



## The Effect of Kinesiotaping on Pain, Functionality and Ultrasound Parameters in Patients with Shoulder Impingement Syndrome: A Randomised Sham-controlled Study

Omuz Sıkışma Sendromlu Hastalarda Kinezyobantlamanın Ağrı, Fonksiyonellik ve Ultrason Parametreleri Üzerine Etkisi: Randomize Sham-kontrollü Bir Çalışma

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### Abstract

**Objective:** We investigated the effect of kinesiotaping on pain, functionality and ultrasound parameters in patients with shoulder impingement syndrome (SIS).

**Materials and Methods:** Seventy-five patients with SIS were randomly classified into the following three groups: Kinesiotaping (KT), conventional exercise (CE) and sham-kinesiotaping (sham-KT). Each group underwent a two-week treatment programme. The patients were then evaluated in terms of pain analysed using the visual analogue scale (VAS), joint range of motion and disabilities of the arm, shoulder and hand (DASH) questionnaire, before and after treatment. In addition, the supraspinatus tendon (SsT) thickness and acromiohumeral distance (AHD) parameters were measured using ultrasonography (US).

**Results:** There was a statistically significant difference between the groups in terms of post-treatment VAS scores obtained at night and during activity. According to the post-hoc analysis, these differences were observed in the CE group ( $p<0.016$ ). Because there was a difference in the pre-treatment DASH scores between the groups, the groups were examined for changes in the DASH score before and after treatment; a significant difference between sham-KT and KT groups and between sham-KT and CE groups favouring KT and CE groups was observed. US revealed that the CE group was superior to the sham-KT group in terms of both reduction in SsT thickness and increase in AHD ( $p<0.016$ ). Furthermore, KT was effective in increasing AHD.

**Conclusion:** KT was superior to sham-KT in terms of all parameters except pain. KT was also found to be as effective as CE in all parameters. In addition, US objectively revealed that the supraspinatus tendinitis can be reduced and AHD can be increased. KT is an alternative treatment option in patients with SIS and can be used alone or in combination with CE in patients who do not comply with CE.

**Keywords:** Kinesiotaping, shoulder impingement syndrome, ultrasound

### Öz

**Amaç:** Omuz impingement (sıkışma) sendromu (OİS) olan hastalarda kinezyobantlamanın (KT) ağrı, fonksiyonellik ve ultrason parametreleri üzerine etkisini araştırmayı amaçladık.

**Gereç ve Yöntem:** Toplam 75 OİS hastası randomize olarak üç grupta sınıflandırıldı: KT, konvansiyonel egzersiz (KE) ve sham-kinezyobantlama (sham-KT). Her gruba iki haftalık tedavi programı uygulandı. Hastalar tedavi öncesi ve sonrası görsel analog skala (GAS), eklem hareket açıklığı, kol, omuz ve el sorunları (DASH) anketi kullanılarak değerlendirildi. Ayrıca supraspinatus tendonu (SsT) kalınlığı ve akromiohumeral mesafe (AHM) parametreleri ultrasonografi (US) kullanılarak ölçüldü.

**Bulgular:** Gruplar arasında tedavi sonrası gece ve aktivite-GAS skorları açısından istatistiksel olarak anlamlı fark vardı. Post-hoc analize göre, bu farklılıklar KE grubu lehine gözlenmiştir ( $p<0,016$ ). Gruplar arasında tedavi öncesi DASH skorlarında bir fark olduğu için, gruplar tedavi öncesi ve sonrası DASH skorundaki değişiklikler açısından incelendi; sham-KT ve KT ile sham-KT ve KE grupları arasında KT ve KE grupları lehine anlamlı bir fark tespit edilmiştir. US, KE grubunun hem SsT kalınlığındaki azalma hem de AHD'deki artış açısından sham-KT grubundan daha üstün olduğunu ortaya koymuştur ( $p<0,016$ ). Ayrıca, KT AHD'nin artmasında etkili olmuştur.

**Sonuç:** KT, ağrı dışındaki tüm parametrelerde sham-KT'den üstün görülmüştür. KT'nin tüm parametrelerde KE kadar etkili olduğu bulunmuştur. Ek olarak, US nesnel olarak supraspinatus tendinitinin azaltılabileceğini ve AHD'nin artırılabilirliğini ortaya koymuştur. KT, OİS'li hastalarda alternatif bir tedavi seçeneğidir ve KE ile uyumlu olmayan hastalarda tek başına veya KE ile kombinasyon halinde kullanılabilir.

**Anahtar kelimeler:** Kinezyobantlama, omuz sıkışma sendromu, ultrason

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**Received/Geliş Tarihi:** 02.01.2020 **Accepted/Kabul Tarihi:** 16.04.2020

## Introduction

Shoulder impingement syndrome (SIS) is one of the most common causes of shoulder pain (1). Shoulder pain associated with SIS ranges from simple pathologies such as subacromial bursitis to rotator cuff tendinopathy and full fold tear (2). Conservative treatment methods include non-steroidal anti-inflammatory drugs, steroid injections to the subacromial region, physical therapy modalities, and conventional exercise (CE) (3-5). Kinesiotaping (KT) has recently become more widespread and has been used to treat various musculoskeletal pathologies (6,7). Studies have shown that KT reduces pain and increases the range of motion (ROM) in SIS, particularly in the early period (8). Kinesiotape is an advantageous method as it is non-invasive and can be applied easily and in a short time (8). Clinical parameters have been used for evaluation in previous studies comparing the effectiveness of KT (8). The aim of the current study, was to compare the efficacy of KT with both sham application and exercise therapy using ultrasonographic measurements as objective data in addition to clinical parameters.

## Materials and Methods

### Participants

The study included patients aged 45-70 years who were admitted to our outpatient clinic with complaints of shoulder pain, were diagnosed with SIS based on physical examination and magnetic resonance imaging results, and were eligible

based on the inclusion and exclusion criteria. The inclusion criteria were as follows: (1) at least three positive results in the Hawkins-Kennedy, Neer, empty can, drop-arm, and lift-off tests, (2) magnetic resonance imaging findings, and (3) age between 45 and 70 years. Patients were excluded from the study if they had received physical therapy for the shoulder region within the past three months, had a history of injections to the shoulder joint, had any cervical pathologies, clinical conditions accompanied by neuromotor or sensory dysfunction, a history of malignancy, were pregnant, had a partial or total rupture in the supraspinatus tendon (SsT), adhesive capsulitis, diabetes, chronic liver disease, or kidney failure. Demographic data (age, gender, dominant side, disease duration, and occupational status) were recorded at the beginning of the study.

All patients provided written consent prior to study initiation. The details of the numbers of included and excluded patients through to the final data analysis are shown in Figure 1 as a flowchart. Approval for the study was granted by the Haydarpaşa Numune Training and Research Hospital Clinical Research Ethics Committee (decision no: HNEAH-KAEK 2016/98).

### Design

This prospective, randomized, single-blind trial was conducted at a single center between April 2017 and October 2017. A total of 75 patients were included in the study and were randomly assigned to one of the following three groups using a table of random numbers: KT, CE, and sham-KT groups. The study was completed with no drop-outs.

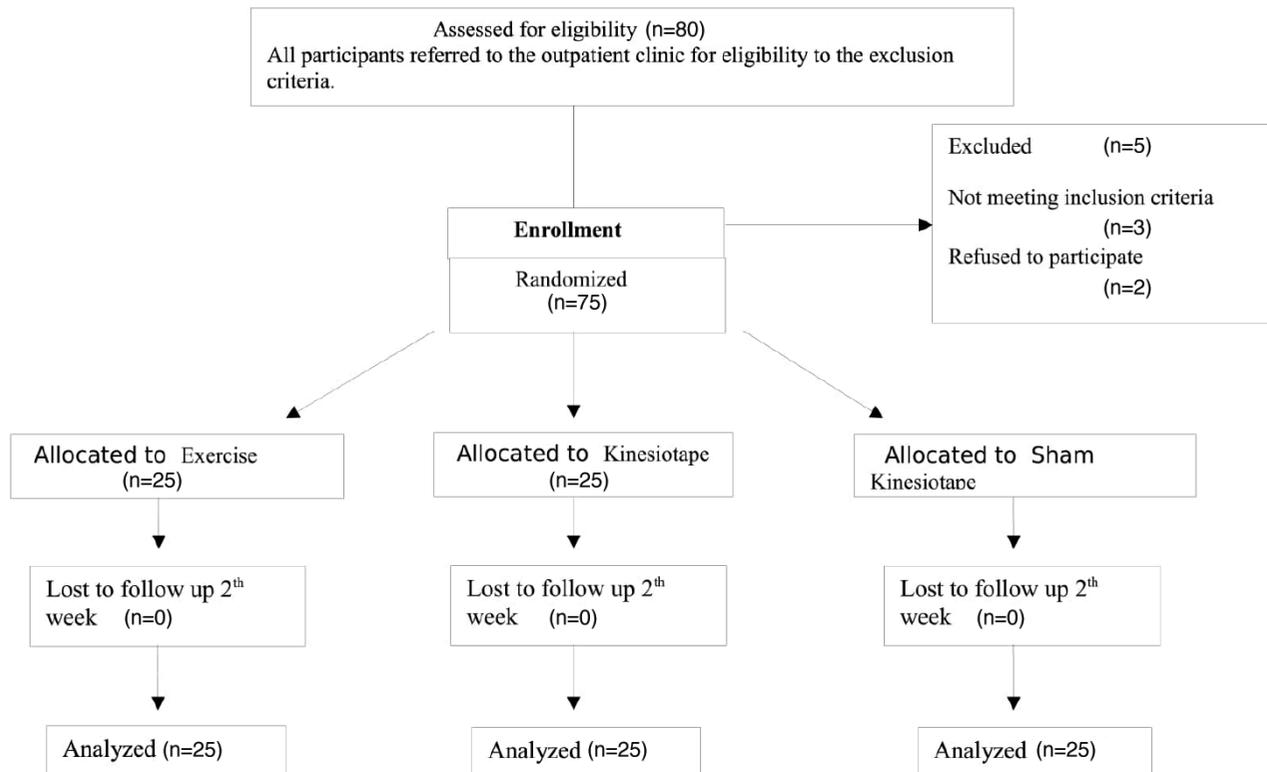


Figure 1. Flow diagram for study participants

All participants were assessed by the same clinician (F.B.) at baseline and at 2 weeks after completing the interventions using a visual analog scale (VAS), ROM and disabilities of the arm, shoulder and hand (DASH) questionnaire. All the KT and sham-KT applications were made by a single clinician who was blinded to the group allocation (N.M.). The ultrasonographic (US) measurements were taken by a single clinician (D.G.K.), who was blinded to the group allocation.

In the power analysis, based on the VAS score determined during activity and considering a dropout rate of 20%, 25 patients in each group were expected to result in a 5% type 1 error rate. The power of the test was expected to be 80%. MedCalc Statistical Software version 12.7.7 (MedCalc Software Bvba, Ostend, Belgium; <http://www.medcalc.org>; 2013) was used for these measurements.

### Interventions

**Group 1:** Cold application, KT treatment (2 sessions with a 5-day interval)

**Group 2:** Cold application, CE treatment (3 sessions per day for 2 weeks)

**Group 3:** Cold application, sham-KT treatment (2 sessions with a 5-day interval)

### Treatments Applied to the Patients

**Cold application:** At the beginning of each treatment session, gel ice packs were wrapped in a damp towel and applied to the affected shoulder joint for 20 minutes.

**Group 1:** KT was applied to the deltoid muscle using the inhibition and mechanical correction technique and to the supraspinatus muscle using the inhibition technique (Two sessions with a 5-day interval) (9).

**Deltoid muscle inhibition technique:** The Y-shaped kinesiotape was applied 3 cm below the deltoid tuberosity of the humerus without tension. The Y-band was applied to the forearm with 15%-25% light stretching along the outer edge of the anterior deltoid up to the acromioclavicular joint and with 15%-25% light stretching along the outer edge of the posterior deltoid up to lateral aspect of the acromion. The last 3-5 cm of the tails was applied to the adhesion sites without any tension. When performing the procedure, the shoulder was maintained in transverse extension and external rotation (ER) for the forearm of the Y-strip and 45° of transverse flexion (FLX) and horizontal adduction for the posterior arm of the Y-strip (Figure 2a).

**Supraspinatus muscle inhibition technique:** The Y-shaped kinesiotape was applied 3 cm below the greater tubercle of the humerus without tension. The patient was instructed to adduct the shoulders simultaneously bringing the neck to lateral FLX on the opposite side. The initial section of the tape was adhered to the greater tubercle below the acromion with submaximal (75%) tension. The lower arm of the Y-strip was applied to the medial edge of the scapula along the spina scapula without tension. The upper arm was applied to the spina scapula superior between the upper and lower trapezoidal muscles along the medial edge of the scapula in the supraspinatus fossa without stretching. The

shoulder was in protraction and internal rotation (IR) during the application (Figure 2b).

**Mechanical correction technique:** The Y-shaped kinesiotape was applied with the arm in the neutral position along the torso. The application was started from the coracoid process without stretching. The upper arm of the Y-strip was applied over the acromioclavicular joint to the outer edge of the posterior deltoid with maximal stretching, with the last 3-5 cm applied without any stretching; the lower arm of the Y-strip was applied several centimeters below the upper arm with the same technique (Figure 2c).

**Group 2:** CE treatment was administered for 10 days with 3 sessions/day. A triphasic exercise program was administered to the patients. Before starting the exercise program, the patients were instructed not to perform overhead movements exceeding 90°. The exercise program was initiated using Codman pendulum, passive joint motion range (with a 1-m stick), and posterior capsule stretching exercises. Shoulder wheel, finger ladder, and shoulder strengthening exercises using a theraband were added to the exercise programs of patients with full or near total ROM and pain relief. Exercise was administered twice a week under supervision and the patients were advised to exercise at home on the other days with 20 repetitions of each exercise. The patients were followed up via telephone to make sure they were adhering to the exercise program (10).

**Group 3:** Sham-KT was applied in 10 cm I-shaped strips on the sagittal plane over the acromioclavicular joint without stretching and on the transverse plane distal to the deltoid area. The kinesiotape was applied twice with a 5-day interval (Figure 2d).

### Evaluation Parameters

All patients were evaluated before and after treatment.

**1. Pain level:** The severity of shoulder pain (resting, activity, and night pain) of the patients was evaluated using the VAS score. The patients were asked to mark the average severity of the pain they felt during the past week on a 10 cm ruler, where 0: no pain and 10: the most severe pain. The marked points were recorded.

**2. Functional status:** DASH questionnaire comprises three sections. The first section includes 30 items: 21 items assess the patient's difficulties in performing daily activities, 5 assess symptoms (pain, activity-related pain, tingling, stiffness, and weakness), and each of the remaining 4 items assesses social function, work, sleep, and self-confidence. All items are rated on a 5-point Likert-type scale (1: no difficulty, 2: mild difficulty, 3: moderate difficulty, 4: extreme difficulty, 5: cannot perform at all). The total score ranges from 0 to 100 (0: no disability, 100: maximum disability). Turkish reliability and validity studies have been performed for this questionnaire (11).

**3. Joint range of motion measurements:** FLX, abduction (ABD), IR and ER were measured using a goniometer (Saehan goniometer - plastic).

**4. Ultrasonography:** US was performed using a 7.5-mHz linear probe in the B mode (Mindray-China). SsT thickness was

measured at three different points (10, 15, and 20 mm) lateral to the tendon after identifying the biceps tendon in the transverse section and the average of measurements was recorded (Figure 3a). Acromiohumeral distance (AHD) was assessed by measuring the linear distance between the inferior of the acromion from the anterior of the shoulder and the superior of the humeral head (Figure 3b) (12).

### Statistical Analysis

All analyses were performed using the MedCalc Statistical Software version 12.7.7 (MedCalc Software bvba, Ostend, Belgium; <http://www.medcalc.org>; 2013). Descriptive statistics were used to describe continuous variables (average, standard deviation, minimum, median, and maximum values). Conformity of the data to normal distribution was assessed with the Kolmogorov-Smirnov test. As the data were not normally distributed, the relationship between two dependent continuous variables was investigated using the Wilcoxon Signed Rank test, and the Kruskal-Wallis test was used to compare continuous variables among many groups. In cases where the Kruskal-

Wallis test revealed a significant difference, post-hoc analysis was conducted using the Mann-Whitney U test with Bonferroni correction. A value of  $p < 0.016$  was considered statistically significant.

The correlation between dependent and categorical variables was examined using the McNemar test. The chi-square or Fisher's Exact test, as appropriate, was used to examine the correlation between independent categorical variables. The comparison of two independent variables was performed using the Mann-Whitney U test. The level of statistical significance was set at  $p < 0.05$ .

### Results

**1. Demographic data:** No significant difference was determined between the groups in terms of demographic characteristics (Table 1).

**2. Pain parameters:** VAS: No difference was determined between the groups in respect of the resting, activity and night-VAS scores before treatment. In the post-treatment night- and



**Figure 2.** Application of kinesiotaping. (a) Deltoid muscle inhibition technique; (b) supraspinatus muscle inhibition technique; (c) deltoid muscle mechanical correction technique; (d) sham application technique



**Figure 3.** (a) Ultrasonographic measurement of supraspinatus tendon thickness. (b) Ultrasonographic measurement of acromiohumeral distance

activity-VAS scores, a statistically significant difference was determined between the groups. According to the post-hoc analysis, these differences were found between the sham-KT and CE groups and in favor of the CE group ( $p < 0.016$ ) (Table 2).

**3. Functional status:** As there was a difference between the groups in the pre-treatment DASH scores, the changes in DASH scores after treatment were examined in terms of differences between the groups. A significant difference was found between the sham-KT and KT groups and between the sham-KT and CE groups in favor of the KT and CE groups, respectively (Table 2).

**4. Joint range of motion measurements:** Pre-treatment, there was no difference between the groups in terms of joint ROM in all directions. Following treatment, a statistically significant difference

was determined between the groups in respect of the FLX angles. According to the post-hoc analysis, this difference was found between the sham-KT and KT groups and between the sham-KT and CE groups in favor of the KT and CE groups, respectively ( $p < 0.016$ ) (Table 3). There were statistically significant differences between the groups in terms of the post-treatment ABD and ER angles. According to the post-hoc analysis, this difference was found between the sham-KT and CE groups in favor of the CE group ( $p < 0.016$ ). There was no difference between the groups in terms of the post-treatment IR angles.

**5. Ultrasonographic measurements:** As there was a difference between the groups in terms of the pre-treatment SsT thickness and AHD measurements, it was analyzed

**Table 1. Demographic features of the groups**

Parameters		KT group (n=25)	CE group (n=25)	Sham-KT group (n=25)	<sup>a</sup> p
Dominant side, n (%)	Right	24 (96%)	25 (100%)	23 (92%)	0.769
	Left	1 (4%)	0 (0.0)	2 (8%)	-
Patient side, n (%)	Right	17 (68%)	17 (68%)	14 (56%)	0.717
	Left	8 (32%)	8 (32%)	11 (44%)	-
Gender, n (%)	Woman	16 (64%)	15 (60%)	19 (76%)	0.555
	Man	9 (36%)	10 (40%)	6 (24%)	-
Occupation, n (%)	Housewife	10 (40%)	11 (44%)	15 (60%)	0.104
	Retired	5 (20%)	9 (36%)	6 (24%)	-
	Civil servant	5 (20%)	0 (0.0)	0 (0.0)	-
	Worker	5 (20%)	5 (20%)	4 (16%)	-
Age, years, range		50 (46-70)	58 (52-70)	56 (54-70)	<sup>b</sup> 0.571
BMI, (kg/m <sup>2</sup> ), range		27 (17-35)	26 (21-36)	27 (15-37)	0.952

BMI: Body mass index, KT: Kinesiotaping, CE: Conventional exercise, <sup>a</sup>Ki-kare test, <sup>b</sup>Mann-Whitney U test

**Table 2. Visual analogue scale and the disabilities of the arm, shoulder and hand scores at pre- post-treatment visit**

Parameters	KT group (n=25)	CE group (n=25)	Sham-KT group (n=25)	<sup>a</sup> p
	Median (min-max)	Median (min-max)	Median (min-max)	
VAS rest, mm (baseline)	6 (0-10)	7 (0-8)	5 (0-10)	0.794
VAS rest, mm (2 <sup>nd</sup> week)	4 (0-8)	2 (0-5)	5 (0-10)	0.110
<sup>b</sup> p	0.001	<0.001	0.011	-
VAS night, mm (baseline)	8 (0-10)	8 (5-10)	8 (0-10)	0.945
VAS night, mm (2 <sup>nd</sup> week)	4 (0-10)	3 (0-7)	7 (0-10)	0.017
<sup>c</sup> p	<0.001	<0.001	0.005	-
VAS activity, mm (baseline)	8 (1-10)	8 (0-10)	7 (3-10)	0.947
VAS activity, mm (2 <sup>nd</sup> week)	5 (0-9)	3 (0-8)	6 (3-10)	0.001
<sup>c</sup> p	<0.001	<0.001	0.005	-
Q-DASH (baseline)	60 (29-90)	67 (38-80)	55 (28-75)	0.023
Q-DASH (2 <sup>nd</sup> week)	30 (0-65)	25 (0-80)	48 (23-71)	0.001
<sup>c</sup> p	<0.001	<0.001	0.001	-
<sup>c</sup> Q-DASH pre-post (difference)	30 (0-53)	38 (0-59)	2 (0-52)	<0.001

VAS: Visual analogue scale, Q-DASH: The disabilities of the arm, shoulder and hand questionnaire, KT: Kinesiotaping, CE: Conventional exercise, min: Minimum, max: Maximum, <sup>a</sup>Kruskal-Wallis test, <sup>b</sup>Wilcoxon Signed Rank test, <sup>c</sup>Post-hoc analysis-Bonferroni test

whether the post-treatment changes in these measurements differed between the groups. A significant difference was found between the groups for both parameters. According to post-hoc analyses, there was a statistically significant difference between the sham-KT and CE groups and between the KT and CE groups in favor of the CE group in terms of SsT thickness decrease ( $p < 0.016$ ). In terms of AHD increase, there was a significant difference between the sham-KT and KT groups and the sham-KT and CE groups in favor of the KT and CE groups ( $p < 0.016$ ) in Table 4.

### Discussion

Assessments were made of pain, functionality, ROM, physical examination findings, SsT thickness and AHD in patients with SIS and it was observed that the application of KT resulted in significant improvements in all parameters. CE is reportedly effective as a treatment for pain, ROM, and functionality in patients with SIS (13,14). Therefore, the aim of this study was

to investigate the efficacy of KT application in comparison with not only sham-KT application but also with CE, which is known to be effective in patients with SIS. KT is frequently used as a component of rehabilitation programs for SIS and rotator cuff tendinitis (15). In these pathologies, the primary aim is to reduce edema and pain and increase the ROM and muscle activity.

Studies have shown that KT reduces pain in patients with SIS and increases the joint ROM, particularly in the early period (16-19). Thelen et al. (18) assessed the efficacy of KT application on pain, disability, and painless active joint ROM in patients with SIS or rotator cuff tendinitis. KT showed a similar effect to that of sham-KT in all parameters except for painless shoulder ABD. In contrast, Şimşek et al. (19) compared the efficacy of KT and sham-KT in patients with SIS and found that KT was superior to the sham-KT group in terms of pain and functionality. In this study, CE treatment was applied to both groups. Kaya et al. (20) compared KT and manual therapy in patients with SIS in terms of disability and pain and reported that compared with manual

**Table 3. Comparison of joint range of motion measurements at pre- post-treatment visit**

Parameters	KT group (n=25)	EX group (n=25)	Sham-KT group (n=25)	ap
	Median (min-max)	Median (min-max)	Median (min-max)	
FLX, degree (baseline)	180 (90-180)	180 (90-180)	150 (120-180)	0.680
FLX, degree (2 <sup>nd</sup> week)	180 (140-180)	180 (120-180)	160 (120-180)	0.001
<b>bp</b>	0.003	0.003	0.011	-
ABD, degree (baseline)	150 (80-180)	130 (90-180)	150 (80-180)	0.484
ABD, degree (2 <sup>nd</sup> week)	170 (90-180)	180 (140-180)	150 (90-180)	0.007
<b>p</b>	0.003	<0.001	0.017	-
IR, degree (baseline)	70 (20-90)	70 (20-70)	70 (40-70)	0.600
IR, degree (2 <sup>nd</sup> week)	70 (40-90)	70 (40-70)	70 (40-70)	0.556
<b>p</b>	0.041	0.007	0.317	-
ER, degree (baseline)	90 (30-90)	90 (30-90)	90 (40-90)	0.140
ER, degree (2 <sup>nd</sup> week)	90 (60-90)	90 (40-90)	90 (40-90)	0.008
<b>p</b>	0.018	0.066	0.317	-

FLX: Flexion, ABD: Abduction, IR: Internal rotation, KT: Kinesiotaping, ER: External rotation, min: Minimum, max: Maximum, <sup>a</sup>Kruskal-Wallis test, <sup>b</sup>Wilcoxon Signed Rank test

**Table 4. Comparison of ultrasonographic measurements at pre- post-treatment visit**

Ultrasound measurements	KT group (n=25)	CE group (n=25)	Sham-KT group (n=25)	p
SsT thickness, mm median (min-max) (baseline)	6.6 (5.3-8.8)	7.6 (4.2-13)	7.3 (5-12)	0.043
SsT thickness, mm median (min-max) (2 <sup>nd</sup> week)	6.6 (5-8.8)	7.5 (4.2-13)	7 (5-12)	0.099
<b>p</b>	0.024	<0.001	0.017	-
AHD, mm median (min-max) (baseline)	11.7 (9.8-16.7)	13.5 (9.9-16.3)	13 (11.1-16.4)	0.020
AHD, mm median (min-max) (2 <sup>nd</sup> week)	12 (10-16.7)	13.8 (10.5-16.6)	13 (11.1-16.4)	0.022
<b>p</b>	0.002	<0.001	0.577	-
<sup>c</sup> SsT Thickness, mm (baseline -2 <sup>nd</sup> week difference)	-0.2 (0-0.5)	0 (0-0.4)	0 (0-0.3)	0.002
AHD, mm (baseline -2 <sup>nd</sup> week difference)	0.2 (-1-0.2)	0 (-0.4-0.4)	0 (0-0.9)	<0.001

SsT: Supraspinatus tendon, AHD: Acromiohumeral distance, KT: Kinesiotaping, CE: Conventional exercise, min: Minimum, max: Maximum, <sup>a</sup>Kruskal-Wallis test, <sup>b</sup>Wilcoxon Signed Rank test, <sup>c</sup>Post-hoc analysis - Bonferroni test

therapy, KT had a considerable effect on pain reduction after the first week of treatment initiation, although both groups showed similar improvements in the pain and functionality parameters after the second week. In the current study, both groups received CE treatment. Kaya et al. (20) considered this early pain reduction effect observed with KT to be a positive advantage because it would also increase the performance of CE application. In the current study, considerable improvements were observed in the pain and disability scores of all three groups. When the groups were compared in terms of pain scores, it was observed that the CE group but not the KT group was superior to the sham-KT group. A significant improvement was also observed in the functionality scores in all three groups, and the improvements in both the KT and CE groups were determined to be superior to those of the sham-KT group.

There are certain theories explaining the role of KT in pain relief. The first theory is the reduction of subcutaneous nociceptor pressure in the skin. KT also provides afferent stimulation on soft tissue. Thus, the gate control mechanism is activated with afferent feedback. By regulating superficial and deep fascia functions, it reduces edema and inflammation, thereby producing analgesic effects (17,21). In the current study, the pain parameters in the sham-KT group also showed improvement. Sham-KT applied to the same muscle can produce analgesic effects because of the appropriate sensory feedback during the movement of the muscle. This in turn reduces mechanical irritation in soft tissues (22,23). Furthermore, this sensory feedback simultaneously increases patient awareness and compliance with ergonomic principles (17,21). Therefore, sham-KT application is recommended using a different band or to a different area. However, application to a different area may disrupt the blinding process of a study. Using tapes with different characteristics would be more suitable in prospective studies. Of the aforementioned studies related to SIS, only the study conducted by Thelen et al. (18) directly compared isolated KT and sham-KT applications. In other studies, the patients were also administered CE treatment, which eliminates the chance of observing the isolated effect of KT application on pain and functionality. The limitation of the study by Thelen et al. (18) was that it did not compare KT application with another treatment method with proven efficacy in terms of evidence-based medicine. In the current study, this limitation was considered and a third group was treated with CE therapy alone, as it has proven efficacy on SIS. The results of this study, which directly compared KT application with CE treatment, concluded that KT is as effective as CE in terms of improving the pain scores and shoulder disability index.

The current study results showed an increase in all joint ROM parameters in the KT group. In terms of the joint ROM, the ABD angles were found to be significantly higher in the CE group than in the sham-KT group. In the sham-KT group, the FLX and ABD angles showed improvement. The strengthening of the motor unit caused by an increased proprioceptive stimulus by the tape

and a subsequent increase in motion may lead to an increase in the ABD angle (17,24). The improvements in the joint ROM can be considered to have occurred through the restoration of damaged and irregular fascia by KT application and pain relief. Fascial correction with KT allows the fascia movements to be guided to the desired direction and alignment (25). From the results of the current study, it was thought that the shoulder was guided to the glenohumeral motion arch with the applied KT method, and that there was a simultaneous reduction in mechanical irritation in the affected soft tissue structures, leading to increased joint ROM (26,27).

The AHD and SsT thickness measurements were also evaluated using US. Michener et al. (12) also used US to evaluate SsT thickness and AHD in patients with SIS and reported SsT thickness as 6.6 mm and AHD as 10.8 mm. In the control group, SsT thickness and AHD were measured as 6 and 11.4 mm, respectively. In previous studies investigating patients with SIS, SsT thickness has been observed to be between 5.6 and 8.1 mm. In the current study, the pre-treatment SsT thickness measured using US was 7.1 mm, which is consistent with values reported in literature. Pre-treatment AHD was measured as 12.7 mm, which was slightly higher than the value reported by Michener et al (12). In two previous studies, SsT thickness was found to be 1.1-1.5 mm thicker than in the control group (28,29). Kaya et al. (20) compared KT and manual therapy in patients diagnosed with SIS, evaluated SsT thickness with pre-treatment and post-treatment US, and found no significant changes. In the current study, a decrease in SsT thickness was detected in all three groups. When the groups were compared, the reduction was found to be considerably higher in the KT group than in the sham-KT group. Pre-treatment AHD measurements were 13, 11.7, and 13.5 mm (mean, 12.7 mm), whereas post-treatment AHD measurements were 13, 12, and 13.8 mm (mean, 12.9 mm). AHD was significantly increased in the CE and KT groups. To the best of our knowledge, no other study has yet evaluated AHD using US before and after treatment in patients with SIS. Therefore, this study, in which SsT thickness and AHD were evaluated using US before and after treatment in patients with SIS, can be considered to contribute to the literature. The decrease in SsT thickness and the increase in AHD were considerably higher in the CE group than in the other groups, which was thought to be due to the effect of the CE program administered to patients at the time of SIS diagnosis. Strengthening of the muscles providing glenohumeral stabilization eliminated the strain on the supraspinatus muscle, thereby decreasing tendinitis symptoms, and this was observed as a decrease in the tendon thickness in the US measurements. The increase in AHD observed in the KT group was attributed to the effect of the inhibition technique applied to the deltoid muscle, which elevated the humerus, and the mechanical correction technique, which restored the protracted position of the shoulder.

## Study Limitations

Limitations of the study; patients were evaluated at baseline and at 2 weeks after treatment only and there was no follow-up evaluation.

## Conclusion

The results of this study demonstrated that with the exception of the pain parameter, KT was superior to sham-KT in patients with SIS in all the other parameters. It was also found to be as effective as CE in terms of all parameters. In addition, the US measurements performed in this study objectively revealed that tendinitis in the supraspinatus could be relieved and that AHD could be increased. KT is an alternative treatment option in patients with SIS and can be used alone as well as safely in combination with CE.

## Ethics

**Ethics Committee Approval:** Approval for the study was granted by the Haydarpaşa Numune Training and Research Hospital Clinical Research Ethics Committee (decision no: HNEAH-KAEK 2016/98).

**Informed Consent:** All patients provided written consent prior to study initiation.

**Peer-review:** Externally and internally peer-reviewed.

## Authorship Contributions

Surgical and Medical Practices: F.B., D.G.K., N.M., M.H.T., Concept: F.B., D.G.K., N.M., M.H.T., Design: F.B., D.G.K., N.M., M.H.T., Data Collection or Processing: F.B., M.H.T., Analysis or Interpretation: F.B., D.G.K., N.M., M.H.T., Literature Search: F.B., D.G.K., Writing: F.B., D.G.K.

**Conflict of Interest:** No conflict of interest was declared by the authors.

**Financial Disclosure:** The authors declared that this study received no financial support.

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