective cross-sectional analysis, further longitudinal and prospective studies are needed to confirm the results of the current study.

Additional Information

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. Bursa City Hospital Clinical Research Ethics Committee issued approval 2022-1/23. This study was approved by the Institutional Review Board (Approval Number: 2022-1/23). Animal subjects: All authors have confirmed that this study did not involve animal subjects or tissue. Conflicts of interest: In compliance with the ICMJE uniform disclosure form, all authors declare the following: Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. Financial relationships: All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. Other relationships: All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

- Zhang J, Zhao F, Wang FL, et al.: Identification of lumbar disc disease hallmarks: a large cross-sectional study. Springerplus. 2016, 5:1973. 10.1186/s40064-016-3662-7
- 2. Ropper AH, Zafonte RD: Sciatica. N Engl J Med. 2015, 372:1240-8. 10.1056/NEJMra1410151
- 3. Frymoyer JW: Back pain and sciatica. N Engl J Med. 1988, 318:291-300. 10.1056/NEJM198802043180506
- 4. Huang W, Qian Y, Zheng K, Yu L, Yu X: Is smoking a risk factor for lumbar disc herniation? . Eur Spine J. 2016, 25:168-76. 10.1007/s00586-015-4103-y
- Yang X, Guo X, Huang Z, et al.: CHRNA5/CHRNA3 gene cluster is a risk factor for lumbar disc herniation: a case-control study. J Orthop Surg Res. 2019, 14:243. 10.1186/s13018-019-1254-2
- Mateos-Valenzuela AG, González-Macías ME, Ahumada-Valdez S, Villa-Angulo C, Villa-Angulo R: Risk factors and association of body composition components for lumbar disc herniation in Northwest, Mexico. Sci Rep. 2020, 10:18479. 10.1038/s41598-020-75540-5
- Kim YK, Kang D, Lee I, Kim SY: Differences in the incidence of symptomatic cervical and lumbar disc herniation according to age, sex and national health insurance eligibility: a pilot study on the disease's association with work. Int J Environ Res Public Health. 2018, 15: 10.3390/ijerph15102094
- 8. Yang Z, Griffith JF, Leung PC, Lee R: Effect of osteoporosis on morphology and mobility of the lumbar spine. Spine (Phila Pa 1976). 2009, 34:E115-21. 10.1097/BRS.0b013e3181895aca
- Tosun O, Fidan F, Erdil F, Tosun A, Karaoğlanoğlu M, Ardıçoğlu O: Assessment of lumbar vertebrae morphology by magnetic resonance imaging in osteoporosis. Skeletal Radiol. 2012, 41:1583-90. 10.1007/s00256-012-1435-0
- Geng J, Wang L, Li Q, et al.: The association of lumbar disc herniation with lumbar volumetric bone mineral density in a cross-sectional Chinese study. Diagnostics (Basel). 2021, 11:10.3390/diagnostics11060938
- Keser N, Atici A, Celikoglu E, Akpinar P, Ramazanoglu AF, Aktas İ: Effect of bone mineral density on lumbar discs in young adults: A case-control study. Medicine (Baltimore). 2017, 96:e7906. 10.1097/MD.000000000007906
- 12. Del Grande F, Maus TP, Carrino JA: Imaging the intervertebral disk: age-related changes, herniations, and radicular pain. Radiol Clin North Am. 2012, 50:629-49. 10.1016/j.rcl.2012.04.014
- Hou Y, Luo Z: A study on the structural properties of the lumbar endplate: histological structure, the effect of bone density, and spinal level. Spine (Phila Pa 1976). 2009, 34:E427-33. 10.1097/BRS.0b013e3181a2ea0a
- Mattei TA: Osteoporosis delays intervertebral disc degeneration by increasing intradiscal diffusive transport of nutrients through both mechanical and vascular pathophysiological pathways. Med Hypotheses. 2013, 80:582-6. 10.1016/j.mehv.2013.01.030
- 15. Fardon DF, Williams AL, Dohring EJ, Murtagh FR, Gabriel Rothman SL, Sze GK: Lumbar disc nomenclature:

- version 2.0: Recommendations of the combined task forces of the North American Spine Society, the American Society of Spine Radiology and the American Society of Neuroradiology. Spine J. 2014, 14:2525-45. 10.1016/j.spinee.2014.04.022
- Zhou L, Li C, Zhang H: Correlation between bone mineral density of different sites and lumbar disc degeneration in postmenopausal women. Medicine (Baltimore). 2022, 101:e28947. 10.1097/MD.000000000028947
- Zhang Y, Patiman, Liu B, Zhang R, Ma X, Guo H: Correlation between intervertebral disc degeneration and bone mineral density difference: a retrospective study of postmenopausal women using an eight-level MRIbased disc degeneration grading system. BMC Musculoskelet Disord. 2022, 23:833. 10.1186/s12891-022-05793-w
- Salo S, Leinonen V, Rikkonen T, et al.: Association between bone mineral density and lumbar disc degeneration. Maturitas. 2014, 79:449-55. 10.1016/j.maturitas.2014.09.003
- Wang YX, Griffith JF, Ma HT, et al.: Relationship between gender, bone mineral density, and disc degeneration in the lumbar spine: a study in elderly subjects using an eight-level MRI-based disc degeneration grading system. Osteoporos Int. 2011, 22:91-6. 10.1007/s00198-010-1200-y
- Livshits G, Ermakov S, Popham M, Macgregor AJ, Sambrook PN, Spector TD, Williams FM: Evidence that bone mineral density plays a role in degenerative disc disease: the UK Twin Spine study. Ann Rheum Dis. 2010, 69:2102-6. 10.1136/ard.2010.131441
- Poureisa M, Daghighi MH, Mesbahi S, Hagigi A, Fouladi DF: End plate disproportion and degenerative disc disease: a case-control study. Asian Spine J. 2014, 8:405-11. 10.4184/asj.2014.8.4.405
- 22. Pye SR, Reid DM, Adams JE, Silman AJ, O'Neill TW: Radiographic features of lumbar disc degeneration and bone mineral density in men and women. Ann Rheum Dis. 2006, 65:234-8. 10.1136/ard.2005.038224
- Atalay A, Kozakcioglu M, Cubuk R, Tasali N, Guney S: Degeneration of the lumbar spine and dual-energy Xray absorptiometry measurements in patients without osteoporosis. Clin Imaging. 2009, 33:374-8.
 10.1016/j.clinimag.2008.12.005
- Ding Y, Chen JY, Yang JC, Li RY, Yin YJ, Chen JT, Zhu QA: Disc degeneration contributes to the denser bone
 in the subendplate but not in the vertebral body in patients with lumbar spinal stenosis or disc herniation.
 Spine J. 2023, 23:64-71. 10.1016/j.spinee.2022.09.010
- Güngör Ö, Gezer NS, Özdamarlar U, Balcı A: The effect of bone mineral density on development of Schmorl's nodes in young patients. Acta Orthop Traumatol Turc. 2020, 54:287-92.
 10.5152/j.aott.2020.03.577
- Takahashi H, Aoki Y, Inoue M, et al.: Characteristics of relief and residual low back pain after discectomy in patients with lumbar disc herniation: analysis using a detailed visual analog scale. BMC Musculoskelet Disord. 2021, 22:167. 10.1186/s12891-021-04015-z
- Shambrook J, McNee P, Harris CE, Kim M, Sampson M, Palmer KT, Coggon D: Clinical presentation of low back pain and association with risk factors according to findings on magnetic resonance imaging. Pain. 2011, 152:1659-65. 10.1016/j.pain.2011.03.011
- Wáng YX, Griffith JF, Deng M, Yeung DK, Yuan J: Rapid increase in marrow fat content and decrease in marrow perfusion in lumbar vertebra following bilateral oophorectomy: an MR imaging-based prospective longitudinal study. Korean J Radiol. 2015, 16:154-9, 10:3348/kir 2015, 16:1, 154
- Wang YX, Griffith JF: Effect of menopause on lumbar disk degeneration: potential etiology . Radiology. 2010, 257:318-20. 10.1148/radiol.10100775