The Role of Ultrasound in Diagnosis of the Causes of Low Back Pain: a Review of the Literature

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Context: Low back pain (LBP) is among the most prevalent musculoskeletal conditions in the developed countries. It is a common problem causing disability and imposing a huge economic burden to individuals and state organizations. Imaging plays an important role in diagnosis of the etiology of LBP.

Evidence Acquisition: The electronic databases included: PubMed (1950 to present), Ovid SP Medline (1950 to present) and ISI (1982 to present) and Google Scholar. In every search engine another search was performed using various permutations of the following keywords: ultrasonography, ultrasound imaging, low back pain, back muscles, paraspinal muscles, multifidus, transverse abdominis, muscle size, spinal canal, sacroiliac joint and spondylolisthesis.

Results: Magnetic resonance imaging (MRI) is widely used in evaluation of patients with LBP; however, high costs, limited availability and contraindications for its use have restricted MRI utilization. In a quest for a less expensive and readily available tool to investigate LBP, clinicians and researchers found ultrasonography (US) as an alternative. In this review we discuss the US application in diagnosis of some common causes of non-specific chronic LBP. Discussed topics include evaluation of spinal canal diameter, paraspinal and transabdominal muscles, sacroiliac joint laxity, pregnancy related LBP, sacroiliitis, and spondylolisthesis using US in patients with LBP.

Conclusions: While the first researches on employing ultrasound in diagnosis of patients with LBP had been focused on spinal canal diameter, recent studies have been mostly performed to evaluate the role of transabdominal and paraspinal muscles on core stability and thereby LBP occurrence. On the other side, Doppler ultrasonography has recently played an important role in objective measurement of joint laxity as a common etiology for LBP. Doppler imaging also in pregnant patients with LBP has been recommended as a safe and sensitive method. As conclusion, according to recent and most prestigious studies, focusing more on transabdominal muscle thickness can be considered as future approach in investigations.

Keywords: Ultrasonic Diagnosis; Low Back Pain; Review Literature
since its inception. The electronic databases included: PubMed (1950 to present), Ovid SP Medline (1950 to present) and ISI (1982 to present) and Google Scholar. In every search engine another search was performed using various permutations of the following keywords: ultrasonography, ultrasound imaging, low back pain, back muscles, paraspinal muscles, multifidus, transverse abdominis, muscle size, spinal canal, sacroiliac joint and spondylolisthesis. Secondary searching (or PEARLing) was however undertaken, whereby the reference lists of the selected articles were reviewed for additional references not identified in the primary search. The full text of all potentially relevant articles were retrieved and screened by the authors, in order to determine the eligibility of the paper for inclusion in the review.

3. Results

3.1. Ultrasonography and Spinal Canal Diameter

Employing US in measurement of spinal canal diameter was firstly reported by Porter et al. (24) who measured the diameter of spinal canal by placing the transducer of ultrasound in an oblique midsagittal plane 1 centimeter lateral to midline (24, 26-28). Porter et al.(24) reported that spinal diameter in trefoil shaped vertebrae such as L5 is less than other vertebrae and the incidence of LBP due to bony stenosis or laminar hypertrophy might be more in trefoil shaped vertebrae in comparison to other vertebrae. Porter et al.(26) also found that mean spinal canal diameter in patients with LBP was 1.44 cm, compared to a diameter of 1.61 in normal subjects and this difference was found to be statistically significant. It was suggested that a diameter of <1.4 cm increases the risk of LBP and subjects with a wider spinal canal can escape from the root entrapment on account of the greater space in their spinal canal. In the latter study, however, the patients of two groups were not well-matched demographically (e.g. age which is a determinant in the size of spinal canal was higher in LBP group). In a large scale study (27), a significant correlation was found between the incidence of neurologic claudication, disc lesion and nerve root entrapment and the spinal canal diameter of the subjects. Subsequent studies showed that the extreme of small spinal canal diameter is correlated with number of days one is absent from work; more than 32% of days lost due to the LBP were related to men whose spinal canal size was less than the 10th percentile of the measurements achieved from the subjects (29). The odds ratio of a spinal canal size in the lower 10th percentile spinal canal size and absence from work was reported to be 10.7 (30). Regarding the measurement of spinal canal diameter employing ultrasonography, reliability studies have reported the ease of learning the technique and rapid development of operator’s skill in performing the procedure (25, 29, 31). It was also demonstrated that the technique has a high inter and intra observer reliability with an average error of <0.2 mm between measurements (24, 32); this was, however, a non-uniform finding and a number of studies concluded that the technique is operator dependent (33), lacks acceptable reliability in elderly patients (34), and has a high interobserver (0.9 to 1.5 mm) and intraobserver error (0.6 mm to 0.9 mm) (35).

Transabdominal ultrasonography was also used in measurement of spinal canal size (36, 37); this technique was suggested to have an acceptable role in measurement of spinal canal size both in epidemiologic and pre-employment screening tests (36, 37). Missere et al. (36) reported a sensitivity of 84% and specificity of 60% for transabdominal ultrasound in diagnosis of LBP with a cut-off spinal diameter of 14 mm; nevertheless, the use of transabdominal ultrasonography is limited due to anatomical restrictions like different abdominal tissues and oblique direction of the intervertebral discs as the major boundaries against ultrasonographic pulses at the level of L3-4 and L5-S1 (37).

Finally, well-controlled, prospective studies demonstrated that although spinal canal size might be a risk factor for LBP, ultrasound measurement of spinal canal size has no practical role in prediction and/or estimation of the prognosis of LBP, neither in workers nor in general population (38, 39). It seems that small spinal canal diameter only plays as a facet of a multifactorial matrix in development of LBP. In addition, spinal canal size measurement using ultrasound has not gained wide attention since it requires quite a lot of expertise of the operator and the proficiency of the operator is the major determinant of the accuracy of measurements.

3.2. Ultrasonography and Paraspinal Muscles

Osseoligamentous lumbar spine is inherently unstable (40, 41) and is dependent on the integrated function of the muscles (especially paraspinal muscles) and neural subsystems for stability and movement (42, 43). Among the paraspinal muscles lumbar multifidus (LM) has a unique role in spinal stabilization and contributes to almost 2/3 of lumbar spine stability especially in the lower lumbar section (44) and is the predominantly affected paraspinal muscle in patients with LBP (45). In healthy subjects, the LM muscles are round or oval in shape, symmetrical between sides and increase in size cephalocaudally (46-49).

The most commonly used imaging studies for evaluation of paraspinal muscles are CT, MRI and rehabilitative ultrasound imaging (RUSI). Important aspects of muscles assessed using RUSI are muscle size, density and muscle contraction (50). The reliability of using RUSI to measure size of paraspinal musculature has been shown to be fair to excellent (ICC = 0.72-0.98), which is acceptable for clinical application (47, 51-53). It also has shown to have a reasonable inter-rater reliability among novice raters (54). Many authors reported muscle cross-sectional area (CSA) as the indicator of
Healthy (10)

LBP (41)

CSA of LM

.linear

Subject Position

Year

Results

Heidari P et al.

Healthy

LBP

The altered function of LM might be due to failure of a neuromuscular self-regulatory mechanism to regulate muscle contraction in order to meet the postural demands, which may in turn predispose the patient to LBP.

PMI for this purpose, however, remains to be specified in future studies; achievement of this goal might require sticking to strict criteria, as image brightness is affected by gain setting on the ultrasound machine.

Measurement of changes in muscle activation associated with LBP can lead to development of selective interventions to reverse the identified impairment. Pursuing this goal, USI was used to evaluate functional impairments in muscle contraction. It was demonstrated that there is a linear relation \((r = 0.79 \ P = 0.001)\) between LM thickness change and EMG activity across a narrow span of activation range \((19-34\%\) of maximum voluntary isometric contraction \((MVIC))\) \((52)\). USI was able to show that patients with LBP were not able to voluntarily contract the LM at the vertebral level with muscle atrophy \((65,66)\). It was demonstrated that the ability of patients with chronic LBP to activate the LM at the affected lumbar section is reduced, as evidenced by smaller increases in thickness on RUSI images during contraction compared to contralateral normal side muscle or asymptomatic control subjects \((58,67)\).

**Table 1. Details of Studies Using Ultrasound for Evaluation of Paraspinal Muscles**

<table>
<thead>
<tr>
<th>Author</th>
<th>Year</th>
<th>Subject (No.)</th>
<th>Measuring Elements (Muscle)</th>
<th>Probe Position</th>
<th>Subjects Position</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hides et al.</td>
<td>1992</td>
<td>Healthy (48)</td>
<td>Linear dimensions and CSA of LM</td>
<td>Transducer placed transversely over the spinous process and moved directly laterally.</td>
<td>Prone with the head in the midline position with a small roll placed under the forehead and two rolls under the shoulders. The lower lumbar spine was made flat by placing pillows under the hips, then gradually decreases during forward flexion while; in patients with chronic LBP forward flexion produces a further increase in CSA ((59)). The altered function of LM might be due to failure of a neuromuscular self-regulatory mechanism to regulate muscle contraction in order to meet the postural demands, which may in turn predispose the patient to LBP. USI showed that patients with LBP were not able to voluntarily contract the LM at the vertebral level with muscle atrophy ((65,66)). The validation and reliability of USI for this purpose, however, remains to be specified in future studies; achievement of this goal might require sticking to strict criteria, as image brightness is affected by gain setting on the ultrasound machine. Measurement of changes in muscle activation associated with LBP can lead to development of selective interventions to reverse the identified impairment. Pursuing this goal, USI was used to evaluate functional impairments in muscle contraction. It was demonstrated that there is a linear relation ((r = 0.79 \ P = 0.001)) between LM thickness change and EMG activity across a narrow span of activation range ((19-34%) of maximum voluntary isometric contraction ((MVIC))) ((52)). USI was able to show that patients with LBP were not able to voluntarily contract the LM at the vertebral level with muscle atrophy ((65,66)). It was demonstrated that the ability of patients with chronic LBP to activate the LM at the affected lumbar section is reduced, as evidenced by smaller increases in thickness on RUSI images during contraction compared to contralateral normal side muscle or asymptomatic control subjects ((58,67)).</td>
<td></td>
</tr>
</tbody>
</table>

| Kennedy et al. | 1993 | Scoliosis (20) | Linear dimensions and CSA of LM | Transducer placed transversely over the spinous process and was held against the skin surface at 90° and moved laterally. | Prone with a rolled towel under their forehead and shoulder. A pillow was placed under the hips to eliminate the lumbar lordosis. | It was shown that for different curve types in lumbar scoliosis, a pattern of asymmetry in LM exists. |

| Hides et al.   | 1994 | Healthy/LBP (51/26) | Linear dimensions and CSA of LM | Transducer was placed transversely over the spinous process and moved directly laterally. | Prone with the head in the midline position with a small roll placed under the forehead and two rolls under the shoulders. The lower lumbar spine was made flat by placing pillows under the hips. | Most of the patients showed greatest wasting at the level of 5. Asymmetry of CSA in patients was significantly different from between-side differences in control group. This asymmetry was greater in female patients. |

| Hides et al.   | 1995 | Healthy (10) | Linear dimensions and CSA of LM | Transducer was located adjacent to demarcated spinous process of the level to be examined. | Subjects were positioned in a comfortable and relaxed prone position, with their hips flexed to 35°. | In terms of LM muscle CSA no significant differences were found between MRI and US in young female adults. |

| Hides et al.   | 1996 | LBP (41) | CSA of LM | Transducer was placed transversely over the spinous process and moved directly laterally. | Prone with the head in the midline position with a small roll placed under the forehead and two rolls under the shoulders. The lower lumbar spine was made flat by placing pillows under the hips. | In the group that only received medical treatment LM muscle recovery was not spontaneous on remission of painful symptoms in patients. After 10-week follow-up examination patients in this group still had decreased LM muscle size. |

<table>
<thead>
<tr>
<th>Eisele et al. (69)</th>
<th>1998</th>
<th>Healthy/Lumbar Disk Disorder/Unknown (30/20/40)</th>
<th>Texture analysis and CSA of paraspinal lumbar muscle</th>
<th>Not Stated</th>
<th>Not Stated</th>
<th>Using US all patients with lumbar spinal history were detected. LM texture analysis can be a good and rapid investigation in patients with discogenic and structural disorders. Assessing CSA of LM muscle at the level of L5 can be made at both prone and side lying positions.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coldron et al. (70)</td>
<td>2003</td>
<td>Healthy (20)</td>
<td>CSA of LM</td>
<td>The transducer was placed longitudinally over the skin marking for spinous process and then moved directly laterally.</td>
<td>Not Stated</td>
<td>Not Stated</td>
</tr>
<tr>
<td>Stokes et al. (48)</td>
<td>2005</td>
<td>Healthy (120)</td>
<td>Linear dimensions and CSA of LM</td>
<td>The transducer was first placed longitudinally over the lower lumbar spine, in the mid-line. The transducer was then rotated through 90° to lie transversely in the midline and the spinous processes and laminae were identified on a cross-sectional scan. The transducer was then moved laterally to each side.</td>
<td>Prone: the subject lay with the head in the midline position, with one pillow under the lower legs and another under the hips to reduce the lumbar lordosis. 2: Side lying position: subjects lay on their left side and the transducer was placed behind the subject. One pillow was placed under the head and another between the knees, with the hips and knees positioned in sufficient flexion. A rolled towel was placed under the waist.</td>
<td>It was found that CSA of LM is larger in males and age didn’t have any effect. Both in males and females the CSA was larger in L5 than L4. Linear measurements multiplied (A × Lat) correlated highly with CSA.</td>
</tr>
<tr>
<td>Lee et al. (59)</td>
<td>2006</td>
<td>Healthy/LBP (99/16)</td>
<td>CSA of LM</td>
<td>The transducer was held perpendicular to the skin surface of the subjects’ lower back.</td>
<td>Prone: Subjects were prone and a small pillow was inserted below their abdomen. 2: Subjects were upright standing, and 25° and 45° forward stooping.</td>
<td>In different positions, CSA changes in LBP group had a reverse pattern in comparison to healthy subjects.</td>
</tr>
<tr>
<td>Pressler et al. (71)</td>
<td>2006</td>
<td>Healthy (30)</td>
<td>Linear dimensions and CSA of LM</td>
<td>The transducer was held orthogonal to the surface of the body and moved slowly from the left or right PSIS to the SI spinous process.</td>
<td>Prone with 35° of hip flexion and no lumbar lordosis. A manually adjusted treatment table with the hip joints placed along the hinge was used.</td>
<td>US seems to be a reliable way for imaging LM at the level of SI by newly trained assessors.</td>
</tr>
<tr>
<td>Vasseljen et al. (72)</td>
<td>2006</td>
<td>Healthy (10)</td>
<td>LM muscle activity onset</td>
<td>The probe was transversally oriented to the fiber direction and placed on a line running from the PSIS to the L1L2 interspinous space. Subjects stood relaxed with their arms beside the body.</td>
<td>US can detect muscle activity onset accurately but it has a small systematic delay that should be corrected for determining activity onset.</td>
<td></td>
</tr>
<tr>
<td>Kiesel et al. (52)</td>
<td>2007</td>
<td>Healthy (5)</td>
<td>Thickness of LM</td>
<td>Transducer was placed along the spine with the mid-point over the L4 spinous process. It was moved laterally and angled slightly medially until the L4/S zygapophyseal joint could be identified.</td>
<td>Prone position. An inclinometer was placed longitudinally over the lumbosacral junction and pillows were used to flatten the lumbar curve to less than 10°.</td>
<td>In a narrow range of LM muscle contraction RUS showed to be a valid method of measurement.</td>
</tr>
<tr>
<td>Wallwork et al. (51)</td>
<td>2007</td>
<td>Healthy (10)</td>
<td>Thickness of LM</td>
<td>The transducer was placed longitudinally where the zygapophyseal joints, the overlying multifidus muscle bulk at 2 to 3 vertebral levels, and the thoracolumbar fascia could be visualized.</td>
<td>Prone, with a pillow placed under the abdomen to minimize the lumbar lordosis.</td>
<td>Reliable evaluations of CSA of LM muscle in two vertebral levels were performed by both novice and experienced assessors.</td>
</tr>
<tr>
<td>Hides et al. (71)</td>
<td>2008</td>
<td>Athletes (26)</td>
<td>CSA of LM</td>
<td>The transducer placed transversely over the spinous process of the vertebral level being measured.</td>
<td>Prone with a pillow placed under the abdomen to minimize lumbar lordosis.</td>
<td>Even in highly active individuals with LBP atrophy of LM can exist. Improvement in CSA of LM was concomitant with a decrease in pain.</td>
</tr>
<tr>
<td>Hides et al. (74)</td>
<td>2008</td>
<td>Healthy/LBP (40/50)</td>
<td>CSA of LM</td>
<td>The transducer placed transversely over the spinous process of L2 to L5.</td>
<td>Prone with pillows under the hips to eliminate the lumbar lordosis.</td>
<td>Level of L5 was the greatest site of asymmetry in LM in patients with unilateral pain. The reported side of pain was the side that LM was smaller.</td>
</tr>
<tr>
<td>Wallwork et al. (58)</td>
<td>2009</td>
<td>Healthy/LBP (7/7)</td>
<td>Thickness and CSA of LM</td>
<td>The transducer placed transversely over the spinous process of L2 to L5.</td>
<td>Prone, with a pillow placed under the abdomen to minimize the lumbar lordosis.</td>
<td>At the level of L5, smaller CSA of LM muscle was reported for subjects of CLBP group than control group and percent thickness contraction was smaller in CLBP group.</td>
</tr>
<tr>
<td>Dicks et al. (75)</td>
<td>2010</td>
<td>Healthy (15)</td>
<td>Thickness of LM</td>
<td>The transducer was placed on the spinous processes and then moved lateral allowing visualization of the zygapophyseal joints, multifidus muscle and thoracolumbar fascia.</td>
<td>Prone with pillows under the abdomen to minimize the lumbar lordosis. An inclinometer ensured that the lumbar curve was less than 10°.</td>
<td>Muscle thickness increase during contraction decreased when unilateral pain was induced at a segmental level.</td>
</tr>
</tbody>
</table>
3.3. Ultrasonography and Sacroiliac Joint Dysfunction

SIJ has been implicated as the primary source of pain in 10% to 25% of the patients with LBP (77, 78). Abnormal biomechanics of the SIJ is considered a potential source of LBP (79-81); however, evaluation of SIJ dysfunction is still a challenge to clinicians. Intra-articular injection of steroids and analgesics to the SIJ is the gold standard in the diagnosis of SIJ dysfunction (82, 83). Nevertheless, the technique is invasive and requires specialized equipment and expertise. Neither the SIJ clinical presentation nor diagnostic procedures like X-rays and MRI are accurate enough for evaluation of SIJ dysfunction (82, 84, 85). Thus, a non-invasive objective method, which can be routinely applied in the clinic, was needed. Buyruk et al. (86) developed a technique called Doppler imaging of vibrations (DIV), to objectively measure the laxity of the SIJ. The laxity of the SIJ is quantified as threshold units (TU) which is assumed to be representative of the degree of vibration intensity attenuation through the SIJ, measured by color Doppler ultrasound (CDUS). TU is directly related to the density of the tissue that the vibrations pass through; for instance, a large difference in TU between sacrum and ilium indicates a lax joint and a small difference or an absence of it is an indicator of a stiff joint (87-92). The laxity measurements of the SI joint with DIV, after specific training of the operator, seems to be reliable and accurate (92). Moreover, DIV has proven to be of clinically relevant in evaluation of the patients with pregnancy related pelvic pain (88-91).

3.3.1. Pregnancy

According to different studies, twenty to eighty percent of women complain of some sort of back pain during pregnancy (93-99). This pain may persist, or arise, after delivery (100), and will, in some patients, lead to severe disability (97, 101-105). Two major patterns of back pain during pregnancy have been identified; pregnancy-related lumbar pain (PRLP) and pregnancy-related posterior pelvic pain (PRPPP) (106).

These mechanisms of back pain during pregnancy have been a matter of debate (107-112) and the precise mechanism of pregnancy back pain development is not fully understood (113). Studies have suggested that both PRLP and PRPPP might be related to sacroiliac joint (SIJ) dysfunction (89, 114). Recent introduction of DIV has made the evaluation of SIJ laxity feasible (115). DIV was first used in a cross-sectional study to compare SIJ stiffness in patients with peripartum pelvic pain including both PRPPP and PRLP patients and healthy subjects. No significant difference in the stiffness of SIJ was found between pelvic pain group and healthy subjects. Nevertheless, asymmetry in stiffness of the SIJ was significantly higher in patients with pelvic pain (88). These findings were further confirmed in succeeding studies (113, 116) where the authors showed the asymmetry in laxity rather than stiffness in SIJ is related to moderate to severe PRPPP (116). It was also demonstrated that asymmetric laxity during pregnancy measured using DIV, can predict persistence of pelvic pain to the postpartum period in patients with moderate to severe PRPPP; patients with asymmetrical laxity of SIJ during pregnancy had a three-fold increase in moderate to severe pain persisting into the postpartum period compared with subjects with symmetrical laxity. The sensitivity, specificity, and positive predictive value of SIJ asymmetric laxity during pregnancy for PRPPP persisting postpartum were 65%, 83%, and 77%, respectively (113).

3.4. US in Sacroiliac Joint Inflammation

Sacroiliitis is a frequent and early manifestation of the spondyloarthropathies (SpA) (117). Inflammatory back pain (IBP) due to sacroiliitis is the key symptom of axial involvement in SpA and is present in the majority of patients with SpA (118-120). Diagnosis of sacroiliitis is mainly based on clinical findings and X-ray studies, which lack specificity and are poorly reproducible (121). MRI can demonstrate early pre-destructive alterations of SIJ, and thus provide an early diagnosis of sacroiliitis (122-126). The availability of MRI, however, is limited, and the technique is time consuming and costly (123, 125, 127).

Color Doppler ultrasonography (CDUS), a technology that is widely used for detection of blood flow, has been used for diagnosis of sacroiliitis. Early studies found a high sensitivity (100%) for CDUS to detect vascularity around and/or inside SIJ in patients with active sacroiliitis (128); but this finding lacks specificity as vascularity was also present in some of the patients with osteoarthritis and in the control subjects. However, in sacroiliitis the resistive index (RI) of the vasculature was significantly lower than those in osteoarthritis (P < 0.001) and volunteers (P < 0.001) (128). Nevertheless, not all the studies using CDUS showed such a high sensitivity for detection of sacroiliitis. Klauser et al. (129) could only detect sacroiliitis in 18% of the patients who had MRI confirmed inflammation of SIJ.
which is likely to be explained by the flow signal criteria that was used to detect SIJ involvement. Microbubble contrast agents have been shown to improve the detection of increased vascularity in inflamed synovium by CDUS (130-132). In sacroiliitis contrast enhanced CDUS had dramatically improved sensitivity (18% before contrast vs. 94% after contrast administration) and negative predictive value (72% before contrast vs. 97% after contrast administration) to a level that is comparable with MRI (129). Clustered receiver operating curve (ROC) analysis also demonstrated that enhanced CDUS was significantly better than unenhanced CDUS for the diagnosis of active sacroiliitis (Az = 0.61, P < 0.0001) (129). The high sensitivity and negative predictive value of contrast-enhanced CDUS may obviate the need for MRI in screening of the IBP patients for SIJ involvement (129). An advantage of CDUS is that it can be used to quantitate response to therapy; the increase of the RI value and the decrease of symptoms are indicative of response to medical treatment (34, 129). This method, however, has its own limitations; false negative results may arise when inflammation is at the anterosuperior portion only or when there is prominent spur formation dorsally, because of limited ultrasound beam penetration (129).

Most recently, ultrasound had been implemented for detecting sacroiliac joint effusion in spondyloarthropathies (SpA). US showed joint effusion in 38.9% of SIJs of patients with SpA and only in 1.7% SIJ of the controls (P < 0.0001). SIJ effusion assessed by US had a positive likelihood ratio of 2.67 for the presence of IBP (133). This implies that high resolution US might be useful in the assessment of SIJ involvement in SpA especially when IBP is present. Ultrasonographic detection of SIJ effusion is easy to perform, requires only standard and not sophisticated US equipment, and is an inexpensive tool suggesting that this diagnostic method could represent a relevant tool in clinical practice. Nevertheless, the value of US as a routine diagnostic tool for IBP due to sacroiliitis has to be further confirmed by future studies.

3.5. Ultrasonography in Spondylolisthesis

Spondylolisthesis refers to slipping of a vertebra relative to an adjacent vertebra. Early detection of progressive disease by repeated radiologic evaluation is a key factor in the successful management of this entity (134, 135). However, the relatively high cumulative radiation dose of long-term follow-up is of concern especially in pediatric patients (134). Ultrasound has the advantage of being devoid of harmful effects of radiation and could be easily used by the physician for repetitive follow-ups of spondylolisthesis. It was shown that ultrasonic measurement of vertebral dislocation has a mean error of only 1.3 mm (range 0·3 mm) and a high correlation (r = 0.976, P < 0.001) with X-ray in the measurement of degree of slip (134). Therefore, it appears that the accuracy of ultrasonography should be sufficient for clinical use. However, at the moment there is only limited evidence to support the use of ultrasound in serial follow-up of spondylolisthesis. Future studies will elucidate the role of ultrasound in the management of spondylolisthesis.

3.6. Ultrasonography and Transabdominal Muscles

The role of transverse abdominal muscle (TrA) in LBP has been thoroughly investigated over the past decade. The first clues to biomechanical involvement of TrA in patients with LBP were reports that demonstrated TrA muscle contractions prior to limb movement in asymptomatic subjects (66, 136, 137), while such response was delayed in patients with LBP (138). Alteration in motor control of the abdominal muscles, particularly TrA, in patients with LBP was reported to cause the delay (139). It was therefore concluded that TrA may play an essential role in the stability of spinal column, and thereby in LBP (140). These findings were further confirmed by finding that TrA strengthening exercises reduce the pain intensity in patients with LBP (141, 142). Moreover, an asymmetric pattern for TrA muscle thickness (143) and an asymmetric increase in TrA thickness in patients with LBP (144) provided additional evidence of TrA derangements in development of LBP. Studies that investigated transabdominal muscles using US are presented in Table 2.

The reliability of US in measuring TrA thickness in LBP has been extensively tested (145); according to Hodges et al. (146) US has an acceptable reliability in the measurement of TrA thickness if the transducer is positioned in the right place using a belt. This method of transducer fixation has also shown to have reasonable intra-class correlation co-efficient for measurement of abdominal muscle thickness during active tasks (147). Furthermore, it was found that US is also a reliable way in measuring the controlled contraction of TrA thickness (51, 148-150). It was demonstrated that there is a linear relation between level of contraction (measured by EMG) and thickness of TrA in up to 30-40% of MVIC of TrA (151, 152).

In a nutshell, it appears that the TrA dysfunction plays a major role in development and increase of severity of LBP. Ultrasound proved to be a reliable tool in the measurement of TrA thickness and contraction. However, well-controlled and better designed prospective studies are required to understand the role of ultrasound measurement of TrA thickness and activity in prediction of LBP in predisposed patients.

4. Conclusions

Ultrasonography has been used widely in diagnosis and even rehabilitation of patients with LBP; however, lack of conclusive evidence to generalize the use of ultrasonography in clinical setting cannot be dismissed. While the first researches on employing ultrasound in diagnosis of patients with LBP had been focused on spinal canal diameter, recent studies have been mostly performed to evaluate the role of transabdominal and paraspinal muscles on core stability and thereby LBP occurrence.
#### Table 2. Details of Studies Using Ultrasound for Evaluation of Abdominal Wall Muscles

<table>
<thead>
<tr>
<th>Author</th>
<th>Year</th>
<th>Subject (No.)</th>
<th>Measuring Elements (Muscle)</th>
<th>Probe Position</th>
<th>Subjects Position</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bunc et al. (145)</td>
<td>2002</td>
<td>Healthy (22)</td>
<td>TrA</td>
<td>The transducer was located between the 12th rib and the iliac crest over the anterolateral abdominal wall.</td>
<td>1: Supine; 2: Standing; 3: Walking on treadmill</td>
<td>US was found as a reliable way for measurement of TrA thickness.</td>
</tr>
<tr>
<td>Critchley (142)</td>
<td>2002</td>
<td>Healthy (28)</td>
<td>EO, IO, TrA</td>
<td>Transducer was located at a point 2.5 cm anterior to the midpoint between the ribs and ilium on the midaxillary line.</td>
<td>Low abdominal hollowing in four-point kneeling with and without pelvic floor contraction.</td>
<td>Co-contraction of pelvic floor with abdominal hollowing in maneuver may lead to greater increase of TrA thickness compared to abdominal hollowing in maneuver alone.</td>
</tr>
<tr>
<td>Kidd et al. (148)</td>
<td>2002</td>
<td>Healthy (11)</td>
<td>TrA</td>
<td>Not Stated</td>
<td>Lying and standing; no more details are described.</td>
<td>US imaging provides a reliable measure of controlled contraction of TrA.</td>
</tr>
<tr>
<td>Hodges (155)</td>
<td>2003</td>
<td>LBP (3)</td>
<td>EO, IO, TrA</td>
<td>The transducer was placed transversely across the abdominal wall along a line midway between the inferior angle of the rib cage and the iliac crest.</td>
<td>The subjects had to sit in a reclining chair when their hips were flexed to approximately 30°.</td>
<td>In terms of muscle activity US seems to detect low levels of muscle activity. Moderated and high muscle activity cannot be distinguished employing US.</td>
</tr>
<tr>
<td>Ferreira et al. (144)</td>
<td>2004</td>
<td>Healthy/LBP (10/10)</td>
<td>EO, IO, TrA</td>
<td>The transducer was placed in a transverse plane just superior to the left iliac crest along the axillary line.</td>
<td>Supine with arms crossed over the chest, the hips flexed to 50°, and knees flexed to 90°.</td>
<td>A positive correlation between EMG and US findings in those with and without LBP was found. Also changes in TrA control in patients with LBP comparing to other group was concluded.</td>
</tr>
<tr>
<td>McMeeken et al. (152)</td>
<td>2004</td>
<td>Healthy (11)</td>
<td>TrA</td>
<td>25 mm antero-medial to the midpoint between the ribs and ilium on the midaxillary line and parallel to transversus abdominis.</td>
<td>Supine with a pillow under the head and the knees bent to approximately 20° over two pillows.</td>
<td>Reliability of US measurements as well as a positive correlation between US and EMG findings were reported in this study.</td>
</tr>
<tr>
<td>Teyhen et al. (149)</td>
<td>2005</td>
<td>LBP (30)</td>
<td>EO, IO, TrA</td>
<td>The transducer was placed in a transverse plane just superior to the left iliac crest along the axillary line.</td>
<td>1: Quadeduped; 2: Seated; 3: Supine; 4: Hook-lying.</td>
<td>A high inter-reliability for transabdominal muscle measurement of those with and without LBP was achieved. Short-term abdominal drawing in maneuver did not influence the thickness of TrA.</td>
</tr>
<tr>
<td>Ainscough et al. (153)</td>
<td>2006</td>
<td>Healthy (30)</td>
<td>IO, TrA</td>
<td>The transducer was placed on the skin halfway between the anterior or superior or iliac spine and the lower rib cage in the anterior axillary line.</td>
<td>1: Supine lying; 2: Relaxed sitting on a chair; 3: Relaxed sitting on a gym ball with both feet on the ground; 4: Sitting on a gym ball lifting the left foot.</td>
<td>There was no difference in muscle thickness between relaxed sitting on chair and sitting on a gym ball. At the end of aspiration the muscles were thicker.</td>
</tr>
<tr>
<td>Hides et al. (150)</td>
<td>2006</td>
<td>Healthy (11)</td>
<td>IO, TrA</td>
<td>A transverse image of the anterolateral abdominal wall was obtained just inferior to the level of the umbilicus for left and right sides.</td>
<td>Supine</td>
<td>There was a positive correlation between MRI and US findings in measurement of IO and TrA. Anterior abdominal fascia of TrA moved laterally during weight bearing.</td>
</tr>
<tr>
<td>Rankin et al. (154)</td>
<td>2006</td>
<td>Healthy (123)</td>
<td>EO, IO, TrA, RA</td>
<td>1: Immediately below the ribcage in direct vertical alignment with the ASIS; 2: Halfway along a line joining the ASIS to just below the ribcage in the midaxillary line.</td>
<td>Subjects lay supine with two pillows under their knees.</td>
<td>In terms of relative thickness of the muscles the pattern was as follows: RA &gt; IO &gt; EO &gt; TrA. There was no asymmetry for all muscles relative thickness.</td>
</tr>
<tr>
<td>Springer et al. (143)</td>
<td>2006</td>
<td>Healthy (32)</td>
<td>EO, IO, TrA</td>
<td>The center of the transducer was placed in a transverse plane just superior to the iliac crest, in line with the midaxillary line.</td>
<td>Bilaterally while the subjects were at rest, and while they performed the abdominal drawing-in maneuver.</td>
<td>Bilateral symmetry in the lateral abdominal muscles in those without LBP.</td>
</tr>
<tr>
<td>Hides et al. (155)</td>
<td>2007</td>
<td>Healthy (19)</td>
<td>IO, TrA</td>
<td>The transducer was along a line midway between the inferior angle of the rib cage and the iliac crest.</td>
<td>Supine with their right heel against a footplate linked to a force transducer. Each subject performed a static simulated weight-bearing task of the right lower extremity.</td>
<td>A greater TrA than IO thickness was found. There was no significant differences between right and left abdominal muscles.</td>
</tr>
<tr>
<td>Hides et al. (156)</td>
<td>2007</td>
<td>Healthy (19)</td>
<td>IO, TrA</td>
<td>The transducer was along a line midway between the inferior angle of the rib cage and the iliac crest.</td>
<td>Supine hook-lying position, with their hips in 45° of flexion.</td>
<td>RUSI showed a high reliability in three measurements and also a fair to high reliability was stated across two days.</td>
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<td>Authors</td>
<td>Year</td>
<td>Group(s)</td>
<td>Procedure Description</td>
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<tr>
<td>Kiesel et al. (157)</td>
<td>2007</td>
<td>Healthy/LBP, TrA</td>
<td>Transducer placed along the lateral abdominal wall, just superior to the iliac crest, along the midaxillary line.</td>
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<td>Supine hook-lying position</td>
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<td>TrA muscle thickness significantly changed during the abdominal drawing-in maneuver.</td>
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<tr>
<td>Raney et al. (158)</td>
<td>2007</td>
<td>LBP (9), IO, TrA</td>
<td>Transducer was superior to the iliac crest, along the right midaxillary line in the transverse plane.</td>
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<td>Cases were positioned in the supine hook-lying position.</td>
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<td>In 6 of 9 patients increased ability to improve TrA thickness during drawing-in maneuver was demonstrated. The thickness of TrA at rest decreased in 5 patients. This decrease was also noted in IO muscle in 4 of the patients.</td>
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<tr>
<td>Kiesel et al. (159)</td>
<td>2008</td>
<td>Healthy (6), TrA</td>
<td>Transducer was placed just superior to the iliac crest along the axillary line.</td>
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<td>Supine hook-lying position</td>
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<td>TrA thickness changed during the abdominal draw-in activity. In terms of thickness changes, control group with pain was significantly different to no pain group.</td>
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<tr>
<td>Mannion et al. (160)</td>
<td>2008</td>
<td>Healthy/LBP, EO, IO, TrA</td>
<td>Transducer was positioned 2.5 cm anteromedial to the mid point between the iliac crest and the costal margin on the mid-axillary line.</td>
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<td>Supine hook-lying position (hips in 30° flexion)</td>
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<td></td>
<td>Using US there was no significant between-day differences in thickness of any muscle during rest and hollowning.</td>
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<tr>
<td>Hides et al. (159)</td>
<td>2009</td>
<td>Healthy/LBP, IO, TrA</td>
<td>Midway between the inferior angle of the rib cage and the iliac crest of both sides.</td>
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<td>Supine, lying on a near-frictionless surface with the heel of the test limb resting on a foot plate.</td>
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<td>Impairment of TrA and IO contraction in those without LBP in comparison to those with LBP.</td>
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<tr>
<td>Koppenhaver et al. (161)</td>
<td>2009</td>
<td>LBP (30), TrA</td>
<td>Transducer positioned just superior to the iliac crest along the midaxillary line.</td>
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<td>Supine, with hips and knees extended at rest and were instructed to “raise your leg off of the table approximately 8 inches (20 cm) without bending your knee“.</td>
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<td>RUSI showed to be a reliable method to measure TrA thickness based on the mean of two measures. A high reliability was demonstrated when the measures were taken by a single examiner, and the reliability employing different examiners was also adequate.</td>
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<td>Reeve and Dilley (140)</td>
<td>2009</td>
<td>Healthy (20), TrA</td>
<td>Between the iliac crest and the lowest rib along the anterolateral line.</td>
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<td>Posture may influence the measured thickness of TrA using US.</td>
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<tr>
<td>Kordi et al. (162)</td>
<td>2011</td>
<td>Healthy (63), EO, IO, TrA</td>
<td>A point 25 mm anteromedial at the mid point between the inferior rib and the iliac crest.</td>
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<td>2: A point immediately under the rib cage in direct vertical alignment with the ASIS.</td>
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<td>Subjects were positioned in a crook-lying position with pillows under the head and the knees.</td>
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<td>After food consumption thickness values significantly reduced in all measured abdominal muscles.</td>
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<tr>
<td>Noormohammadpour et al. (163)</td>
<td>2012</td>
<td>Healthy (19), EO, IO, TrA</td>
<td>Transducer was at A point 25 mm anteromedial to the midpoint between the inferior rib and the iliac crest on the mid-axillary line.</td>
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<td>Crook lying position while pillows were placed under their head and knees.</td>
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<td>After 12 weeks of concurrent energy restricted diet and abdominal resistance training increase in muscle thickness during drawing-in maneuver was demonstrated.</td>
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<tr>
<td>Rostami et al. (164)</td>
<td>2013</td>
<td>Healthy (50), EO, IO, TrA</td>
<td>The point of probe position was set at 25 mm anteromedial to the midpoint between the inferior rib and the iliac crest on the mid-axillary line, as it was previously used in other studies.</td>
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<td>Supine hook-lying position (supine position with hips flexes to almost 30°) where small pillows were laid under their knees and head.</td>
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<td>A significant positive relation was found between EO thickness and weight, mass index, waist circumference and skin fold thickness. IO muscle thickness decreased with higher values of mass index, waist circumference and skin fold thickness but weight did not have a significant correlation with IO thickness. These measurements of fatness showed no significant relation to TrA thickness.</td>
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</table>

Use of ultrasonography in the measurement of transabdominal muscles particularly TrA has led to findings which can be employed in rehabilitation of patients as a curative procedure. The rehabilitation would be established for the patients via biofeedback. It seems more controlled imaging studies should be carried out to recommend strengthening of transabdominal muscles in LBP treatment.

RUSI assessment of LM as the predominantly affected muscle in LBP is of clinical importance in diagnosis of LBP. Impaired morphology of LM has been investigated both in acute and chronic LBP. Atrophy and infiltration of fatty tissue into the muscle are seen in LM of individuals suffering LBP. The validity and reliability of US for this purpose, however, remains to be specified in future studies. US can also be used to evaluate functional impairments in muscle contraction in addition to morphological abnormalities. The use of USI for observation of contraction of paraspinal muscles which is reduced during chronic LBP has been recently validated.

On the other side, Doppler ultrasonography has recently played an important role in objective measurement of joint laxity as a common etiology for LBP. It was proposed that large difference in TU, obtained by DIV, between sacrum and ilium indicates a joint laxity and a smaller space in obtained view or an absence of it is an
indicator of a stiff joint. However conclusions based on measurements with DIV should be made with great care until further studies further validate the technique. There are also studies implying usefulness of US in assessment of SIJ in SpA especially when IBP is present. CDUS with microbubble contrast agent has been recently shown to be beneficial in diagnosis of sacroiliitis in light of detection of increased vascularity in inflamed joints. Nevertheless, in case of using Doppler ultrasonography easiness and inexpensiveness of usage cannot be well mentioned.

Doppler imaging also in pregnant patients with LBP has been recommended as a safe and sensitive method. However, it should be noted that there is only limited evidence regarding the use of US in spondylolisthesis and this method has not yet been widely used in clinical practice. As conclusion, according to recent and most prestigious studies, focusing more on transabdominal muscle thickness can be considered as future approach in investigations.

References


66. Hodges PW, Richardson CA. Feedback contraction of transversus abdominis is not influenced by the direction of arm movement. Exp Brain Res. 1997;114(2):362–70.


70. Coldron I, Stokes M, Cook K. Lumbar multifidus muscle size does not differ whether ultrasound imaging is performed in prone or side lying. Man Ther. 2003;8(3):361–5.


