# Is There a Relationship Between Joint Hypermobility and Lumbar Disc Degeneration and Also Low Back Pain? A Multidisciplinary Clinical Study

Eklem Hipermobilitesi ile Lomber Disk Dejenerasyonu ve Bel Ağrısı Arasında İlişki Var mı? Bir Multidisipliner Klinik Çalışma

ABSTRACT

**Objective:** Hypermobility (HM) is a hereditary or acquired condition of connective tissue in which the synovial joints have range of movement beyond their normal limits and its effects on disc degeneration (DD) is not fully known. In our study, We aimed to reveal the relationship between joint hipermobility, lumbar DD and also low back pain.

**Method:** The cases aged between 20-50 years with low back and/or leg pain were included in the study. Their lumbar MRIs were evaluated using Pfirrmann grading system. For the cases meeting the study criteria, the Beighton score was used for evaluating the generalized joint laxity, and the 1998 Brighton criteria were used to assess the benign joint hypermobility syndrome. The cases were also evaluated prospectively for pain by using the visual analog scale (VAS) and for disability using the Oswestry disability index (ODI).

**Results:** Hundred and seventy-two cases with a mean age of 36.82±7.62 years including 112 female (65.1%), and 60 male (34.9%) patients met the inclusion and the exclusion criteria, While 14 % of cases had joint HM, and HM was not encountered in 86% of them. There was no statistically significant difference in the values of Pfirrmann grade parameters at all lumbar disc levels between the groups with and without HM (p> 0.05). Likewise, there was no statistically significant difference between the groups concerning the VAS and the ODI values (p>0.05).

**Conclusion:** These findings suggest that HM may not be a risk factor for increased lumbar DD grades, VAS and ODI scores in patients aged between 20-50 years.

**Keywords:** low back pain, benign joint hypermobility syndrome, disc degeneration, hypermobility, generalized joint hypermobility, lumbar spine

### ÖZ

Amaç: Hipermobilite (HM) sinovial eklemlerin normal sınırının ötesinde hareket yeteneğinin olduğu bağ dokusunun çoğunlukla kalıtsal ya da edinilmiş bir durumu olup lomber disk dejenerasyonu (DD) üzerine etkileri tam olarak bilinmemektedir. Çalışmamızda eklem hipermobilitesi ile lomber DD ve ayrıca bel ağrısı arasındaki ilişkinin araştırılması amaçlandı.

Yöntem: Bel ve/veya bacak ağrısı yakınması olan ve son altı ay içinde çekilmiş lomber MRİ'lerinde DD saptanılan 20-50 yaş arasındaki olgular çalışmaya dahil edildiler. DD değerlendirimi Pfirrmann derecelendirme sistemine göre yapıldı. Çalışma kriterlerine uyan tüm olgular Beighton skoru kullanılarak generalize eklem hipermobilitesi yönünden değerlendirilmiş, benign eklem hipermobilitesi sendromu değerlendiriminde ise, 1998 Brighton kriterleri kullanılmıştır. Ayrıca, olguların ağrı derecesi değerlendiriminde vizüel analog skala (VAS) kullanılmış, dizabilite değerlendirimi ise Oswestry dizabilite indeksi (ODI) kullanılarak prospektif olarak yapılmıştır.

Bulgular: İçerleme ve dışlama kriterlerine uyan 172 olgunun 112'si kadın (%65,1), 60'ı erkek (%3.9), olguların yaşları ortalaması ise 36,82±7,62 yıl idi. Bu olguların 24'ünde hipermobilite varken (%14), 148'inde HM'ye rastlanılmadı (%86). HM görülenler ve görülmeyenler arasında tüm lomber disk düzeylerinde Pfirrmann grade parametreleri değerleri açısından istatistiksel olarak anlamlı bir farklılık bulunmadı (p>0.05). Aynı şekilde gruplar arasında VAS değerleri ve ODİ değerleri açısından da istatistiksel olarak anlamlı bir farklılık yoktu (p>0.05).

Sonuç: Bu sonuç 20-50 yaş aralığındaki olgularda, normal şartlar altında, HM'nin lomber DD'u ile VAS ve ODİ artışında bir risk faktörü olmayabileceğini düşündürmüştür.

Anahtar kelimeler: bel ağrısı, benign eklem hipermobilite sendromu, disk dejenerasyonu, hipermobilite, jeneralize eklem hipermobilitesi, lomber omurga



© Telif hakkı İstanbul Kanuni Sultan Süleyman Eğitim ve Araştırma Hastanesi'ne aittir. Logos Tıp Yayıncılık tarafından yayınlanmaktadır. Bu dergide yayınlanan bütün makaleler Creative Commons Atıf-Gayri Ticari 4.0 Uluslararası Lisansı ile lisanslanmıştır.

© Copyright İstanbul Kanuni Sultan Süleyman Research and Training Hospital. This journal published by Logos Medical Publishing. Licenced by Creative Commons Attribution-NonCommercial 4.0 International (CC BY-NC 4.0) Neşe Keser @ Esin Derin Çiçek @ Arzu Atıcı @ Pınar Akpınar @ Özge Gülsüm İlleez @ Ahmet Eren Secen @

Received: 8 May 2019 Accepted: 20 January 2020 Publication date: 31 January 2020

**Cite as:** Keser N, Çiçek ED, Atıcı A, Akpınar P, İlleez ÖG, Seçen AE. Is there a relationship between woint hypermobility and lumbar disc degeneration and also low back pain? A multidisciplinary cilincal study. IKSSTD 2020;12(1):1-6.

Neşe Keser

İstanbul Fatih Sultan Mehmet EAH Beyin Cerrahisi Bölümü E5 Karayolu Üzeri Ataşehir 34752 İstanbul - Türkiye Makeser@gmail.com ORCID: 0000-0002-6024-8855

E. D. Çiçek 0000-0002-0391-3003 İstanbul Fatih Sultan Mehmet EAH Radyoloji Bölümü İstanbul - Türkiye

A. Atıcı 0000-0002-2069-4017 P. Akpınar 0000-0002-7796-0750 Ö.G. İlleez 0000-0002-3824-4428 İstanbul Fatih Sultan Mehmet EAH Fizik Tedavi ve Rehabilitasyon Bölümü İstanbul - Türkiye

A. E. Seçen 0000-0003-2185-020X İstanbul Fatih Sultan Mehmet EAH Beyin Cerrahisi Bölümü İstanbul - Türkiye

## **INTRODUCTION**

Hypermobility (HM) is an observed phenomenon which can be inherited or rarely acquired and in which the synovial joints can have a range of motion (ROM) beyond the reasonable limits. Its prevalence varies by sex, age, and racial characteristics, but can reach 15% in adults and 30% in childhood <sup>(1)</sup>. HM is a serious condition since it is associated with chronic pain complaints, premature osteoarthrosis (OA), osteoporosis (OP), vascular and spinal disorders <sup>(2)</sup> and cannot be detected if it is not investigated <sup>(3)</sup>. Although in childhood and adolescence HM is referred to as generalized joint laxity (GJL) and since it is associated with the clinic in adults, it is referred to as benign joint hypermobility syndrome (BJHS), and different diagnostic criteria have been used <sup>(4)</sup>. While Beighton criteria <sup>(3,5)</sup> are used to detect GJL (Appendix 1), Brighton criteria are used for the diagnosis of BJHS (Appendix 2)<sup>(6)</sup>. The Brighton criteria differentiate BJHS from other connective tissue disorders and the rare causes of HM such as Ehlers-Danlos syndrome, Marfan's syndrome and osteogenesis imperfecta<sup>(3)</sup>.

#### Appendix 1. Beighton hypermobility score\*

Right	Left
1	1
1	1
1	1
1	1
1	
9	
	<b>Right</b> 1 1 1 1 1 9

#### Total

\* Note: A threshold cutoff value of four in total nine points was used to classify generalized joint laxity.

#### Appendix 2. Revised 1998 Brighton criteria.

#### Major criteria

- Beighton score of 4/9 or greater (either currently or historically)
- Arthralgia for longer than three months in 4 or more joints

#### Minor criteria

- Beighton score of 1, 2, or 3/9 (0, 1, 2, or 3/9 if aged >50 years)
- Arthralgia in one to three joints or back pain for more than three months, spondylolysis, spondylolysis/spondylolisthesis
- Dislocation/subluxation in more than one joint or one joint on more than one occasion
- Soft tissue rheumatism with >3 lesions (e.g., bursitis, tenosynovitis, epicondylitis)
- Marfanoid habitus (tall, slim, span/height ratio >1.03, upper/lower segment ratio <0.89, arachnodactyly (positive Steinberg/wrist signs)
- Cutaneous lesions: striae, hyperextensibility, thin skin
- Eye signs: Drooping eyelids, myopia, or antimongoloid slant
- Varicose veins or a hernia or uterine/rectal prolapse

**Required diagnostic criteria for hypermobility:** Two major criteria or, one major and two minor criteria or, four minor criteria or, the presence of BJHS in first-degree relatives and two minor criteria.

Low back pain (LBP) is one of the severe health problems and, lumbar disc degeneration (LDD) is one of tts most common causes <sup>(7)</sup>. Although many factors have been identified that aggravate disk degeneration (DD) <sup>(8)</sup> the effect of HM on DD and LBP is not fully understood. Currently, the MRI is the most valid noninvasive examination tool used to assess the lumbar spine morphologically and to detect the association of these morphological changes with LBP <sup>(9)</sup>. Pfirrmann grading (PG) system is comprehensive, with intra- and interobserver reliability sufficient to discriminate between the different grades <sup>(10)</sup>.

We aimed to investigate the effect of HM on DD grades at all levels of the lumbar IVDs and also LBP, in the non-operated and the non-traumatized cases. As far as we know, this is the first study in English literature which is used the PG for this purpose.

## **MATERIAL and METHODS**

TThis study was conducted with the approval of the ethics committee of Fatih Sultan Mehmet Training and Research Hospital (decision date, and number: 11.23.2017/112). The informed consent form was obtained from all patients included in the study. The cases with waist and leg pain, aged between 20-50 years, were evaluated prospectively. The effects of the HM on the lumbar IVDs in the MRI were compared with the case-control study. Patients with a history of lumbar surgery, tumor, rheumatic disease or infection, stroke, dementia or diseases requiring intensive care, and women in their menopausal period were not included in the study. During the

N. Keser ve ark., Is There a Relationship Between Joint Hypermobility and Lumbar Disc Degeneration and Also Low Back Pain? A Multidisciplinary Clinical Study

#### Table 1. Evaluation of demographic data according to the presence of hypermobility.

		Hypermobility		
		Present n (%)	Absent n (%)	р
Age (year)	20-30 31-40 41-50	8 (33) 10 (42) 6 (25)	29 (20) 62 (42) 57 (38)	<sup>1</sup> 0,241
Gender	Male Female	5 (21) 19 (79)	55 (37) 93 (63)	<sup>2</sup> 0,185
dof	Working Not Working	10 (42) 14 (58)	84 (57) 64 (43)	<sup>2</sup> 0,247
Body Mass Index	≤25 >25	11 (46) 13 (54)	51 (35) 97 (65)	<sup>2</sup> 0,397
Fingertip-to-Floor	Touching Not Touching	12 (50) 12 (50)	98 (66) 50 (34)	<sup>2</sup> 0,192
Waist Circumference (cm)	Male ≤102 >102 Female ≤88 >88	5 (100) 0 (0) 8 (42) 11 (58)	38 (69) 17 (31) 31 (33) 52 (57)	<sup>3</sup> 0,309 <sup>2</sup> 0,640
Smoking	>88 <5 ≥5	11 (58) 17 (71) 7 (29)	62 (67) 90 (61) 57 (39)	<sup>2</sup> 0,500
Previous Illness	Preseent Absent	12 (50) 12 (50)	62 (42) 86 (58)	<sup>2</sup> 0,602
Consanguineous Marriage	Present Absent	19 (79) 5 (21)	117 (79) 31 (21)	²1,000

<sup>1</sup> Chi-square Test

<sup>2</sup> Continuity (Yates) Correction

<sup>3</sup> Fisher's Exact Test

study period between December 2017 and May 2018, 172 of 2088 patients complied with our criteria. The patients filled the questionnaires, and their demographic data, waist circumference values, and fingertip-to-floor (FTF) distances were recorded, body mass indexes (BMIs) were calculated, and lumbar MRIs obtained within the last six months were evaluated. The evaluation HM was made using the Beighton score (Appendix 1), and the revised Brighton (Brighton 1998) criteria (Appendix 2) <sup>(5,6)</sup>. A cut-off value of four in a total of nine points was used to classify GJL. A 1: 6 equivalence design was used to compare the cases with (Group A) and without hypermobile (Group B) joints. The severity of pain was assessed using the visual analog scale (VAS) (11), and the clinical outcome using the Oswestry Disability Index (ODİ) <sup>(12)</sup>. The MRİs were evaluated blindly and all lumbar IVD levels, and the severity of DD was classified using the PG system. In this grading system, evaluation of DD degree is made at five grades (G I-V) considering the disk density and height <sup>(10)</sup>. While grade I corresponds to healthy intervertebral disc (homogenous bright white structure), grades II and III to mild, grade IV to moderate and, grade V to severe degeneration of IVD, accordingly. A point was given for every grade in a direct proportion to the grade (1 point to grade 1 degeneration, 5 points to grade 5 degeneration), the mean± standard deviations of these five grades were calculated and the values were compared between the two groups.

For statistical analysis, data were evaluated using the SPSS for Windows version 22.0 software (IBM Corp., Armonk, NY, USA) program. Descriptive statistical methods (mean, standard deviation, frequency) were used for the evaluation of data. Chi-square

		Hypermobility		
		Present n (%)	Absent n (%)	р
Visual Analog Scale	≤5	14 (58)	80 (54)	<sup>1</sup> 0,865
	>5	10 (42)	68 (46)	
Oswestry Disability Index	≤30	1 (4)	12 (8)	<sup>2</sup> 0,474
	31-50	12 (50)	51 (35)	
	51-75	9 (38)	58 (39)	
	76-100	2 (8)	27 (18)	
Brighton Criteria	Absent	7 (29)	102 (69)	<sup>1</sup> 0,000*
	Present	17 (71)	46 (31)	

Table 2. Evaluation of visual analog scale, Oswestry disability index parameters and Brighton Criteria according to the presence of hypermobility.

<sup>1</sup> Continuity (Yates) Correction <sup>2</sup> Fisher Freeman Halton Test

\*p<0.05

test, Fisher's Exact test, Fisher Freeman Halton test and Yates continuity correction test were used for the comparison of qualitative data. A p value of <0.05 was accepted as statistically significant.

#### RESULTS

Of the 172 cases that met the inclusion and exclusion criteria, 60 cases (34.9%) were male and 112 cases (645.1%) were female, and the mean age was 36.82± 7.62 years. HM was detected in 24 (13.8%), and was not detected in 148 (86.2 %) cases There was no difference between these groups regarding age, gender, smoking status, previous illnesses, WC, BMI, FTF distance, and presence of consanguineous marriage (p>0.05) (Table 1). Also, there was no statistically significant intergroup difference as for VAS scores and ODI values (Table 2).

## DISCUSSION

At present, MRI is the best noninvasive examination tool is used to evaluate lumbar vertebrae <sup>(9)</sup>. It has been reported that reduced signal intensity is related to dehydration and decreased proteoglycan content, but the latter issue continues to be the subject of debate <sup>(13)</sup>. Pfirrmann et al. used both signal intensity and disc height categories in their classification and, showed that the low signal intensity in the MRI is mostly associated with dehydration <sup>(10)</sup>.

All functional spinal unit elements, such as IVDs, facet joints, bony structures, and spinal ligaments

may undergo degenerative changes, but mainly IVDs are responsible for the instability  $^{(14,15)}$ . In autopsies of the patients under 50 years of age, 85-95% of these cases were reported to have LDD, suggesting that the DD may also occur other than the age factor such as mechanical or metabolic injury  $^{(7,8)}$ .

It has been claimed that the deteriorated proprioception in the hypermobile cases increases the susceptibility to sports injuries and the traumatized joints lead to more serious problems (3,16,17). Some of the vertebral joints lie between relatively fixed and movable regions. While the movement in one region increases as a result of HM which may rapidly lead to faster and more severe degeneration of the joint which lies between the relatively less mobile and the hypermobile parts. Likewise, this situation is seen both in the presence of lumbosacral transitional vertebra and in the adjacent segment disease (ASD), which is a complication of the lumbar stabilization <sup>(18)</sup>. It has been reported that restriction of movement with rigid posterior lumbar interbody fusion in the cases with HM leads to overload in the adjacent segment, and in these cases, ASD is seen more frequently than the others <sup>(19)</sup>. Therefore, before the spinal surgery, patients should be assessed regarding HM and rigid stabilization should be avoided in these cases.

Although HM is not commonly referred to as a disease, the predominant complaint is usually widespread and long-lasting pain, and it is also the most common cause of unexplained joint pain (3). But, the relationship between increased lumbar segmental motion and LDD and also LBP is still controversial (20-23). Our results suggest that HM does not lead to increase in DD, VAS, and ODI values (Table 2) which can be an indicator of the importance of maintaining the joint motion harmonization especially for the cases who have hypermobile joints. Since the evaluation of HM is difficult in the elderly, we included the men under 50 years old and premenopausal women in our study. Also, we selected the cases having the same demographic characteristics such as nonobese individuals without history of trauma (Table 1). Our study results also suggest that hypermobile cases should be informed about to start lumbar paravertebral muscle strengthening exercises at an earlier age and, avoid both obesity and any trauma.

One of the limitations of our study is that radiological evaluations are made only on IVDs. To elucidate the relationship between HM and LBP, radiological evaluations should also be performed on facet joints and posterior ligamentous complexes. Other limitations are limited number of hypermobile cases and the large age range. Studies performed on greater number of cases in every decade can give healthier results.

**Ethics Committee Approval:** Approval of the Clinical Research Ethics Committee of SBU Istanbul Fatih Sultan Mehmet Training and Research Hospital was obtained (decision number: 23.11.2017 / 112). **Conflict of Interest:** There is no conflict of interest.

**Funding:** There is no financial support.

**Informed Consent:** Informed consent was obtained from each patient.

# REFERENCES

- Baeza-Velasco C, Gély-Nargeot MC, Pailhez G, Vilarrasa AB. Joint hypermobility and sport: a review of advantages and disadvantages. Curr Sports Med Rep. 2013;12:291-5. https://doi.org/10.1249/JSR.0b013e3182a4b933
- Murray KJ. Hypermobility disorders in children and adolescents. Best Pract Res Clin Rheumatol. 2006;20:329-51. https://doi.org/10.1016/j.berh.2005.12.003
- Simpson MR. Benign joint hypermobility syndrome: evaluation, diagnosis, and management. J Am Osteopath Assoc. 2006;106:531-6.
- Ross J, Grahame R. Joint hypermobility syndrome. BMJ. 2011;342:c7167.

https://doi.org/10.1136/bmj.c7167

5. Beighton P, Solomon L, Soskolne CL. Articular mobility in an

African population. Ann Rheum Dis. 1973;32:413-8. https://doi.org/10.1136/ard.32.5.413

- Grahame R, Bird HA, Child A. The revised (Brighton 1998) criteria for the diagnosis of benign joint hypermobility syndrome (BJHS). J Rheumatol. 2000;27:1777Y9.
- Middendorp M, Vogl TJ, Kollias K, Kafchitsas K, Khan MF, Maataoui A. Association between intervertebral disc degeneration and the Oswestry Disability Index. J Back Musculoskelet Rehabil. 2017;30:819-23. https://doi.org/10.3233/BMR-150516
- Hangai M, Kaneoka K, Kuno S, et al. Factors associated with lumbar intervertebral disc degeneration in the elderly. Spine J. 2008;8:732-40.

https://doi.org/10.1016/j.spinee.2007.07.392

 Saleem S, Aslam HM, Rehmani MA, Raees A, Alvi AA, Ashraf J. Lumbar disc degenerative disease: disc degeneration symptoms and magnetic resonance image findings. Asian Spine J. 2013;7:322-34.

https://doi.org/10.4184/asj.2013.7.4.322

- Pfirrmann CW, Metzdorf A, Zanetti M, Hodler J, Boos N. Magnetic resonance classification of lumbar intervertebral disc degeneration. Spine. 2001;26:1873-8. https://doi.org/10.1097/00007632-200109010-00011
- Hawker GA, Mian S, Kendzerska T, French M. Measures of adult pain: Visual Analog Scale for Pain (VAS Pain), Numeric Rating Scale for Pain (NRS Pain), McGill Pain Questionnaire (MPQ), Short-Form McGill Pain Questionnaire (SF-MPQ), Chronic Pain Grade Scale (CPGS), Short Form-36 Bodily Pain Scale (SF-36 BPS), and Measure of Intermittent and Constant Osteoarthritis Pain (ICOAP). Arthritis Care Res (Hoboken). 2011;63 Suppl 11:S240-52. Review. https://doi.org/10.1002/acr.20543
- Fairbank JC, Pynsent PB. The Oswestry Disability Index. Spine (Phila Pa 1976). 2000;25:2940-52; discussion 2952. https://doi.org/10.1097/00007632-200011150-00017
- Sambrook PN, MacGregor AJ, Spector TD. Genetic influences on cervical and lumbar disc degeneration: a magnetic resonance imaging study in twins. Arthritis Rheum. 1999;42:366-72.

https://doi.org/10.1002/1529-0131(199902)42:2<366::AID-ANR20>3.0.CO;2-6

- Inoue N, Espinoza Orías AA. Biomechanics of intervertebral disk degeneration. Orthop Clin North Am. 2011;42:487-99, vii. https://doi.org/10.1016/j.ocl.2011.07.001
- Kirkaldy-Willis WH, Farfan HF. Instability of the lumbar spine. Clin Orthop Relat Res. 1982;(165):110-23. https://doi.org/10.1097/00003086-198205000-00015
- 16. Olivier B, Taljaard T, Burger E, et al. Which Extrinsic and Intrinsic Factors are Associated with Non-Contact Injuries in Adult Cricket Fast Bowlers? Sports Med. 2016;46:79-101. https://doi.org/10.1007/s40279-015-0383-y
- 17. Kim HJ, Yeom JS, Lee DB, Kang KT, Chang BS, Lee CK. Association of benign joint hypermobility with spinal segmental motion and its clinical implication in active young males. Spine. 2013;38:E1013-9.

https://doi.org/10.1097/BRS.0b013e31828ffa15 18. Farshad-Amacker NA, Herzog RJ, Hughes AP, Aichmair A,

- Farshad-Amacker NA, Herzog RJ, Hugnes AP, Alchmair A, Farshad M. Associations between lumbosacral transitional anatomy types and degeneration at the transitional and adjacent segments. Spine J. 2015;15:1210-6. https://doi.org/10.1016/j.spinee.2013.10.029
- Lee SM, Lee GW. The impact of generalized joint laxity on the clinical and radiological outcomes of single-level posterior lumbar interbody fusion. Spine J. 2015;15:809-16.

https://doi.org/10.1016/j.spinee.2014.12.013

- Kraus VB, Li YJ, Martin ER, et al. Articular hypermobility is a protective factor for hand osteoarthritis. Arthritis Rheum. 2004;50:2178-83. https://doi.org/10.1002/art.20354
- Corey JM. Genetic disorders producing compressive radiculopathy. Semin Neurol. 2006;26:515-22. https://doi.org/10.1055/s-2006-951624
- 22. Kim KT, Park SW, Kim YB. Disc height and segmental motion as risk factors for recurrent lumbar disc herniation. Spine (Phila Pa 1976). 2009;34:2674-8.
  - https://doi.org/10.1097/BRS.0b013e3181b4aaac
- 23. Teraguchi M, Yim R, Cheung JP, Samartzis D. The association of high-intensity zones on MRI and low back pain: a systematic review. Scoliosis Spinal Disord. 2018;13:22. https://doi.org/10.1186/s13013-018-0168-9