

Accuracy of the Diagnostic Tests of Sacroiliac Joint Dysfunction



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ABSTRACT

Objective: The purpose of this study was to assess the reliability and validity of motion palpation and pain provocation compared with sacroiliac joint (SIJ) block as the gold-standard assessment method of patients with sacroiliac joint dysfunction (SIJD).

Methods: A cross-sectional study was conducted in the Department of Sports and Exercise Medicine at Rasool Akram Hospital. Forty-eight patients suspected of having SIJD were selected from a total of 150 patients on the basis of a combination of symptoms, physical tests, and magnetic resonance imaging findings. The patients suspected of having SIJD received the SIJ block, to which the accuracy of all the physical tests was compared. Sensitivity, specificity, and positive and negative predictive values were calculated for each test. The receiver operating characteristic curve and the area under the receiver operating characteristic curve were measured.

Results: The Flexion, Abduction and External Rotation (FABER) test had the highest specificity and positive predictive values of the physical tests. Furthermore, the combination of the FABER test and the thigh thrust test improved overall diagnostic ability more so than any of the other test combinations.

Conclusion: A combination of the motion and provocation tests increased specificity and positive predictive values, and the FABER test had the highest of these single values. The palpation tests did not change after the SIJ block, suggesting that their accuracy cannot be determined using this method. (J Chiropr Med 2020;19:28-37)

Key Indexing Terms: *Diagnostic Tests; Sensitivity and Specificity; Predictive Value of Tests*

INTRODUCTION

Lower back pain (LBP) is one of the most common musculoskeletal complaints; almost 80% of people experience pain in this area at least once in their lives.¹ Despite its high prevalence, the etiology of LBP is not well known and is nonspecific in approximately 85% of cases.²

The disorders affecting the sacroiliac joint (SIJ), which is an area extending inferiorly in the medial part of the posterior superior iliac spine (PSIS),^{3,4} are defined using different terms that include *sacroiliac strain*, *sacroiliac instability*, *sacroiliac arthritis*, and *sacroiliac joint*

dysfunction (SIJD). The latter is used in the case of a noninflammatory condition of the SIJ that is characterized by a reversible decreased mobility of the joint, resulting from articular causes.⁵

Sacroiliac joint dysfunction accounts for 10% to 27% of the causes of mechanical lower back or buttock pain,⁶⁻¹⁰ and one of its hallmarks is local tenderness in the SIJ.⁵ Sacroiliac joint dysfunction can be the sole disorder, or it can be accompanied by disc herniation or spinal stenosis.^{11,12}

It is difficult to make an exact diagnosis of SIJD, not only because historical, physical, or radiological evidence is not absolute, but also because the symptoms can be the result of other common conditions, such as facet syndrome and disc herniation.¹³⁻¹⁶ Because there is no widely accepted reference standard for diagnosing SIJD, the SIJ block is the preferred method for this purpose.¹⁷ With this method, an anesthetic agent is injected into the SIJ under fluoroscopic guidance. The rationale for using the SIJ block is that the SIJ has many nerves, which, when stimulated, can generate pain. Moreover, the level of evidence for the specificity and validity of this diagnostic test is considered moderate (level III).^{6,18-28} A number of authors recommend single-injection diagnostic block for clinical studies,^{29,30} although others suggest double (confirmatory) diagnostic block more accurately determines the source of pain by using 2 different local anesthetics with different durations of action.^{5,13,18,31-42}

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However, despite such recommendations, SIJ block is not cost-effective, nor is it practical for practitioners who lack training in intra-articular injections. These considerations have led clinicians to employ physical tests, with the belief that even negative findings can be used in diagnosing SIJD. The fact that there is a wide range of physical tests, many of which are not as accurate as the SIJ block,^{6,40,43-49} calls into question several factors, including which tests are most helpful, whether the lack of agreement on clinical criteria for an SIJD diagnosis makes utility of these tests controversial,¹⁴⁻¹⁶ and whether using a combination of such tests can lead to better diagnosis.

The present study aimed to determine which physical tests have the highest sensitivity, specificity, and predictive values in determining the presence of SIJD compared with the SIJ block, in addition to which combination of physical tests has the closest diagnostic value to the SIJ block.

METHODS

This cross-sectional study was carried out between 2016 and 2018 and used convenience-based sampling to recruit patients with lower back or buttock pain. A written consent compatible with the International Council for Harmonisation of Technical Requirements for Pharmaceuticals for Human Use was obtained from all participants. Ethical approval for this study was obtained from an independent ethics committee at Iran University of Medical Sciences.

Patients with lower back or buttock pain were evaluated by a sports medicine specialist through a 2-step screening process. First, physical musculoskeletal examinations were used to identify patients who were subject to an SIJD diagnosis. Patients were then evaluated for pathologies of the lumbar spine and pelvis; those with pain in these areas that emanated from other causes, such as lumbar discopathy, spinal stenosis, and spondylolisthesis, were excluded. Eligible patients were subsequently evaluated using lumbosacral and sacroiliac magnetic resonance imaging (MRI) scans to rule out other sources of pain and to further verify the results of the physical examinations.

Inclusion Criteria

The following inclusion criteria were adopted during the physical examinations and screening:

- Aged 20 to 70 years
- Presence of lower back or buttock pain with or without radiation to lower extremities for at least 6 weeks before study enrollment
- Presence of pain or local tenderness in the SIJ region (ie, the joint between the sacrum and the ilium bones of the pelvis, covering an area extending inferiorly in the medial part of the PSIS)

- Presence of pain exacerbated as a result of bending laterally or backward
- Positive results on at least 2 of the pain-provocation tests (ie, Flexion, Abduction, and External Rotation [FABER], thigh thrust, Gaenslen, Yeoman, compression, distraction, and Newton tests) and one of the motion palpation tests (ie, Gillet and forward flexion tests)

Exclusion Criteria

Patients were excluded if they met any of the following criteria:

- Pregnancy
- Receiving physical therapy modality and nonsteroidal anti-inflammatory drugs over a 72-hour period before the study period
- A history of back surgery during the 6 months before the study period
- Malignant tumors in the spine or pelvis
- Sacroiliitis and infections of the SIJ
- Presence of any fracture in the spine or pelvis
- Presence of other causes of LBP such as lumbar discopathy and spinal stenosis discovered via clinical examination and MRI scanning

Generally, SIJD-suspected participants were defined as patients with lower back or buttock pain whose symptoms indicated SIJD and who had positive results on at least 2 of the pain provocation tests (ie, FABER, thigh thrust, Gaenslen, Yeoman, compression, distraction, and Newton tests) and 1 of the motion palpation tests (ie, Gillet and forward flexion tests) in the absence of other causes of pain according to MRI test results.

Background Data

Collected patient characteristics included sex, age, and body mass index (BMI). To determine the intensity of the SIJ pain felt during each subjective test, a 100-mm visual analog scale (VAS) was used, where 0 represented no pain and 100 denoted the most severe pain. The VAS scores were recorded mainly to measure the decline in pain level as a result of administering the SIJ block.

Motion Palpation Tests

Gillet Test. To perform the Gillet test, the examiner stood behind the patient with one thumb on the PSIS and the other thumb on the sacrum. Then, the patient was instructed to bend and pull up the leg corresponding to the PSIS being palpated. The test was repeated on the other side and compared bilaterally.

The test was considered negative if the thumb on the PSIS moved inferiorly to the thumb placed on the sacrum. In contrast, no movement on the PSIS or superior movement to the other thumb on the sacrum was taken as a positive result.^{45,50-53}

Forward Flexion Test. During the flexion test, the patient was asked to slowly bend forward as much as possible with the examiner's thumbs on their left and right PSIS. Then, the symmetry of the movement in the thumbs was assessed. A positive test result was defined as any superiority in the movement of the 2 thumbs, indicating hypomobility of the ipsilateral SIJ.^{44,53}

Pain Provocation Tests (Subjective)

FABER Test (Patrick's Test). During the FABER test, the patient was asked to lie supine on the examination table. The examiner brought the hip joint into the FABER positions. One knee was flexed 90°, and the affected-side foot was rested on the opposite-side knee. Subsequently, the examiner pressed the contralateral anterior superior iliac spine (ASIS) against the table and pushed the bent knee down toward the table.^{3,6} The test was considered positive if the patient felt pain in the SIJ on the side where the knee was flexed. At this point, pain in the buttocks was suggestive of SIJD, whereas pain in the inguinal region could have indicated hip pathology.

Thigh Thrust Test (Posterior Shear Test). With the thigh thrust test, the patient lay in a supine position while the tested-side hip joint was flexed to approximately 90° by the examiner. An anteroposterior shear force was applied to the SIJ through the axis of the femur.^{3,43} Resulting pain indicated the test was positive.

Gaenslen Test. The patient lay supine, with the tested-side leg hanging over the edge of the table and the other leg flexed to the chest. The examiner applied firm pressure to the flexed knee, and a counterpressure was applied to the knee of the hanging leg. The procedure was then repeated on the opposite side.^{29,34} The test was considered positive if the patient felt pain in the hanging-leg side.

Yeoman Test. With this test, the patient was asked to lie prone. The examiner lifted the tested-side knee by extending it to 90° and then extended the hip joint with one elbow on the patient's buttock.^{3,47} Pain in the hanging-leg side was considered a positive result.

Compression Test. The patient lay in the lateral decubitus position, with the affected side up, and faced away from the examiner, who applied a downward pressure to the ipsilateral iliac crest and ASIS. The test was considered positive if the patient felt pain in the SIJ on the contralateral side.^{3,48}

Distraction Test. The patient was placed supine on the table. With the patient's forearms crossed, the examiner applied slow and steady outward pressure to the left and

right ASIS, spreading them away. The test was considered positive if the patient's pain in the SIJ increased.^{3,48,55}

Newton's Test. The patient lay in a supine position. The examiner fully flexed and pressed the tested-side hip and knee joints toward the abdomen. The test was considered positive if the patient experienced increased pain in the SIJ.³

SIJ Block (SIJ Injection)

Patients who met the criterion for positivity were suspected to have SIJD⁴⁷ and were transferred to the pain procedure room within an hour for the double SIJ block. This test was performed by a pain specialist with more than 10 years of experience in spinal injections and who was blind to the results of the physical tests. For the SIJ block, the patient lay in a prone position with a pillow placed under the abdomen at the iliac crests. After prepping and draping, a spinal 22G needle was inserted and positioned in the SIJ. Next, 1 mL of iodixanol (Visipaque) was injected as the radiocontrast agent. The placement of the needle and the spread of iodixanol was documented via fluoroscopy (a lateral view and a 3-quarter view). Additionally, 1.5 mL of lidocaine 2% was used in the initial injection, and 1.5 mL of bupivacaine 0.5% was employed in the confirmatory block,^{5,6,18,26,31-42} producing a double SIJ block within an hour of the SIJ block. The physical tests were repeated, and VAS pain scores were obtained again to measure possible pain relief compared with the pre-SIJ block state.²⁷ A pain reduction of at least 60% indicated the presence of SIJD, and a reduction of smaller than 60% denoted the absence of SIJD.^{28,39,56}

To measure the diagnostic validity of the physical tests, they were compared with the SIJ block in the SIJD-suspected patients. The comparison was performed for each individual test and different combinations.

The flowchart of the study is presented in [Figure 1](#).

Statistical Methods

The findings of the evaluations were analyzed using SPSS 23 (IBM Corporation, New York). The prevalence of SIJD in the patients with lower back or buttock pain was determined while considering their sex, age, and BMI. Also, 2 × 2 contingency tables were created. Moreover, sensitivity, specificity, and positive and negative predictive values were calculated for each individual test and different combinations of these tests based on the formulas presented in [Table 1](#).^{57,58}

The receiver operating characteristic curve is an overall measure of diagnostic efficacy. It is a plot of the true-positive rate against the false-positive rate of a diagnostic test. This curve, which is defined as a plot of test sensitivity as the y coordinate vs test specificity or false-positive rate as the x coordinate, is an effective method for evaluating the validity of diagnostic tests.

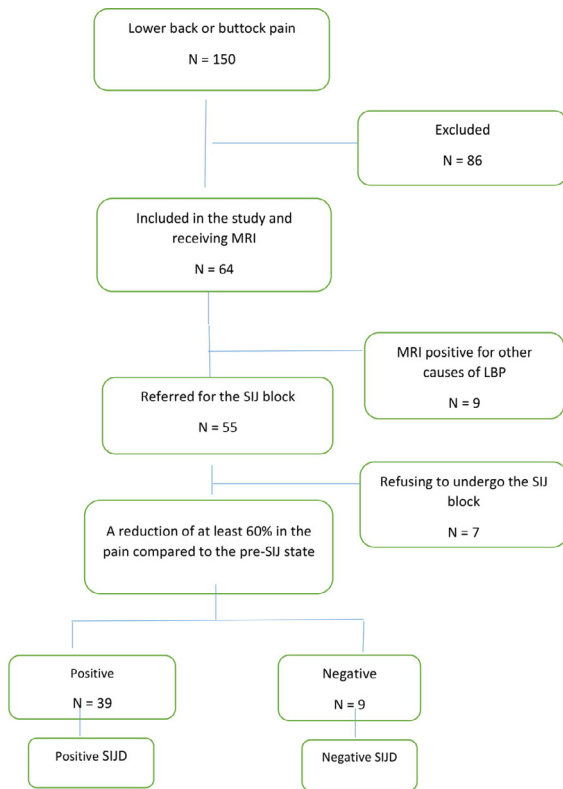


Fig 1. The flowchart of the study. LBP, lower back pain; MRI, magnetic resonance imaging; SIJ, sacroiliac joint.

The area under the curve (AUC) is the area under the receiver operating characteristic curve and a measure of how well a parameter can distinguish between 2 diagnostic groups (affected vs non-affected). The AUC can range from 0.5 (useless model) to 1.0 (perfect discrimination). A value higher than 0.7 can be interpreted as reasonable or fair; a value higher than 0.8 is considered acceptable.⁵⁷ The surface of the curve was calculated and measured using SPSS.

Table 1. Calculation of the Sensitivity, Specificity, and Positive and Negative Predictive Values of the Diagnostic Tests

	Positive SIJ Block	Negative SIJ Block
Positive physical test	A (true positive)	B (false positive)
Negative physical test	C (false negative)	D (true negative)

$$\text{Sensitivity} = (A / (A + C)) \times 100$$

$$\text{Specificity} = (D / (B + D)) \times 100$$

$$\text{Positive Predictive Value} = (A / (A + B)) \times 100$$

$$\text{Negative Predictive Value} = (D / (C + D)) \times 100$$

RESULTS

A total of 150 patients with lower back or buttock pain were examined at the beginning of the study. Sixty-four of these patients were selected to undergo an MRI. Of them, 9 patients were excluded as their LBP was the result of other causes. Further, 7 patients refused to undergo the SIJ block. Of the 48 remaining patients, 6 were male and 42 were female, ranging in age from 23 to 69 years old (average of 47.7 years) and a mean BMI of 28.4.

Of the 48 patients undergoing the SIJ block, 39 experienced a reduction of at least 60% in pain and were placed in the SIJD-positive group, for a prevalence of 81.25%. Although the pre-SIJ block VAS scores ranged from 60 to 100, with an average of 77, the post-SIJ block scores varied between 10 and 30, with an average of 18 (Fig 2). No adverse effects of the SIJ block were observed.

Nine patients had a reduction in pain that was less than 60% after the SIJ block and were assigned to the SIJD-negative group. Of these patients, the pre-SIJ block VAS scores ranged from 50 to 90, with an average of 74, whereas the post-SIJ block scores ranged from 40 to 70, with the average being 46 (Fig 3).

Tables 2 and 3 show the sensitivity, specificity, and positive and negative predictive values for each individual physical test and different combinations of these tests,

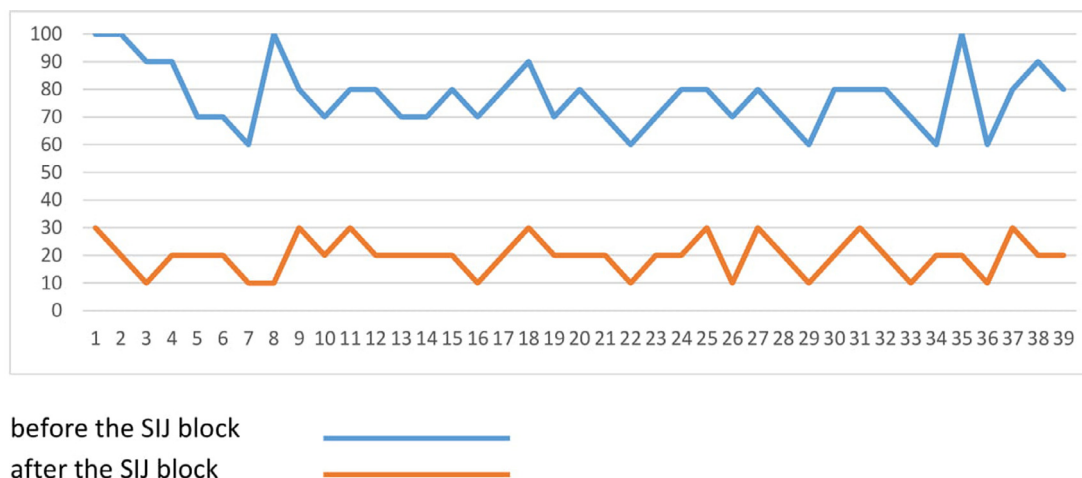


Fig 2. VAS scores before and after the SIJ block in SIJD-positive patients. SIJD, sacroiliac joint dysfunction; VAS, visual analog scale.

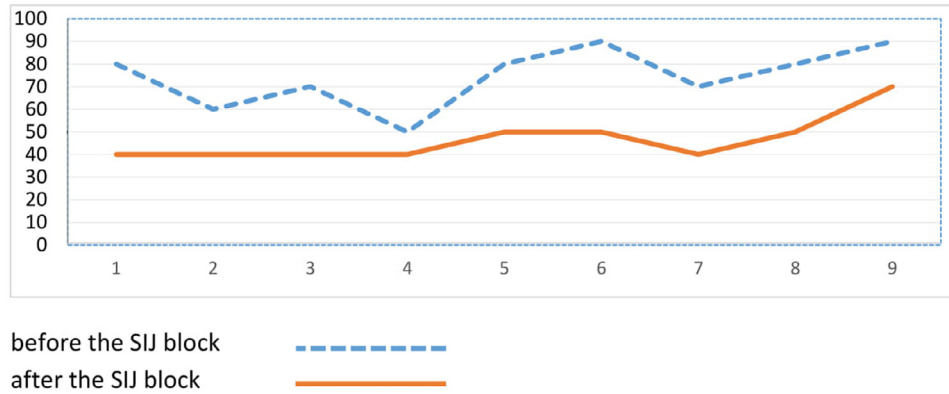


Fig 3. VAS scores before and after the SIJ block in SIJD-negative patients. SIJD, sacroiliac joint dysfunction; VAS, visual analog scale.

Table 2. Sensitivity, Specificity, and Positive and Negative Predictive Values for Each Physical Test

	FABER Test	Thigh Thrust Test	Gaenslen Test	Yeoman Test	Gillet Test	Forward Flexion Test
Sensitivity (%)	71.8	74.4	61.5	64.1	100	100
Specificity (%)	66.7	44.4	33.3	33.3	0	0
PPV (%)	90.3	85.3	80	80.6	81	81
NPV (%)	35.3	28.6	16.7	17.6	-	-

FABER, Flexion, Abduction and External Rotation; NPV, negative predictive value; PPV, positive predictive value.

respectively. The FABER test had the highest specificity and positive predictive values.

Tables 4 and 5 present the distribution of the positivity of each individual physical test and different combinations of these tests, respectively, in SIJD-positive and SIJD-negative patients.

The results for dual combinations of pain provocation tests are shown in Table 6. A combination of FABER and thigh thrust tests showed the highest improvement in the overall diagnostic power (as measured by improvement in the AUC). More specifically, this combination resulted in an AUC of 69.2%, a sensitivity value of 71.7%, and a specificity value of 66%.

Table 3. Sensitivity, Specificity, and Positive and Negative Predictive Values for Different Combinations of Physical Tests

	Three or More Positive Tests	Four or More Positive Tests	Five or More Positive Tests
Sensitivity (%)	94.9	92.3	59
Specificity (%)	11.1	22.2	55.5
PPV (%)	82.2	83.7	85.1
NPV (%)	33.3	40	23.8

NPV, negative predictive value; PPV, positive predictive value.

Table 4. The Positivity of Each Physical Test in SIJD-Positive and SIJD-Negative Patients

Test		SIJD-Positive	SIJD-Negative
FABER test	Positive	28	3
	Negative	11	6
Thigh thrust test	Positive	29	5
	Negative	10	4
Gaenslen test	Positive	15	6
	Negative	24	3
Yeoman test	Positive	25	6
	Negative	14	3
Gillet test	Positive	39	9
	Negative	0	0
Forward flexion test	Positive	39	9
	Negative	0	0

FABER, Flexion, Abduction, and External Rotation; SIJD, sacroiliac joint dysfunction.

Table 5. The Positivity of Different Combinations of Physical Tests in SIJD-Positive and SIJD-Negative Patients

Combinations of Physical Tests	SIJD-Positive	SIJD-Negative
Fewer than 3 positive tests	0	0
Three or more positive tests	37	8
Fewer than 4 positive tests	3	2
Four or more positive tests	36	7
Fewer than 5 positive tests	16	5
Five or more positive tests	23	4
Fewer than 6 positive tests	29	8
Six positive tests	10	1

SIJD, sacroiliac joint dysfunction.

DISCUSSION

The findings of this study revealed that the positive predictive values of the provocation tests were high (larger than 80%) and the sensitivity values of these tests exceeded 60%. Moreover, the thigh thrust test was the most sensitive (74.4%), and the FABER test was the most specific (66.7%).

Studies on the accuracy of the FABER test have been inconsistent in their findings. For example, our study found its specificity to be 66.7%, but Dreyfuss et al⁴⁵ reported a 16% specificity for this test. This significant difference can be attributed to that not only did the authors consider a reduction of larger than 90% in VAS scores as SIJD-positive but also administered a single injection of a local anesthetic and a corticosteroid with long-term effectiveness. Similarly, Broadhurst et al⁵⁹ report a 100% specificity of the FABER test. This is the result of the pain relief criterion used, which was lower and thus more lenient in the current study, and also that they injected the patients who had a positive FABER test and not those with a negative FABER result.⁵⁹

Although we observed a sensitivity value of 71.8% for the FABER test, Broadhurst et al⁵⁹ reported this value to be 77% when the diagnostic criterion had been set at a reduction of 70% in pain, but found a sensitivity of 50% when the criterion was 90% pain relief. In contrast, this test had a lower sensitivity (57%) than in the studies by van der Wurff et al,^{60,61} because the SIJ was injected blindly; blocking under fluoroscopic guidance can increase sensitivity by raising the probability of infiltration into the SIJ.

In this study, the positive and negative predictive values of the FABER test were 90.3 and 35.3%, respectively. In the review article by Cattley et al,⁵² the FABER test was referred to as an unreliable and invalid test in SIJD diagnosis. The authors cited methodological quality, technique application, and VAS pain parameters as reasons for this.

The thigh thrust test had a sensitivity of 74.4% and a specificity of 44.4% in our study. The sensitivity and specificity of this test were 36% and 50%, respectively, in Dreyfuss et al.⁴⁵ Broadhurst et al⁵⁹ reported a sensitivity value of 80% with the pain relief criterion having been set at 70%, and a sensitivity of 69% using a pain relief criterion of 90%. In both cases, study authors observed a specificity value of 100%. The sensitivity, specificity, and positive and negative predictive values obtained in the study by Laslett et al¹³ for the thigh thrust test were 50%, 69%, 58%, and 92%, respectively, using a pain reduction threshold of 80%. The larger cutoff value was associated with a lower reported sensitivity; however, to reduce the probability of false positivity, it is better to use a cutoff value that is as close to 100% as possible in all of the physical tests.

The Gaenslen's test had a sensitivity value of 61.5% and a specificity value of 33.3% in our study. The sensitivity, specificity, and positive and negative predictive values found in Laslett et al¹³ for this test were 37%, 71%, 47%, and 76%, respectively. The specificity of this test in the study by Broadhurst⁵⁹ was reported as 100%, which may be attributed to the use of a different protocol (eg, setting a higher cutoff and the injection of 4 cc of lidocaine being

Table 6. Sensitivity, Specificity, and Positive and Negative Predictive Values and AUC Levels in Dual Combinations of Provocative Tests

	FABER and Thigh Thrust	FABER and Gaenslen	FABER and Yeoman	Thigh Thrust and Gaenslen	Thigh Thrust and Yeoman	Gaenslen and Yeoman
Sensitivity (%)	71.7	48.71	43.58	48.71	43.58	35.89
Specificity (%)	66	77.77	66.66	66.66	55.55	55.55
PPV (%)	90.32	90.47	85	86.36	80.95	15.66
NPV (%)	35.29	25.92	21.42	23.07	18.51	77.7
AUC	69.2%	63.2	55.1	57.7	49.6	45.7

AUC, area under the curve; FABER, Flexion, Abduction, and External Rotation; NPV, negative predictive value; PPV, positive predictive value.

restricted to patients with a positive result on the Gaenslen's test).

Regarding the other tests, the observed discrepancies between the studies may be due to the adoption of different executive approaches. A review of the existing literature indicates that studies have used various thresholds of pain reduction after the SIJ block, ranging from 50% to 90%. For instance, Polly⁵⁶ and van der Wurff⁶⁰ used a 50% pain reduction as a diagnostic criterion for SIJD, but Irwin et al²⁶ set a pain reduction of 70% after administering the confirmatory SIJ block. Schwarzer et al⁶² employed a single-injection SIJ block and set a threshold of 75% reduction in pain; Maigne et al⁶ used a reduction of 75% but with a double block. Slipman et al¹⁸ and Young et al⁶³ used a reduction of 80% in pain after a single block. Lastly, Dreyfuss et al⁴⁵ used a single injection of a local anesthetic and corticosteroids, in addition to pain provocation tests, and adopted a threshold of 90% reduction in pain severity. In the present study, the threshold was set at 60% because all of the SIJD-suspected patients received the double block and this study used established, stringent inclusion criteria for identifying SIJD-suspected patients, believing that these measures can justify a threshold lower than those set in more lenient studies.²⁸ Indeed, studies that only used pain reduction without any other diagnostic criteria had to set a higher threshold for diagnosing positive SIJD.^{19,41,46,59}

Some studies used the single block,^{18,19,59} whereas others^{5,6,13,26,39,42,64} (including the current study) adopted the double block approach. This study used the double block because it has been demonstrated that the prevalence of SIJ pain is estimated to range between 10% and 38% using a double block paradigm, whereas the false-positive rate of the single block is 20% to 54%.²⁸

Additionally, in some studies,⁴ corticosteroids were injected instead of or in combination with lidocaine,¹⁹ which can influence the results of the SIJ block. More specifically, corticosteroids have a delayed onset of action but can reduce pain more effectively than lidocaine, which results in almost instant pain relief.⁶⁵

Although the SIJ block is considered the gold standard in the diagnosis of SIJD, in 20% or sometimes up to 50% of cases, the SIJ block yields false-positive results, which can overestimate the sensitivity of the alternative diagnostic tools.^{16,28,66,67,68} These false results are caused by the extravasation of the locally injected anesthetic to the surrounding structures, such as ligaments, muscles, and lumbosacral nerve roots, potential sources of pain. The vertical position of the SIJ makes this joint prone to the leakage of the anesthetic, and leakage may occur regardless of the type of SIJ block (eg, fluoroscopy, computed tomography scan, sonography, and MRI) employed.²⁸ Thus, gathering supplemental forms of evidence through multiple tests can help to diagnose SIJD more reliably.

According to the findings of this study, a combination of the motion and provocation tests increased specificity and

positive predictive values but decreased sensitivity and negative predictive values. This is particularly considerable if there are several tests being combined. Furthermore, a combination of 3 or more positive provocation tests plus at least 1 positive motion palpation test produced the best results.

Laslett et al⁴⁰ showed that patients with a positive diagnostic SIJ block are at least 3 × and as much as 20 × as likely to have had a combination of 3 or more positive provocative tests than patients with a negative SIJ block. Similarly, Laslett et al¹³ found that the positivity of 2 of 4 tests (ie, distraction, compression, thigh thrust, or sacral thrust) or 3 or more provocation-motion tests were the best predictors of a positive SIJ block.

In the present study, a combination of FABER and thigh thrust tests was more accurate than any other combination. The next most accurate results were obtained from a combination of FABER and Gaenslen tests. Because FABER and thigh thrust tests had the highest sensitivity and specificity values (see Table 2), it seems reasonable to use a combination of these 2 tests for SIJD diagnosis.

Here, Gillet and forward flexion tests had a sensitivity value of 100% and 0 specificity. This is owing to the inclusion of patients with at least 1 positive palpation test in accordance with our eligibility criteria. It turned out that Gillet and forward flexion tests were both positive in all of the included patients. The positive predictive value of both these tests was 81%.

It was also found that the results of neither the Gillet test nor the forward flexion test changed after the SIJ block. This was because these tests indicate mobility of the ilium and the sacrum and are not influenced by the SIJ block. It seems that in SIJD cases without a biomechanical disturbance, these motion tests are not necessarily positive and also may be positive in patients without SIJD.^{3,50} Thus, it is reasonable to use a combination of provocation tests and motion palpation tests for SIJD diagnosis.

In contrast, Dreyfuss et al⁶⁹ showed the false positivity of motion tests by observing that they were positive in 20% of asymptomatic patients. They also found that the sensitivity and specificity of the standing flexion and Gillet tests were poor¹⁹ because they were compared with the SIJ block as the gold standard. In this regard, other studies^{70,71} compared motion tests with provocation tests and showed that agreement ranged from 67% to 97% for pain provocation tests but was 48% for palpation tests. Furthermore, kappa values varied between 0.43 and 0.84 for provocation tests but were -0.06 for palpation tests. Vanelderden et al also expressed that in the presence of a weak predictive value of provocation tests, combined batteries of physical tests can help ascertain SIJD diagnosis.⁷² Hence, it can be concluded that joint hypomobility leading to positive motion tests does not mean a patient has SIJD, but the positivity of at least 3 provocation tests and at least 1 motion test increases the predictivity of SIJD.

Among the provocation tests, a combination of FABER and thigh thrust tests was more successful in diagnosing SIJD. The authors believe that a combination of physical tests in addition to patient history findings and clinical data can compensate for the low diagnostic power of these tests.

Limitations

A major limitation of the present study is that the adoption of at least 1 positive palpation test as a diagnostic criteria resulted in the 0 specificity of the palpation tests in SIJD-negative patients. Another limitation is the low cutoff point (60%) in pain reduction. If this point was more than 60%, this may have resulted in lower sensitivity in each test. If the number of participants and thus the number of SIJ blocks was higher, there could have been various results regarding the accuracy of the physical tests.

CONCLUSION

This study found that the FABER test had the highest single specificity and positive predictive values of all of the provocation tests under discussion. Further, a combination of the FABER and thigh thrust tests improved the overall diagnostic power. In addition, greater numbers of positive physical tests contribute more to an SIJD diagnosis. It is also recommended that at least 3 provocation tests be used in addition to motion palpation tests when confirming an SIJD diagnosis. Finally, as the existing literature reports inconclusive findings for the use of individual physical tests, it is advisable to use a combination of such tests in conjunction with other sources of data, including patient history, symptoms, and imaging, to diagnose SIJD.

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CONTRIBUTORSHIP INFORMATION

Concept development (provided idea for the research): P.N.
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Supervision (provided oversight, responsible for organization and implementation, writing of the manuscript): P.N., F.I.

Data collection/processing (responsible for experiments, patient management, organization, or reporting data): E.S., M.S.

Analysis/interpretation (responsible for statistical analysis, evaluation, and presentation of the results): E.S., L.N.

Literature search (performed the literature search): E.S., R.M.

Writing (responsible for writing a substantive part of the manuscript): P.N., E.S., L.N.

Critical review (revised manuscript for intellectual content, this does not relate to spelling and grammar checking): P.N., F.I., M.S.

Other (list other specific novel contributions): R.M. (reporting the MRI of the patients)

Practical Applications

- In this study, using 2 palpation tests of Gillet and forward flexion was not suitable for diagnosing sacroiliac joint dysfunction.
- The prevocational tests were more reliable than the Gillet and forward flexion that are a marker of anterior rotation of ilium on the sacrum.
- The combination of the provocative tests had more sensitivity than each test solitarily.

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