

## ORIGINAL ARTICLE

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**Comperative outcomes of the patients undergoing  
percutaneous and open trigger finger release****●Duran Toprak<sup>1</sup>, ●Fatih Dogar<sup>1</sup>, ●Burak Kusu<sup>1</sup>, ●Ali Aydın Karadeniz<sup>1</sup>, ●Okkes Bilal<sup>1</sup>**<sup>1</sup>Kahramanmaraş Sutcu Imam University, Faculty of Medicine, Department of Orthopaedic and Traumatology, Kahramanmaraş, Turkey

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**Abstract**

The aim of this study was to compare patients who underwent open or percutaneous trigger finger release in terms of clinical outcomes, time to return to activities, and recurrence. The records of patients who underwent percutaneous and open trigger finger release between 2012 and 2018 at two different hospitals in the same city were retrospectively reviewed. The patients were divided into two groups: 33 patients who underwent percutaneous trigger finger release (Group PR) and 48 patients who underwent open release of A1 pulley (Group OR). The clinical classification of cases was done according to the Quinell classification. The functional outcomes of the patients were evaluated according to the Quick DASH scale. The mean age of the patients was  $55.95 \pm 11.73$  (27–82) years; 71.6% (n = 58) were female. The left side was involved in 56.8% (n = 46) patients, and 81 patients underwent percutaneous or open trigger finger release with a mean follow-up duration of  $37.40 \pm 16.22$  (12–72) months. The time to start daily activities was shorter in Group PR than in Group OR, and the difference was statistically significant ( $p < 0.001$ ). A comparison of the upper extremity functional scores between the two groups revealed no statistically significant difference (PR;  $15.21 \pm 6.17$ , PO;  $12.99 \pm 6.89$ ,  $p = 0.142$ ). Although the rate of complications was higher in Group OR, there was no statistically significant difference between the two groups (PR; 12.12%, PO: 20.83%,  $p = 0.217$ ). Percutaneous trigger finger release can be preferred in adult trigger finger surgery due to increased risks regarding wound healing and infections associated with advanced age, presence of diabetes and inflammatory arthritis, and the expectation of rapid return to daily activities.

**Keywords:** Trigger finger, percutaneous release, open release**Introduction**

Trigger finger is a common clinic condition characterized by catching, locking, and a decrease in grip strength during finger movements, resulting from incompatibility between the flexor tendons and A1 pulley [1]. Trigger finger, which is one of the most common causes of pain and disability in the hand, affects 2.6% of the people in the general population [1, 2]. It often occurs on the thumb, with a 4–6 times high incidence in females than in males [3].

Trigger finger can be seen at any age, but it is observed more often in patients aged  $\geq 50$  years and in the dominant hand [4]. The main etiological factor is chronic repetitive friction between the A1 pulley and the flexor tendon. It has been reported that some connective tissue diseases occur more often in patients with diabetes and inflammatory arthritis [5–7].

The Quinell grading system is used to clinically assess the severity of trigger finger [8]. Mild cases with early disease onset can be treated by conservative methods (activity modification, splinting, nonsteroidal anti-inflammatory drugs, and local steroid injections) [1, 4, 7].

In delayed cases that do not benefit from conservative methods, open or percutaneous A1 pulley release is performed [4, 8, 9].

Various conservative and surgical methods have been described for the treatment of trigger finger, and successful outcomes have been reported. The aim of this study was to compare patients who underwent open or percutaneous trigger finger release in terms of clinical outcomes, and recurrence. As far as we can evaluate, it is the first study in the literature to compare the two different treatment options of the trigger finger in terms of time to return to daily activities.

**Material and Methods**

Patients who underwent percutaneous and open trigger finger release between 2012 and 2018 at two different hospitals in the same city were retrospectively reviewed after obtaining approval of the local ethics committee (Session: 2019/02, Date:06.02.2019, Decision no:10).

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We included patients aged >18 years who underwent percutaneous or open trigger finger release, did not have neurological deficit, and followed up for  $\geq 12$  months. Children under 18 years of age, patients treated for recurrence, and patients with incisions or scars in the same area before treatment were excluded because they had the potential of influencing the outcomes of the treatment.

A total of 81 patients who met the criteria were included, and their files were retrospectively reviewed. The patients were divided into two groups: 33 patients who underwent percutaneous trigger finger release (Group PR) and 48 patients who underwent open release of A1 pulley (Group OR).

Then, the data on the patient charts including age, sex, direction, dominant hand, patient complaint, etiological cause, course of treatment, follow-up period, time of return to daily activities, and complications were evaluated. The symptoms of the patients were classified according to the Quinnell grading system [8]. The functional outcomes of the patients were evaluated using the QuickDASH (disabilities of the arm shoulder and hand) scale [10].

All patients in Group OR underwent A1 pulley release under local anesthesia, and these patients received oral antibiotics and analgesics for 5 days in the postoperative period. The patients in Group PR received an injection of a mixture of betamethasone sodium phosphate 2 mg/1mL (Diprospan vial) and 1 ml of 2% xylocaine in a proximal-to-distal direction using a 21-gauge needle through a green-tip injector inserted into the flexor tendon sheath at an angle of 45° over the A1 pulley, while the metacarpophalangeal joint of the finger was in flexion. Then, the pulley was released in a longitudinal direction by serial extension and flexion movements.

### Statistical Analysis

A IBM SPSS 20.0 (IBM, Armonk, NY, USA) software package was used to analyze the data obtained from patient files. Categorical variables were presented as frequency (n, %), and quantitative data were expressed mean and standard deviation. A Kolmogorov–Smirnov test was used to assess the fitness of the data for normal distribution. A chi-square ( $\chi^2$ ) test was used to compare categorical variables of the two independent groups, whereas an independent samples t-test was used to evaluate quantitative data. A p-value of <0.05 was considered statistically significant.

### Results

The mean age of the patients was  $55.95 \pm 11.73$  (27–82) years; 71.6% (n = 58) were female. The left side was involved in 56.8% (n = 46) patients, and there were 81 patients who underwent percutaneous or open trigger finger release with a mean follow-up duration of  $37.40 \pm 16.22$  (12–72) months. The thumb was the most commonly affected finger (n = 43, 53.1%), whereas the most common etiological factor was diabetes that was found in 29.6% (n = 24) of the cases. The most common complications were recurrence (n = 7, 8.6%), delayed wound healing (n = 4), wound site infection (n = 2), and tendon rupture (n = 1). Demographic data of the cases by groups are presented in Table 1.

A comparison between the two groups did not reveal a significant

difference in terms of age, sex, rate of dominant hand involvement, and Quinnell grade. The time to start daily activities was shorter in Group PR than in Group OR, and the difference was statistically significant (PR:  $1.60 \pm 1.45$ ; PO:  $16.31 \pm 7.34$ ;  $p < 0.001$ ). A comparison between the two groups in terms of upper extremity functional scores revealed no statistically significant difference (PR:  $15.21 \pm 6.17$ ; PO:  $12.99 \pm 6.89$ ;  $p = 0.142$ ). Although the incidence of complications was high in Group OR, there was no statistically significant difference between the two groups (PR: 12.12%, PO: 20.83%,  $p = 0.217$ ). The comparison between the two groups is presented in Table 2.

Although there was a statistically significant difference between the two groups in terms of the duration of follow-up, this was not considered to have an influence on the outcomes ( $p = 0.048$ ).

**Table 1.** Distribution of Demographic Data by Groups

|                                                   |                         | n=81 (%)                     |
|---------------------------------------------------|-------------------------|------------------------------|
| <b>Age (mean<math>\pm</math>SD (min-max) year</b> |                         | 55.95 $\pm$ 11.13 (27-82)    |
| <b>Sex</b>                                        | Female                  | 58 (%71.6)                   |
|                                                   | Male                    | 23 (%28.4)                   |
| <b>Side</b>                                       | Dominant                | 39 (%48.1)                   |
|                                                   | Non-Dominant            | 42 (%51.9)                   |
| <b>Etiology</b>                                   | No                      | 38 (%46.9)                   |
|                                                   | Diabetes Mellitus       | 24 (%29.6)                   |
|                                                   | Inflammatory Arthritis  | 9 (%11.1)                    |
|                                                   | Heavy Duty              | 2 (%2.5)                     |
|                                                   | Hand crafted            | 8 (%9.9)                     |
| <b>Affected Finger</b>                            | Thumb                   | 43 (%53.1)                   |
|                                                   | Index Finger            | 11 (%13.6)                   |
|                                                   | Middle Finger           | 16 (%19.8)                   |
|                                                   | Ring Finger             | 10 (%12.3)                   |
|                                                   | Little Finger           | 1 (%1.2)                     |
| <b>Complaint</b>                                  | Trigerring              | 29 (%35.8)                   |
|                                                   | Pain                    | 36 (%44.4)                   |
|                                                   | Swelling                | 5 (%6.2)                     |
|                                                   | Loss of Labor           | 11 (%13.6)                   |
| <b>Complications</b>                              | No                      | 67 (%82.7)                   |
|                                                   | Delay in Wound Healing  | 4 (%4.9)                     |
|                                                   | Surgical Area Infection | 2 (%2.5)                     |
|                                                   | Recurrence              | 7 (%8.6)                     |
|                                                   | Tendon Rupture          | 1 (1.2)                      |
| <b>QDASH Scoring</b>                              |                         | 13.90 $\pm$ 6.66 (2.30-29.5) |
| <b>Following Time (mounth)</b>                    |                         | 37.40 $\pm$ 16.22 (12-72)    |

Abbreviation: QDASH; Quick Disabilities of Arm, Shoulder & Hand

**Table 2.** Comparative Outcomes of The Groups

|                                                                 |                            | Group PR<br>(n=33) | Group OR<br>(n=48) | p<br>Value   |
|-----------------------------------------------------------------|----------------------------|--------------------|--------------------|--------------|
| <b>Age (mean±SD (min-max) year</b>                              |                            | 57.81±11.32        | 54.66±10.93        | 0.213        |
| <b>Sex</b>                                                      | Female                     | 24                 | 34                 | 0.853        |
|                                                                 | Male                       | 9                  | 14                 |              |
| <b>Side</b>                                                     | Dominant                   | 18                 | 21                 | 0.339        |
|                                                                 | Non-Dominant               | 15                 | 27                 |              |
| <b>Quinnell<br/>Classification</b>                              | Class 2                    | 2                  | 2                  | 0.405        |
|                                                                 | Class 3                    | 10                 | 22                 |              |
|                                                                 | Class 4                    | 21                 | 23                 |              |
|                                                                 | Class 5                    | 0                  | 1                  |              |
| <b>Time to Return to Daily Activities<br/>(day)</b>             |                            | 1.60±1.45          | C                  | <b>0.000</b> |
| <b>QDASH Scoring (Mean ± SD)</b>                                |                            | 15.21±6.17         | 12.99±6.89         | 0.142        |
| <b>Complications</b>                                            | Delay in Wound<br>Healing  | 0                  | 4                  | 0.217        |
|                                                                 | Surgical Area<br>Enfection | 0                  | 2                  |              |
|                                                                 | Recurrence                 | 4                  | 3                  |              |
|                                                                 | Tendon Rupture             | 0                  | 1                  |              |
| <b>Mean Follow-up Time (month)</b>                              |                            | 33.12±14.64        | 40.35±16.73        |              |
| Abbreviation: QDASH; Quick Disabilities of Arm, Shoulder & Hand |                            |                    |                    |              |

## Discussion and Conclusions

Trigger finger occurs in individuals aged >50 years, often in those with diabetes and autoimmune disorders (such as rheumatoid arthritis) [4-7, 11]. Aging, diabetes, and inflammatory arthritis are known to affect wound healing adversely by reducing cell proliferation, revascularization, growth factors, and collagen deposition and remodeling in the wound site [12-14].

In this study, there was no delay in wound healing and no wound site infection in Group PR. PR of the trigger finger is superior to open release in terms of their effects on wound healing. Thus, percutaneous trigger finger release should be preferred in the treatment of elderly patients with diabetes and inflammatory arthritis who are at risk of a delay in wound healing. Although the complication rate was lower in Group PR than in Group OR, the difference was not statistically significant. This finding is caused by the small sample size. Percutaneous release is superior to open release in terms of complications. A success rate of 45% was reported in trigger finger surgery after a single corticosteroid injection, whereas another study reported successful outcomes in 39% of the patients who did not respond to the first two injections. The patients are advised to undergo surgery following two unsuccessful steroid injections. Most surgeons prefer

percutaneous or open release techniques [15-17].

In trigger finger surgery, open release of A1 pulley is a reliable method yielding a success rate of 92%–100% [18-21]. The efforts toward reducing the size of incision during open release surgery resulted in the introduction of percutaneous release technique, which was described for the first time in 1958 [22]. Although it is reported to be safe in most studies, the reported success rate is 91%–100% [23-25]. In our study, the success and recurrence rate of both treatment methods are consistent with those in the literature. Short follow-up periods may be the cause of high success rates reported in some studies. The studies on patients with and without diabetes suggest that percutaneous release with concurrent corticosteroid injection is more reliable and successful than steroid injections alone [26, 27]. However, in the first one-month period, it has been reported that pain complaints were less common only in the steroid group. A comparison between patients undergoing open and percutaneous release revealed no difference between the two groups in terms of treatment success and complications and that both treatments can be performed successfully [4, 23, 28]. In Group PR, the mean duration of pain and the mean time to recovery of motor functions were shown to be shorter, showing faster healing, in Group PR than in Group OR [23].

In our study, the success of treatment, QDASH scores, and complication rates were similar between the two groups, but the time of return to daily activities was shorter in the percutaneous release group. Percutaneous release technique must be preferred in patients who expect to return to daily activities in the early period. The involvement of two different surgeons in the treatment of patients and the retrospective study design are important limitations of the study. There is a need for randomized and prospective studies on a larger series of patients. In conclusion, percutaneous trigger finger release can be preferred in adult trigger finger surgery due to increased risks regarding wound healing and infections associated with advanced age, presence of diabetes and inflammatory arthritis, and the expectation of rapid return to daily activities.

### Scientific Responsibility Statement

*The authors declare that they are responsible for the article's scientific content, including study design, data collection, analysis and interpretation, writing, some of the main line, or all of the preparation and scientific review of the contents and approval of the final version of the article.*

### Human and Animal Rights Statement

*All procedures performed in this study were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards.*

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### Conflict of interests

*The authors declare that they have no competing interests.*

### Financial Disclosure

*There are no financial supports.*

### Ethical approval

*This study was approved by K.S.Ü Faculty of Medicine Clinical Research Ethics Committee. (Session: 2019/02, date: 06.02.2019, decision no: 10)*

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