

Age and sex-based distribution of lumbar multifidus muscle atrophy and coexistence of disc hernia: an MRI study of 2028 patients

Elif Evrim Ekin
Hülya Kurtul Yıldız
Harun Mutlu

PURPOSE

We aimed to investigate the prevalence of lumbar multifidus muscle (LMM) atrophy in patients having mechanical low back pain with and without disc hernia.

METHODS

In total, 2028 lumbar magnetic resonance imaging scans of low back pain patients (age range, 18–88 years) were re-evaluated retrospectively. LMM atrophy was visually assessed in axial sections of L4-L5 and L5-S1 levels.

RESULTS

LMM atrophy prevalence at both levels was significantly higher in subjects ≥ 40 years compared with younger adults ($P < 0.001$). LMM atrophy was significantly more frequent in women than in men ($P < 0.001$). Among patients with low back pain without hernia, LMM atrophy was significantly more frequent than normal muscle ($n=559$ vs. $n=392$; $P < 0.001$). Frequency of LMM atrophy in low back pain patients without disc hernia was 13%. Hernia was more frequent in patients with LMM atrophy compared with patients without atrophy ($P < 0.001$).

CONCLUSION

LMM atrophy is more common in women; its prevalence and severity are observed to increase with advancing age, and disc hernia is found more frequently in individuals with LMM atrophy.

Low back pain is a common health problem. In the general population, 80% of adults experience mechanical low back pain at least once in a lifetime (1). From a biomechanical perspective, the main stabilizer of lumbar spine is lumbar multifidus muscle (LMM), which plays an important role in chronic low back pain (2, 3). Especially, long-term neurologic inhibition after low back trauma and chronic diseases that lead to muscle atrophy may result in replacement of healthy LMM fibrils with adipose tissue (2). Two main findings of muscle atrophy become evident in radiology: shrinkage of muscle volume and increase in the intramuscular fat storage (4). LMM atrophy can be identified in transverse planes of lumbar magnetic resonance imaging (MRI) (5).

There is a strong relationship between low back pain and paraspinal muscle atrophy (6). In the literature, various studies suggested that LMM atrophy is more common among women and adults (3). It is known that muscle atrophy increases with age (6). However, to the best of our knowledge, a large scale study of age and sex-based LMM atrophy in patients with low back pain and its coexistence with disc hernia is lacking in the literature.

In this study, we aimed to investigate the prevalence of LMM atrophy in patients having mechanical low back pain with and without disc hernia.

Methods

Local ethics committee approval was obtained for this study.

In total, 2166 patients (age range, 18–88 years) who underwent MRI due to low back pain between May 2014 and December 2014 were retrospectively re-evaluated. Of the patients, 138 were excluded from the study due to malignancy, intraspinal mass, spondylitis, previous surgery, and structural scoliosis.

LMM atrophy and disc herniation were evaluated in all patients at L4-L5 and L5-S1 levels. LMM has the largest diameter at these levels, allowing for a better evaluation.

From the Departments of Radiology (E.E.E. ✉ eeeoner@gmail.com, H.K.Y.) and Orthopedics and Traumatology (H.M.), Gaziosmanpaşa Taksim Education and Research Hospital, Istanbul, Turkey.

Received 10 July 2015; revision requested 29 July 2015; last revision received 1 September 2015; accepted 29 September 2015.

Published online 01 April 2016.
DOI 10.5152/dir.2015.15307

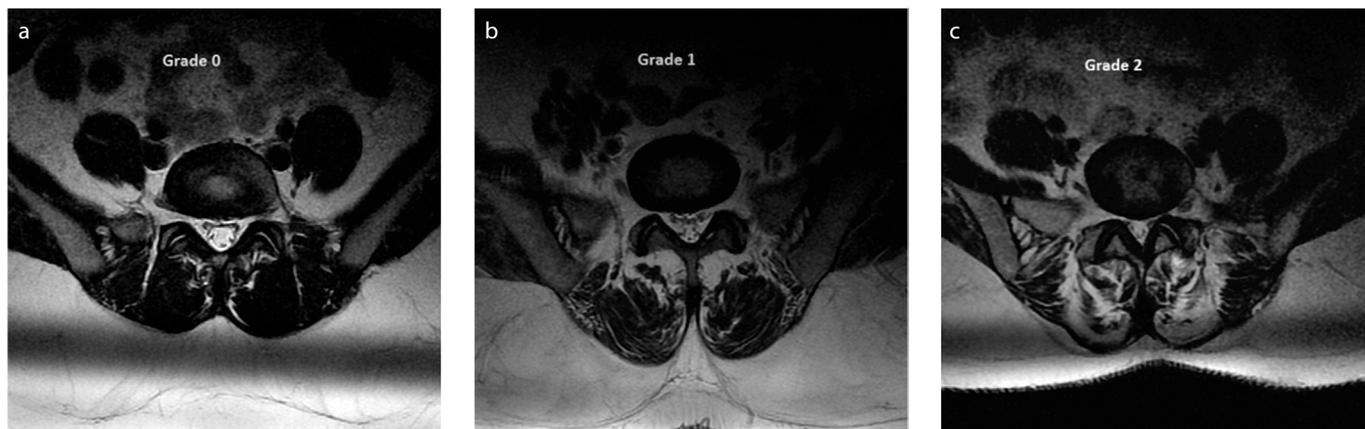


Figure 1. a–c. Examples of lumbar multifidus muscle (LMM) atrophy as seen on axial T2-weighted images. Panel (a) shows normal condition (grade 0); panel (b) shows grade 1 atrophy with slight fat infiltration (10%–50%); panel (c) shows grade 2 atrophy with severe fat infiltration (>50%).

MRI procedure

1.5 T MRI unit (Signa HDxt; General Electric) was used with body surface coil. Lumbar spine was evaluated at L1-S1 levels, and L4-L5 and L5-S1 levels were re-evaluated. Scan sequences included sagittal T1-weighted fast spin-echo (FSE), T2-weighted FSE and an axial T2-weighted FSE (3680/128 repetition time/echo time, 180×256 matrix, 280 mm field of view and 4 mm section thickness, NEX 2).

LMM atrophy was visually examined at L4-L5 and L5-S1 levels. Based on the studies of Parkkola et al. (7) and Kader et al. (8), fatty atrophic changes in LMM were divided into three grades: grade 0, fatty atrophy <10% (Fig. 1a); grade 1, fat infiltration 10%–50% (Fig. 1b); grade 2, fat infiltration >50% (Fig. 1c). All images were evaluated by two experienced radiologists.

Statistical analysis

Normal distribution of the data was tested by univariate Kolmogorov Smirnov test and a histogram graphic was prepared. Normality was obtained by logarithmic transformation of age (log base 10).

Data were presented as mean, standard

Table 1. Distribution of LMM atrophy at L4-L5 and L5-S1 levels in females and males

	Females (n=1263)		Males (n=765)		P
	Grade 0	Grades 1+2	Grade 0	Grades 1+2	
L4-L5 level	796 (63)	467 (37)	586 (76.6)	179 (23.4)	<0.001
L5-S1 level	330 (26.1)	933 (73.9)	367 (48)	398 (52)	<0.001

Data are presented as n (%).
LMM, lumbar multifidus muscle; Grade 0, no atrophy; Grade 1, mild atrophy; Grade 2, advanced atrophy.

deviation, minimum, maximum, frequency, and percent value based on various characteristics. Intergroup comparisons were made by using independent samples t-test or univariate analysis of variance. Post hoc comparisons after ANOVA were made by Tukey HSD test.

Comparison of nominal variables were made by chi-square test. The effects of atrophy and hernia at L4-L5 and L5-S1 levels were tested by two-way analysis of variance.

Two tailed significance level was adjusted to $P < 0.05$. All statistical analyses were performed using the NCSS10 software (NCSS, LLC).

Results

The study group consisted of 1263 women (62.3%) and 765 men (37.7%) with low back pain. The mean age was 43.4 ± 13.7 years (range, 18–88 years).

At L4-L5, 1197 patients had disc hernia and 646 patients had grade 1 or 2 LMM atrophy. At L5-S1, 1077 patients had disc hernia and 1131 patients had grade 1 or 2 LMM atrophy. Hernia was more frequent in patients with LMM atrophy at both levels compared with patients without atrophy ($P < 0.001$, for both levels).

Occurrence of LMM atrophy at L4-L5 level was 37% in women and 23.4% in men,

while occurrence of LMM atrophy at L5-S1 level was 73.9% in women and 52% in men (Table 1). At both levels, LMM atrophy was significantly more frequent in women than in men ($P < 0.001$, for both levels).

LMM atrophy in women and men was significantly more frequent at L5-S1 than at L4-L5 ($P < 0.001$). All patients with atrophy at L4-L5 had atrophy at L5-S1 as well. However, of patients with LMM atrophy at L5-S1, only 33.8% had coexisting atrophy at L4-L5.

Results of LMM atrophy evaluation in subjects <40 years and ≥ 40 years of age are presented in Table 2. LMM atrophy at both L4-L5 and L5-S1 levels was significantly more frequent among men and women ≥ 40 years compared with younger ages ($P < 0.001$).

At L4-L5, LMM atrophy was significantly associated with age ($P < 0.001$), while disc hernia was not ($P = 0.085$); on the other hand, coexistence of hernia and atrophy at L4-L5 was significantly associated with age ($P = 0.048$). At L5-S1, both atrophy and hernia were significantly associated with age ($P < 0.001$), while their coexistence was not ($P = 0.796$).

Table 3 presents the mean ages of patients according to the grade of LMM atrophy at L4-L5 and L5-S1 levels, excluding the patients with disc hernia. At both levels, the mean age was significantly different among

Main points

- The origin of low back pain is not clear, but it is known that lumbar multifidus atrophy (LMM) is strongly associated with low back pain. LMM atrophy may be the only MRI finding in some low back pain patients.
- LMM atrophy is more common in women than in men, and its prevalence and severity increase with age. Forty years of age is especially important for LMM atrophy.
- Disc hernia is more frequent in patients with LMM atrophy.

patients with different grades of LMM atrophy ($P < 0.001$, for both levels). We observed that the severity of LMM atrophy increased with age (Table 3).

When low back pain patients without disc hernia at L5-S1 ($n=951$) were evaluated for LMM atrophy at the same level, grade 0, 1, and 2 LMM atrophy were present in 392, 497, and 62 patients, respectively. Among patients without disc hernia, significantly more patients had LMM atrophy compared with normal LMM ($n=559$ vs. $n=392$; $P < 0.001$).

Discussion

In this study we found that among patients with low back pain, women and patients ≥ 40 years of age were more likely to have LMM atrophy. Disc herniation was frequently found together with LMM atrophy; there were only two patients who had herniation without atrophy. On the other hand, LMM atrophy was present as the only radiologic finding in a significant subset of patients with low back pain ($n=265$).

The underlying etiology and the origin of pain are not clear in most patients with

low back pain (9). In order to diagnose mechanical low back pain, conditions such as malignancy, inflammatory diseases, infectious processes and referred pain of other organs should be excluded (10). Accordingly, we excluded these pathologies to identify patients with mechanical low back pain.

In biomechanical assessment of low back pain, muscular stabilization of "neutral zone" in low back region becomes important. It has been well understood that LMM is the key factor in neutral stabilization (11) and LMM dysfunction is strongly correlated with low back pain (12, 13). Fat involution in LMM muscle leads to muscle dysfunction. The etiology of muscular atrophy includes nutritional disorders, lack of adequate physical activity, immobility, and long-term systemic diseases. Muscle mass starts to decrease progressively after 40 years of age and the reduction in muscle mass is about 8% in each decade (14). In our study, LMM atrophy was significantly higher in subjects ≥ 40 years of age, regardless of the sex. In order to exclude long-term nerve compression-induced LMM atrophy, we

examined patients without disc hernia: the mean ages of Grade 0, 1 and 2 LMM atrophy groups at L5-S1 level were 34.6 ± 10.28 years, 42.90 ± 12.17 years, and 52.21 ± 12.94 years, respectively (Table 3). Severity of LMM atrophy and advancing decades were directly proportional according to our grading system.

Prevalence of LMM atrophy was significantly higher in women than in men. LMM atrophy might be responsible for majority of the mechanical low back pain in women, and it affects female population more than male population.

Kader et al. (8), reported that muscle atrophy in chronic low back pain patients mostly affected the LMM, which is the largest medial muscle in lumbosacral region of lumbar spine. Mobility in lumbar region occurs mostly at L5-S1 level, followed by L4-L5 level (8). Therefore, we assessed LMM muscle at L4-L5 and L5-S1 levels to associate with low back pain. LMM atrophy was significantly more frequent at L5-S1 than L4-L5 in both sexes.

Lumbar disc hernia was found more frequently in patients with LMM atrophy compared with patients without atrophy. However, coexistence of LMM atrophy and disc hernia at L5-S1 level was not associated with age. We found disc sequestration without LMM atrophy in two patients (Fig. 2). On the other hand, 265 patients had isolated LMM atrophy without disc hernia (Figs. 3, 4). In younger patients, lack of muscle atrophy in acute disc hernia may be explained by the longer time period required for fat replacement; but our study is limited as the duration of pain was not included. There were significantly more low back pain patients with grade 1 or 2 LMM atrophy without hernia than patients with normal muscle without hernia. In our study, LMM atrophy at L4-L5 and L5-S1 levels was the single pathologic MRI finding in 13% of patients with mechanical low back pain. Can LMM atrophy be instructive in low back pain patients without disc hernia, advanced degeneration, or spondylolisthesis? Can LMM atrophy explain low back pain in those patients? This problem may be elucidated by a new study investigating the prevalence of LMM atrophy in the general population.

Our study has a number of limitations. We did not include the duration of pain in our analysis. Thus, we could not analyze acute and chronic pain subgroups separately. In addition, there was no age and sex-

Table 2. LMM atrophy at L4-L5 and L5-S1 levels in male and female subjects

L4-L5 level		<40 years (n=937)	≥ 40 years (n=1091)	P
Females	Grade 0	457 (83.2)	339 (47.4)	<0.001
	Grade 1+2	92 (16.7)	375 (52.5)	
Males	Grade 0	360 (9.2)	226 (59.9)	<0.001
	Grade 1+2	28 (7.2)	151 (40)	
L5-S1 level				
Females	Grade 0	225 (41)	105 (14.7)	<0.001
	Grade 1+2	324 (59)	609 (85.3)	
Males	Grade 0	255 (65.7)	112 (29.7)	<0.001
	Grade 1+2	133 (34.3)	265 (70.3)	

Data are presented as n (%).
LMM, lumbar multifidus muscle; Grade 0, no atrophy; Grade 1, mild atrophy; Grade 2, advanced atrophy.

Table 3. Age comparison between patients without hernia according to grade 0–2 LMM atrophy at L4-L5 and L5-S1 levels

	L4-L5			L5-S1		
	n	Age (years)	P*	n	Age (years)	P*
Grade 0	696	34.99 ± 9.97	<0.001	392	34.6 ± 10.28	<0.001
Grade 1	132	45.95 ± 11.56		497	42.90 ± 12.17	
Grade 2	3	68.67 ± 12.50		62	52.21 ± 12.94	

Data are presented as mean \pm standard deviation.
LMM, lumbar multifidus muscle; Grade 0, no atrophy; Grade 1, mild atrophy; Grade 2, advanced atrophy.
* One-way ANOVA (Posthoc Tukey HSD test).

matched control group in our study, due to the difficulty of establishing an adequate control group in such a large sample.

In conclusion, LMM atrophy is more frequent in women than in men, and its prevalence and severity increase with age. For-

ty years of age appears to be a significant threshold for development of LMM atrophy. Hernia is found more frequently in individuals with LMM atrophy. We suggest that LMM atrophy may be the only MRI finding in low back pain patients, since 13% of our patients had LMM atrophy without disc hernia. However, further studies on the general population are needed to support this hypothesis.

Conflict of interest disclosure

The authors declared no conflicts of interest.

References

1. Rubin DI. Epidemiology and risk factors for spine pain. *Neurolog Clin* 2007; 25:353–371. [CrossRef]
2. Freeman MD, Woodham MA, Woodham AW. The role of the lumbar multifidus in chronic low back pain: a review. *PM&R* 2010; 2:142–146. [CrossRef]
3. Kjaer P, Bendix T, Sorensen JS, Korsholm L, Leboeuf-Yde C. Are MRI-defined fat infiltrations in the multifidus muscles associated with low back pain? *BMC Med* 2007; 5:2. [CrossRef]
4. Dannels LA, Vanderstraeten GG, Cambier DC, Witvrouw EE, De Cuyper HJ. CT imaging of trunk muscles in chronic low pain patients and healthy control subjects. *Eur Spine J* 2000; 9:266–272. [CrossRef]
5. Woodham M, Woodham A, Skeate JG, Freeman M. Long-term lumbar multifidus muscle atrophy changes documented with magnetic resonance imaging: a case series. *J Radiol Case Rep* 2014; 8:27–34.
6. Beneck GJ, Kulig K. Multifidus atrophy is localized and bilateral in active persons with chronic unilateral low back pain. *Arch Phys Med Rehabil* 2012; 300–306. [CrossRef]
7. Parkkola R, Rytokoski U, Kormanen M. Magnetic resonance imaging of the discs and trunk muscles in patients with chronic low back pain and healthy control subjects. *Spine* 1993; 18:830–836. [CrossRef]
8. Kader DF, Wardlaw D, Smith FW. Correlation between the MRI changes in the lumbar multifidus muscles and leg pain. *Clin Radiol* 2000; 55:145–149. [CrossRef]
9. Karataş M. Lomber omurganın fiziksel özellikleri ve fonksiyonel biyomekaniği. In: Beyazova M, Gökçe Kutsal Y. eds. *Fiziksel tıp ve rehabilitasyon*. Ankara: Güneş Kitabevi, 2000; 459–480.
10. Özcan Yıldız E. Bel ağrısı. In: Beyazova M, Gökçe Kutsal Y. eds. *Fiziksel tıp ve rehabilitasyon*. Ankara: Güneş Kitabevi, 2000; 1465–1483.
11. Moseley GL, Hodges PW, Gandevia SC. Deep and superficial fibers of the lumbar multifidus muscle are differentially active during voluntary arm movements. *Spine* 2002; 27:29–36. [CrossRef]
12. Wu WW, Hu ZJ, Fan SW, et al. Influencing of chronic low back pain on multifidus muscle atrophy. *Zhongguo Gu Shang* 2014; 27:207–212.
13. Battié MC, Niemeläinen R, Gibbons LE, Dhillon S. Is level and side specific multifidus asymmetry a marker for lumbar disc pathology? *Spine J* 2012; 12:932–939. [CrossRef]
14. Grimby G, Saltin B. The ageing muscle. *Clin Physiol* 1983; 3:209–218. [CrossRef]

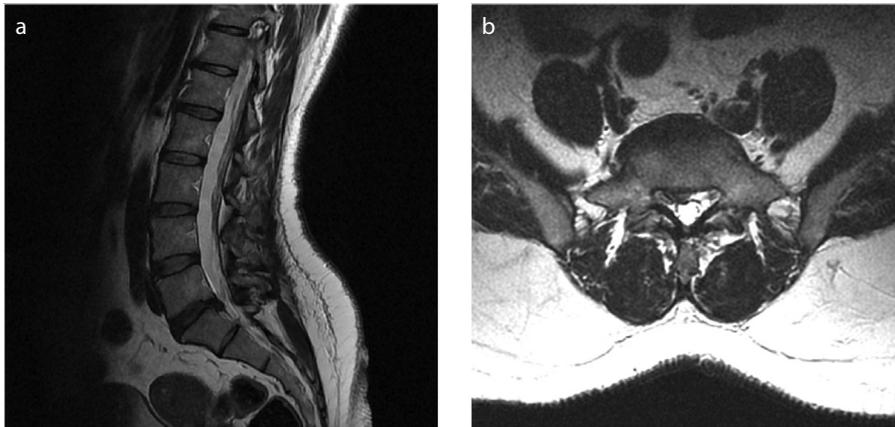


Figure 2. a, b. A 30-year-old female patient with disc hernia without LMM atrophy. Midsagittal T2-weighted image (a) shows sequestered disc hernia at L5-S1 level; dehydration is seen in the disc. Axial T2-weighted image at L5-S1 level (b), shows right lateral compression of the sequestered disc into the dural sac. Multifidus muscle was considered normal (grade 0).

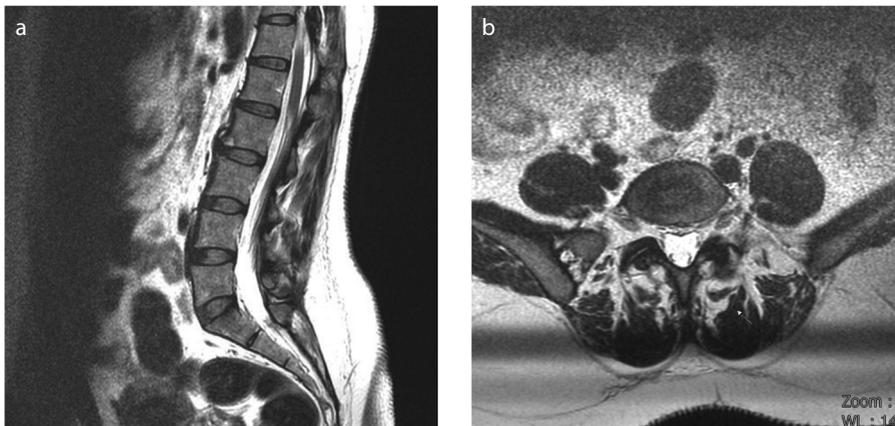


Figure 3. a, b. A 40-year-old female patient with grade 1 LMM atrophy without disc hernia. Midsagittal image (a) shows normal L4-L5 and L5-S1 discs; no significant loss of height or disc hernia is seen. Axial T2-weighted image (b) shows grade 1 LMM atrophy at L5-S1 level.

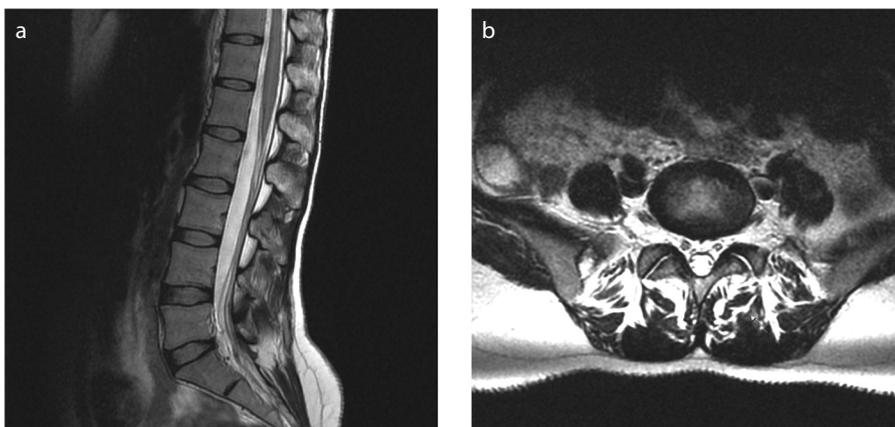


Figure 4. a, b. A 36-year-old male patient with grade 1 LMM atrophy without disc hernia. Midsagittal image (a) shows normal L4-L5 and L5-S1 discs; no significant loss of height or disc hernia is seen. Axial T2-weighted image (b) shows grade 1 LMM atrophy at L5-S1 level.