Comparison of anterior single- and standard two-portal techniques in arthroscopic Bankart repair

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Traumatic recurrent anterior shoulder instability is a common issue, particularly in young active patients.[1] Repair of capsulolabral disruption has been considered the gold standard treatment for this condition. Although open Bankart repair is still used in the management of anterior shoulder instability, many arthroscopic surgical techniques have been developed over the past two decades, with increasing popularity providing a minimally invasive anatomical reconstruction with low operative complication rates.[2-5]

Arthroscopic Bankart repair (ABR) using one posterior and two anterior portals is the standard treatment for recurrent anterior shoulder instability, particularly in cases of isolated Bankart lesions or minimal loss of anterior glenoid rim.[6] However, creating two anterior working portals is challenging in some populations, particularly in patients with small shoulders.[7] Moreover, the standard anterior two-portal technique may lead to iatrogenic nerve

Objectives: This study aims to compare the outcomes of patients undergoing a single anterior portal or a standard two-portal arthroscopic procedure for Bankart repair.

Patients and methods: Between January 2015 and March 2018, a total of 71 consecutive patients (53 males and 18 females; mean age: 33.3±10.3 years; range, 17 to 56 years) who underwent arthroscopic Bankart repair with a minimum two-year follow-up period were included. The patients were divided into two groups according to the arthroscopic technique used: single anterior portal group (Group 1, n=32) and standard two-portal group (Group 2, n=39). Demographic and surgical characteristics of the patients were recorded. Pre- and postoperative clinical and functional outcomes were evaluated using the external rotation degree, as well as Visual Analog Scale (VAS), American Shoulder and Elbow Surgeons (ASES), University of California at Los Angeles (UCLA) Shoulder Rating Scale, Constant-Murley Score (CMS), Oxford Shoulder Instability Score (OSIS), and Rowe scores. The clinical and functional outcomes and revision rates were compared between the groups.

Results: The mean follow-up was 32.0±7.4 months in Group 1 and 38.0±13.4 months in Group 2 (p=0.222). Good-to-excellent postoperative functional and clinical outcomes were achieved in both groups at the final follow-up, compared to baseline (p<0.001 for all). No significant difference was observed in the postoperative outcomes including daily sports activity, VAS, ASES, UCLA Shoulder Rating Scale, CMS, OSIS, and Rowe scores, and external rotation restriction degrees between the groups (p=0.270, p=0.190, p=0.313, p=0.248, p=0.125, p=0.203, p=0.318, p=0.083, respectively). The operative time in Group 1 was significantly lower than that in Group 2 (60.3±8.3 vs. 71.4±7.2, respectively; p=0.001). Four patients (5.6%) experienced recurrent dislocation with no significant difference between the groups (p=0.622). No significant complications occurred in the peri- or postoperative period. Fifty-eight (81.7%) patients returned to their preoperative sports activity level. The mean time to return to sports was 7.2±1.7 months.

Conclusion: Good-to-excellent clinical and functional outcomes can be obtained after arthroscopic Bankart repair, regardless of the use of a single or two anterior working portals. However, the single-portal technique is associated with reduced the operative time, compared to two-portal technique.

Keywords: Arthroscopy, Bankart lesion, joint instability, shoulder dislocation, surgical procedures.
injuries and cannula breakage, as the cannulas have
to be close to each other in small shoulders.[5,7] Over
the past five years, several studies have described
the anterior single-portal technique for ABR.[5,8,9]
This technique is an effective ABR modality with
similar outcomes as that of the anterior two-portal
technique; however, few studies have compared the
two techniques.[5,8,10] In this study, we hypothesized
that the results of ABR might vary based on the
number of working portals used. We, therefore, aimed
to compare the clinical and functional outcomes and
revision rates of patients who underwent ABR using
two different techniques: a single anterior portal and
two standard portals.

**PATIENTS AND METHODS**

This multi-center, retrospective study was conducted
at Kahramanmaras Sutcu Imam University Medical
Faculty and Erciyes University Medical Faculty,
Department of Orthopaedics and Traumatology
between January 2015 and March 2018. A total of
119 patients who underwent ABR for traumatic
anterior shoulder instability were screened. A
surgical team with at least five years of experience
performing arthroscopic shoulder surgery
performed the surgeries for all patients. The data
of 71 consecutive patients (53 males and 18 females;
mean age: 33.3±10.3 years; range, 17 to 56 years)
who underwent ABR with a minimum follow-up
period of two years were retrospectively analyzed.
The patients were divided into two groups based
on the arthroscopic technique performed: Group
1, single anterior portal (n=32) and Group 2, two
standard portals (n=39). Inclusion criteria were
as follows: patients with primary arthroscopic
shoulder stabilization, anterior traumatic
instability with or without a concomitant superior
labrum anterior and posterior (SLAP)[8] lesion, and
Bankart or Bankart-like lesions (Perthes, anterior,
labroligamentous periosteal sleeve avulsion
[ALPSA]) lesion with/without a minimal Hill–Sachs
lesion requiring ABR; those willing to participate in
the study; those compliant with the postoperative
rehabilitation protocol; and those having a
minimum follow-up period of two years. Exclusion
criteria were as follows: having multidirectional
instability without an explicit traumatic episode
(n=4); the presence of preoperative arthropathy
according to the Samilson-Prieto classification[9]
which categorizes osteoarthritis (OA) in four
categories comprising (i) no OA, (ii) osteophytes
measuring <3 mm in greatest distance diameter,
(iii) osteophytes measuring between 3 and 7 mm in
greatest distance diameter and slight glenohumeral
joint irregularity, and (iv) osteophytes measuring
>7 mm in greatest distance diameter, narrowing
of the glenohumeral joint and sclerosis) (n=1);
preoperative rotator cuff injury (n=2); a history
of ipsilateral upper extremity surgery (n=3);
posterior instability repair (n=2); rheumatic disease
(n=2); neurological problems including cervical
myelopathy or any neuropathic disorder affecting
the ipsilateral or contralateral extremity (n=3);
previous surgery for the affected shoulder (n=6);
large Hill-Sachs lesions \(n=3\); anterior glenoid bone loss >17% of the glenoid surface area\(^{[10]}\) \(n=7\); and loss to follow-up \(n=15\). The study flow chart is shown in Figure 1. A written informed consent was obtained from each participant. The study protocol was approved by the Kahramanmaraş Sütçü İmam University Medical Faculty Ethics Committee (Date: 29.04.2020; No: 2020/15). The study was conducted in accordance with the principles of the Declaration of Helsinki.

**Surgical method**

All surgical procedures were performed arthroscopically in a beach chair position under general anesthesia. Diagnostic arthroscopy was conducted using a standard posterior portal. In Group 1, a second single anterior working portal was established through the rotator interval nearly 1 cm superior and 1 cm lateral to the coracoid notch (the best position and angle was determined to allow access to the anteroinferior capsule and glenoid labrum by means of an epidural needle with outside-in technique. The lateral aspect along the superior border of the subscapularis tendon in the appropriate trajectory was the entry point of the needle) (Figure 2a). In Group 2, standard anterosuperior and anteroinferior portals were created. The cannula was inserted in the anterior portal in both groups. After the anteroinferior capsulolabral complex was completely detached from the glenoid (Figure 2b, c), capsulolabral repair was performed through the anterior portal. For single-portal repair technique first a suture transferring system (ACCU-PASS Suture Shuttle, Smith and Nephew, UK) was used to pass the anterior capsulolabral complex taking a healthy bite of the labrum and capsules distally as possible, and further an ample amount of the non-degradable suture of suture shuttle was left in the joint (Figure 3a). The suture shuttle was, then, withdrawn from the anterior portal and a grasper was used to retrieve the transferring suture passed through the labrum (Figure 3b, c). A 1.7-mm, single-loaded suture anchor (SUTUREFIX, Smith and Nephew, London, UK) was placed through the anterior glenohumeral portal (Figure 4a). The first anchor was inserted inferiorly as close as the 5.30 position or 6.30 position in the right and left shoulders, respectively (Figure 4b). Subsequently, one end of the anchor threads was transferred and passed through the capsulolabral complex using the transferring suture (Figure 4c). The sutures were, then, simply tied to fix the capsulolabral complex on the anterior glenoid rim (Figure 4d). Depending on the extent of labrum detachment from the glenoid rim and the presence of concomitant SLAP lesion, two or more anchors were used to complete the repair of the capsulolabral complex. Threads of other additional anchors were transferred and subsequently fixated in a similar manner at an appropriate position. Bankart repair was performed similarly to ALPSA \(n=5\) or Perthes \(n=2\) lesions. Six patients had concomitant SLAP lesions: four patients with Type I SLAP lesion were treated with debridement and two patients with Type V SLAP lesion underwent SLAP repair in addition to ABR.

**Postoperative rehabilitation**

All patients remained in a shoulder immobilizer in slight abduction and neutral positions for six weeks postoperatively. For the first six weeks, pendulum exercises to flex the elbow and wrist...
were allowed; in the subsequent six weeks, active assisted/passive range of motion (ROM) exercises were initiated. After three months, strengthening and resistance exercises were initiated. Return to sports activities were permitted after four months, whereas contact sports were allowed six to nine months after surgery.

Assessments

The demographic characteristics of the patients were recorded from the prospectively collected data. A standard physical examination was conducted to record ROM pre- and postoperatively. For radiological evaluation, standard anteroposterior and axillary views were used. Preoperative axial magnetic resonance imaging (MRI) scans were used to identify the Hill-Sachs lesions and glenoid defects. The patients with glenoid defects on MRI underwent a three-dimensional computed tomography scan, and the defect size was measured by a physician not participating in the surgery according to the method described by Sugaya et al.\textsuperscript{[12]} Hill-Sachs lesions were intraoperatively evaluated and subjectively described as small, medium, or large. Deep lesions involving more than 15% of the humeral articular surface were defined as large lesions. For small- and medium-sized lesions, the remplissage procedure was performed following labral repair, if signs of instability persisted in arthroscopic control. Routine follow-up was performed at 1.5, three, six, and 12 months and annually, thereafter. Operative reports were assessed and pre- and postoperative (at the final follow-up) clinical and functional evaluations were performed. Pre- and postoperative pain and clinical and functional outcomes were measured using the following parameters: the external rotation degree was calculated by comparing the ROM of the patient’s affected side with that of the patient’s healthy side; the Visual Analog Scale (VAS), with scores ranging from 0 to 10 (0: no pain; 10: worst pain ever); the American Shoulder and Elbow Surgeons (ASES) Standardized Shoulder Assessment Form;\textsuperscript{[13]} the University of California at Los Angeles (UCLA)
### TABLE I

**Patient characteristics in single- and two-portal groups**

|                      | Total (n=71)                  | Single anterior portal (n=32) | Double anterior portals (n=39) |
|----------------------|------------------------------|------------------------------|-------------------------------|
|                      | n % Mean±SD Range            | n % Mean±SD Range            | n % Mean±SD Range             |
| Age at surgery (year)| 33.3±10.3 17-56              | 36.0±11.2 17-56              | 31.0±9.0 18-55                |
| Age at first dislocation (year) | 31.4±10.1 16-54           | 33.7±11.2 16-54              | 29.5±8.9 18-53                |
| Sex                  |                              |                              |                               |
| Female               | 18 25.9                      | 11 34.4                      | 7 17.9                       |
| Male                 | 53 74.6                      | 21 65.6                      | 32 82.1                      |
| Side                 |                              |                              |                               |
| Right                | 42 59.2                      | 17 53.1                      | 24 64.1                      |
| Left                 | 29 40.8                      | 15 46.9                      | 14 35.9                      |
| Side dominancy       |                              |                              |                               |
| No                   | 23 32.4                      | 9 28.1                       | 14 35.9                      |
| Yes                  | 48 67.6                      | 23 71.9                      | 25 64.1                      |
| BMI (kg/m²)          | 25.3±2.9                     | 25.9±2.8                     | 24.9±2.9                     |
| Contact sports       |                              |                              |                               |
| -                    | 32 45.1                      | 18 56.2                      | 14 35.9                      |
| +                    | 39 54.9                      | 14 43.8                      | 25 64.1                      |
| Smoking habit        |                              |                              |                               |
| -                    | 41 57.7                      | 18 56.2                      | 23 59.0                      |
| +                    | 30 42.3                      | 14 43.8                      | 16 41.0                      |
| Comorbidity*         |                              |                              |                               |
| -                    | 64 90.1                      | 27 84.4                      | 37 94.9                      |
| +                    | 7 9.9                       | 5 15.6                       | 2 5.1                       |

SD: Standard deviation; BMI: Body mass index; * Asthma, hypertension, diabetes, thyroid disease.

### TABLE II

**Surgical characteristics and injury patterns of all patients and single- and two-portal groups**

|                      | Total (n=71)                  | Single anterior portal (n=32) | Double anterior portals (n=39) |
|----------------------|------------------------------|------------------------------|-------------------------------|
|                      | n % Mean±SD Range            | n % Mean±SD Range            | n % Mean±SD Range             |
| Type of the injury   |                             |                              |                               |
| Training             | 31 43.7                      | 12 37.5                      | 19 48.7                      |
| Sports               | 35 49.3                      | 16 50.0                      | 19 48.7                      |
| Fall                 | 4 5.6                        | 3 9.4                        | 1 2.6                        |
| Accident             | 1 1.4                        | 1 3.1                        | 0 0.0                        |
| Number of instability episodes ≤5 | 30 42.3            | 12 37.5                      | 18 46.2                      |
|                       | 41 57.7                      | 20 62.5                      | 21 53.8                      |
| Time from first dislocation to operation (month) | 17.8±12.8 1-80            | 20.8±16.1 6-80              | 15.3±8.9 4-60                |
| No. of suture anchors | 2.6±0.5 2-4                   | 2.7±0.5 2-4                   | 2.6±0.5 2-4                   |
| Glenoid bone loss    |                             |                              |                               |
| (-)                  | 64 90.1                      | 30 93.7                      | 34 87.2                      |
| (+)                  | 7 9.9                        | 2 6.3                        | 5 12.8                        |
| Hill-Sachs lesion    |                             |                              |                               |
| (-)                  | 52 73.2                      | 23 71.9                      | 29 74.4                      |
| Small/medium         | 19 26.8                      | 9 28.1                       | 10 25.6                      |
| SLAP lesion          |                             |                              |                               |
| No                   | 65 91.5                      | 30 93.8                      | 35 89.7                      |
| Yes                  | 6 8.5                        | 2 6.3                        | 4 10.3                        |
| Perthes lesion       |                             |                              |                               |
| No                   | 69 97.2                      | 31 96.9                      | 38 97.4                      |
| Yes                  | 2 2.8                        | 1 3.1                        | 1 2.6                        |
| ALPSA lesion         |                             |                              |                               |
| No                   | 66 93.0                      | 30 93.8                      | 36 92.3                      |
| Yes                  | 5 7.0                        | 2 6.3                        | 3 7.7                        |
| Operative time (minutes) | 65.3±9.5 45-85            | 60.3±8.3 45-75              | 71.4±7.2 55-85                |

SD: Standard deviation; SLAP: Superior labrum anterior posterior; ALPSA: Anterior labroligamentous periosteal sleeve avulsion.
Shoulder Rating Scale,[14] the Constant-Murley Score (CMS),[15] the Oxford Shoulder Instability Score (OSIS),[16] and the Rowe score.[17] We did not perform routine postoperative MRI. To assess the postoperative recurrent instability, the patients were asked whether they experienced a redislocation after surgery and they needed help to have the shoulder reduced; those who answered “Yes” to both questions were considered to have a dislocation. The demographic data and functional and clinical outcome parameters were compared between the two groups.

### Statistical analysis

Statistical analysis was performed using the IBM SPSS for Windows version 22.0 software (IBM Corp., Armonk, NY, USA). Descriptive data were expressed in mean ± standard deviation (SD), median (min-max) or number and frequency, where applicable. The independent samples t-test and Mann-Whitney U test were used for analyzing independent quantitative data, Wilcoxon’s test for dependent quantitative data, and the chi-square test for independent qualitative data. The Fisher’s

| Evaluation method                        | Total (n=71) | Single anterior portal (n=32) | Double anterior portals (n=39) | p    |
|------------------------------------------|-------------|------------------------------|-------------------------------|------|
| Follow-up period (month)                 | 35.3±11.5   | 32.0±7.4                     | 38.0±13.4                     | 0.222|
| Length of hospital stay (day)            | 1.28±0.48   | 1.2±0.4                      | 1.41±0.56                     | 0.059|
| Daily sports activity (h)                |             |                              |                               | 0.400|
| Preoperative                             | 0.8±0.4     | 0.8±0.4                      | 0.9±0.4                       | 0.270|
| Postoperative                            | 1.5±0.7     | 1.4±0.7                      | 1.5±0.6                       |      |
| p                                        | <0.001      | <0.001                       | <0.001                        |      |
| VAS score                                |             |                              |                               | 0.302|
| Preoperative                             | 4.6±1.3     | 4.5±1.6                      | 4.8±1.0                       | 0.190|
| Postoperative                            | 1.0±1.0     | 0.9±1.0                      | 1.2±1.0                       |      |
| p                                        | <0.001      | <0.001                       | <0.001                        |      |
| ASES score                               |             |                              |                               | 0.186|
| Preoperative                             | 61.6±6.8    | 60.3±6.3                     | 62.7±7.1                      | 0.313|
| Postoperative                            | 87.4±10.8   | 88.4±11.3                    | 86.6±10.5                     |      |
| p                                        | <0.001      | <0.001                       | <0.001                        |      |
| UCLA score                               |             |                              |                               | 0.446|
| Preoperative                             | 18.7±3.5    | 19.3±2.9                     | 18.3±3.8                      | 0.248|
| Postoperative                            | 31.5±4.4    | 30.8±4.7                     | 32.2±4.1                      |      |
| p                                        | <0.001      | <0.001                       | <0.001                        |      |
| CM score                                 |             |                              |                               | 0.299|
| Preoperative                             | 56.6±7.1    | 55.9±6.5                     | 57.2±7.5                      | 0.125|
| Postoperative                            | 89.7±10.5   | 88.3±9.8                     | 90.9±11.1                     |      |
| p                                        | <0.001      | <0.001                       | <0.001                        |      |
| Oxford instability score                 |             |                              |                               | 0.310|
| Preoperative                             | 23.2±4.9    | 23.0±6.1                     | 23.4±3.8                      | 0.203|
| Postoperative                            | 43.0±6.1    | 43.6±6.2                     | 42.5±6.1                      |      |
| p                                        | <0.001      | <0.001                       | <0.001                        |      |
| Rowe score                               |             |                              |                               | 0.805|
| Preoperative                             | 51.1±9.4    | 50.6±8.2                     | 51.5±10.5                     | 0.318|
| Postoperative                            | 89.6±11.5   | 88.7±10.7                    | 90.3±12.3                     |      |
| p                                        | <0.001      | <0.001                       | <0.001                        |      |
| External rotation restriction            |             |                              |                               | 0.065|
| Preoperative                             | 16.6±6.2    | 18.2±5.3                     | 15.3±6.6                      | 0.083|
| Postoperative                            | 8.5±6.4     | 7.0±5.9                      | 9.7±6.5                       |      |
| p                                        | <0.001      | <0.001                       | <0.001                        |      |
| Recurrent instability                    |             |                              |                               | 0.622|
| No                                       | 67          | 94.4                         | 31                            | 96.6 |
| YES                                      | 4           | 5.6                          | 1                              | 3.1  |

SD: Standard deviation; VAS: Visual Analog Scale; ASES: American Shoulder and Elbow Surgeons Standardized Shoulder Assessment Form; UCLA: University at California at Los Angeles Shoulder Rating Scale; CM: Constant-Murley. Bolded p values indicate statistically significant differences between groups (p<0.05).
exact test was used, when the chi-square test requirements were not met. A p value of <0.05 was considered statistically significant.

RESULTS
The mean follow-up period was 35.3±11.5 months (range, 24-60). Of the 71 patients, 32 (45.1%) underwent ABR with the single anterior portal technique (Group 1) and 39 (54.9%) underwent ABR with the two-portal technique (Group 2). No significant differences were observed between the groups in terms of patients’ characteristics and demographic data (Table I). Additionally, injury patterns and surgical characteristics, except for operative time, were similar between the groups (Table II). Operative time was significantly lower in Group 1 than in Group 2 (60.3±8.3 min vs. 71.4±7.2 min; p=0.000). In both groups, patients demonstrated a significant pre- to postoperative improvement in clinical and functional outcome scores (p<0.001); however, this improvement was not significantly different between the groups. Furthermore, the length of hospital stay and recurrent instability rates were not statistically significantly different between the groups (p=0.05). The clinical and functional outcome scores are summarized in Table III. The overall recurrent instability rate at the final follow-up was 5.6% (4/71). One patient (3.1%) in Group 1 and three patients (7.7%) in Group 2 with recurrence underwent revision ABR (p=0.622). After the revision surgery, no redislocations occurred during the remaining follow-up period. Twenty-six patients (81.3%) in Group 1 and 32 patients (82.1%) in Group 2 returned to their preoperative sports activity level (p=0.931). The mean time to return to preoperative sports activity was 7.5±1.7 months in Groups 1 and 2, respectively (p=0.079). No patients developed superficial or deep infections, and no significant complications occurred in the peri- or postoperative period.

DISCUSSION
The main finding of this study is that the anterior single-portal technique is as effective, reliable, and reproducible as the standard two-portal technique, with similar good-to-excellent clinical and functional outcomes. Furthermore, compared to the standard two-portal technique, the anterior single-portal technique was associated with short operative time and low rate of neurovascular injuries.

Over the past two decades, several arthroscopic techniques have been developed, particularly for shoulder surgery [2-5,11,18]. The advantages of ABR are as follows: less postoperative pain, less loss of motion, shorter length of hospital stay, lower morbidity rates, shorter operative time, and better cosmetic appearance [20]. Previously, the standard anterior two-portal technique was used in ABR surgery; however, the single anterior portal technique, which is less invasive, has been introduced and increasingly used over the past five years [2,5,11,18-20]. This present study evaluated the effectiveness of single anterior portal technique as an up-to-date treatment method in ABR. A detailed evaluation was performed using clinical and functional outcome measures and recurrence rates. Compared to the standard two anterior portal, single-portal technique showed similar outcomes concurrent with the literature [5,11,20]. Moreover, the results of the current study showed a similar recurrence rate with a single anterior portal, compared to the standard two-portal technique after at least two years.

Few studies have compared the results of the single anterior portal and standard two-portal techniques [9,11,20]. Cicek et al. [20] compared ABR with the traditional two-portal technique, and ABR with the single anterior portal was associated with less postoperative pain (quantity of analgesics, 200 mg vs. 300 mg, respectively; p<0.001), shorter surgical learning curve, and lower costs (5.7% less than the two-portal technique). They also reported a higher median operative time for the standard two-portal technique (53.5 min vs. 35 min, respectively; p<0.001). However, the authors found no difference in terms of clinical results between these two techniques and emphasized that, compared to the anterior two-portal technique, the single anterior portal technique prevented the instability of the glenohumeral joint due to less damage in the rotator interval. Consistent with the results obtained by Cicek et al. [20], Ghai et al. [5] reported similar functional outcomes, Rowe and OSIS scores, and Tegner activity level in a comparison between the anterior single-portal and anterior two-portal arthroscopic techniques. However, the operative time was significantly lower in the single anterior portal group (46.35 min vs. 68.52 min, respectively; p<0.001), indicating that this technique was an effective treatment modality, with similar outcomes to those of the double technique. Armandgil et al. [21] reported that ABR performed using the single anterior portal technique provided comparable clinical outcomes in terms of postoperative shoulder movements with the two-portal repair technique and emphasized the importance of appropriate patient selection rather than the number of portals. Furthermore, they suggested that, compared to the
two-portal technique, the single anterior portal technique reduced the operative time and was less invasive. A recent prospective study evaluating the effectiveness and safety of ABR revealed that, compared to two anterior portals, the single-portal technique had a shorter operative time and lower costs and that there were no significant differences between the groups in terms of clinical or functional scores, quality of life, or patient satisfaction.[6] In the current study, we found a significant postoperative improvement in clinical and functional outcome scores compared to baseline, regardless of the number of anterior portals; however, in line with the previous findings, the single anterior portal technique had lower operative times.

Postoperative recurrent instability is the most common and undesired complication associated with ABR.[11] Glenoid and humeral bone loss, younger age, participation in contact sports, male sex, a higher number of preoperative dislocations, and bilateral instability have been reported as the leading causes of recurrent instability.[21-26] However, similar outcomes were observed in the study of Cicek et al.[20] that compared the recurrence rate with the number of anterior portals used. In this current study, one (3.1%) patient in Group 1 and three (7.7%) patients in Group 2 had recurrent instability, and there were no significant differences between the groups. The overall dislocation rate in the present study was 5.6%, which is consistent with that reported in the literature.[5,20]

In their study, Cicek et al.[20] reported that the mean length of hospital stay for patients in the traditional two-portal group was significantly longer than that for patients in the single-portal group (1.5 days vs. 1 day, respectively; p<0.001). However, in our patient cohort, no significant difference was observed between the two groups in terms of the length of hospital stay.

Various complications may occur during ABR, particularly during portal opening, anchor placement, and labral repair. The creation of two anterior working portals can be challenging in some populations, particularly in those with small shoulders,[7] iatrogenic nerve injuries and cannula breakage have been reported to occur in the standard anterior two-portal technique due to the overcrowding of cannulas in small shoulders and the difficulty associated with working in these shoulders.[5,7] In our study, no significant complications occurred in the peri- or intraoperative period in any groups.

Studies evaluating the clinical and functional results of the single anterior portal and standard two-portal techniques for ABR are limited. Therefore, the findings of our study are valuable, supporting the existing literature by demonstrating and proving that the single anterior portal technique is safe and can reduce the operative time.

Nonetheless, this study has some limitations. First, it has a retrospective design; however, we prospectively collected the data of patients who were not lost to follow-up to obtain more accurate results. In addition, there may be a selection bias regarding which patients underwent ABR with single and two portals. Second, this study had a short follow-up period, implying that mid- or long-term functional outcomes and the recurrence rate might differ. Third, the number of patients is relatively low. Finally, we did not analyze the cost; however, in the anterior two-portal techniques, additional cannulas are used, which increases the cost of this technique compared to the single-portal technique. Further, prospective randomized-controlled studies with longer follow-up periods are warranted to obtain more accurate results regarding the effect of the number of anterior portals used in ABR on outcomes.

In conclusion, good-to-excellent clinical and functional outcomes can be obtained after ABR, regardless of the use of a single or two anterior working portals. The single anterior portal technique is as effective, reliable, and reproducible as the standard two-portal technique. In addition, the single-portal technique can reduce the operative time and the possible rate of neurovascular injury that may occur during the opening of the second anterior portal.

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