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

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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 (MRI) 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.
**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).

**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).

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.
Lumbar multifidus muscle atrophy and disc hernia

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). Consequently, 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 spondylolysis? 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-

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

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

Data are presented as n (%).

LMM, lumbar multifidus muscle; Grade 0, no atrophy; Grade 1, mild atrophy; Grade 2, advanced atrophy.

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

<table>
<thead>
<tr>
<th></th>
<th>L4-L5</th>
<th>L5-S1</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>Age (years)</td>
</tr>
<tr>
<td>Grade 0</td>
<td>696</td>
<td>34.99±9.97</td>
</tr>
<tr>
<td>Grade 1</td>
<td>132</td>
<td>45.95±11.56</td>
</tr>
<tr>
<td>Grade 2</td>
<td>3</td>
<td>68.67±12.50</td>
</tr>
</tbody>
</table>

Data are presented as mean±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
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]