

Ultrasound-guided percutaneous carpal tunnel release in patients on hemodialysis: early experiences and clinical outcomes

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Purpose: The purpose of the current study was to evaluate the effectiveness of ultrasound-guided percutaneous carpal tunnel release in hemodialysis patients with carpal tunnel syndrome.

Methods: From February 2009 to April 2013, a prospective review of 113 consecutive cases of ultrasound-guided percutaneous carpal tunnel release was carried out in 84 hemodialysis patients. Results were analyzed by clinical subjective scale, two self-administered questionnaires, and functional evaluations at seven time points (1 week and 1, 3, 6, 12, 18, and 24 months).

Results: Satisfactory symptom improvement in patients was 82%, 80%, 86%, 89%, 90%, 91%, and 90% at 1 week and 1, 3, 6, 12, 18, and 24 months postoperatively, respectively. Moderate pain was suffered in 11.5% of patients within 1 week, 8.8% within 1 month, 2.7% within 3 months, and none after 12 months postoperatively. Static two-point discrimination and Semmes-Weinstein monofilament examinations presented significant improvements after 1 week and 1 month postoperatively and with time. Postoperative grip power demonstrated recovery and a significant increase after 3 and 6 months postoperatively. Three-jaw chuck-pinch strength showed significant increase after 1 month postoperatively. There were no operative complications.

Conclusion: Ultrasound-guided percutaneous carpal tunnel release is an effective and safe procedure in hemodialysis patients with carpal tunnel syndrome. The advantages include a less invasive procedure, no tourniquet needed, only limited infiltration anesthesia, minimal soft-tissue exploration, and relatively short operation time. Our data suggest this technique can reliably relieve clinical symptoms, with early restoration of grip and pinch strength.

Keywords: carpal tunnel, hemodialysis, ultrasound, percutaneous release

Introduction

Carpal tunnel syndrome (CTS) is a frequent complication of patients undergoing hemodialysis. It often causes pain, numbness, tingling, or burning sensation in the median nerve distribution, hand weakness, and reduced quality of life.^{1,2,3} CTS in hemodialysis patients is recognized as an unremitting course, and surgical release procedures are often required after a long period of hemodialysis.^{4,5}

Traditionally, open carpal tunnel release (OCTR) is used in CTS patients on long-term hemodialysis, and has the advantage of direct and complete release of pathogens.² Friable skin, arteriovenous fistula (AVF), and coagulopathy make OCTR and extended OCTR prone to create more morbidities in hemodialysis patients than in the general population.⁶ Endoscopic CTR (ECTR) has been applied from idiopathic CTS to hemodialysis-related CTS. Although ECTR has superior short-term

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results, final effectiveness is the same between idiopathic and long-term hemodialysis patients at 3–6 months postoperatively.⁷ However, the potential risk of incomplete release and injury of adjacent neurovascular structures restrict application of ECTR in idiopathic and long-term hemodialysis patients.^{3,8}

Ultrasound is a safe image modality and applied in many medical procedures. Advances in high-frequency musculoskeletal ultrasound have made related therapeutic applications in many musculoskeletal pathogens possible.⁹ The structure of the carpal tunnel, including the transverse carpal ligament (TCL), medial nerve, flexor tendons, proximal palmar arch, and ulnar artery, are easily visible using high-frequency ultrasound. Many studies have recently reported ultrasound application in CTR, from cadaver studies to clinical cases.^{10–14} Chern et al presented their clinical experiences and outcomes of ultrasound-guided percutaneous CTR (PCTR). This has proved to be a effective and safe procedure for idiopathic CTS.¹¹ Nonetheless, there have been few studies of the application of ultrasound-guided PCTR in hemodialysis patients. The purpose of the current work is thus to present the results of ultrasound-guided PCTR in a consecutive series of hemodialysis patients with CTS.

Methods

Patient population

The study was reviewed and approved by the Ethics Committee of National Cheng Kung University Hospital. All participants were aware of the objective of this surgery and signed informed-consent documents. This study was conducted in accordance with the Declaration of Helsinki.

From February 2009 to April 2013, we performed 113 ultrasound-guided PCTRs in 84 hemodialysis patients with CTS. Initial data collected were age, sex, duration of disease, duration of hemodialysis, involved hand, symptom frequency, provocative factors, and comorbidity. CTS was diagnosed in accordance with clinical symptoms of pain, paresthesia and numbness in median nerve distribution, clumsiness, weakness of the hand, positive stimulation tests of Tinel's sign, flick sign, Phalen's test, and positive electrophysiological test results. The indication for surgery was any hemodialysis patient with CTS who had failed on conservative treatment >3 months or clinical evidence of thenar muscle atrophy or weakness. We excluded patients with acute CTS, recurrent CTS, inflammatory arthropathy, and documented peripheral neuropathy from the study.

Operative procedures

All patients had hemodialysis the day before surgery. They all received the same operative procedure: ultrasound guided PCTR under local anesthesia without use of a tourniquet. Firstly, flexor tendons, the TCL, median nerve, and bone landmarks were localized by an ultrasound scanner (5–10 MHz; SonoSite, Bothell, WA, USA) and safe zones determined. Secondly, insertion of the 21-gauge needle for 1% lidocaine infiltration and then creation of an inlet with an 18-gauge needle for the probe was done. Thirdly, the custom-made blunt probe was inserted subcutaneously to create the correct tract, and the hook knife (Ectra; Smith and Nephew, Andover, MA, USA) was introduced to cut the TCL backwardly. After cutting, the probe was inserted to evaluate adequate release. Real-time ultrasound was used to monitor all these procedures¹¹ (Figure 1).

Postoperative care

The small puncture wound of the inlet was covered with a dry dressing without suturing postoperatively. The dressing was able to be removed in the next day. These patients were permitted to engage in all their daily activities and works as tolerated.

Assessing results and outcomes

Arrangements were made for patients to return to the hospital 1 week and 1, 3, 6, 12, 18, and 24 months after the operation. Postoperative assessments were performed by an experienced orthopedic surgeon and a hand therapist. The assessment consisted of an interview with self-administered questionnaires and a clinical examination, including the same strength and sensitivity measurements.

Postoperatively, subjective improvement in symptoms was rated from three grades (poor, fair, and good). Pre- and postoperatively, patients completed the Carpal Tunnel Syndrome Symptom Severity Score (CTS-SSS) and Carpal Tunnel Syndrome Functional Status Score (CTS-FSS) questionnaires, which scaled the patients' symptoms from 1 to 5, with 5 representing the greatest symptoms or the most functional difficulty.¹⁵ Mean scores were recorded.

Digital sensibility was assessed by static two-point discrimination and Semmes–Weinstein monofilament tests over the proximal and distal phalanges of the thumb, middle and distal phalanges of the index, middle, and ring fingers, and the little finger for comparison. The grip strength of involved and uninvolved hands was assessed by all five settings of a hand dynamometer

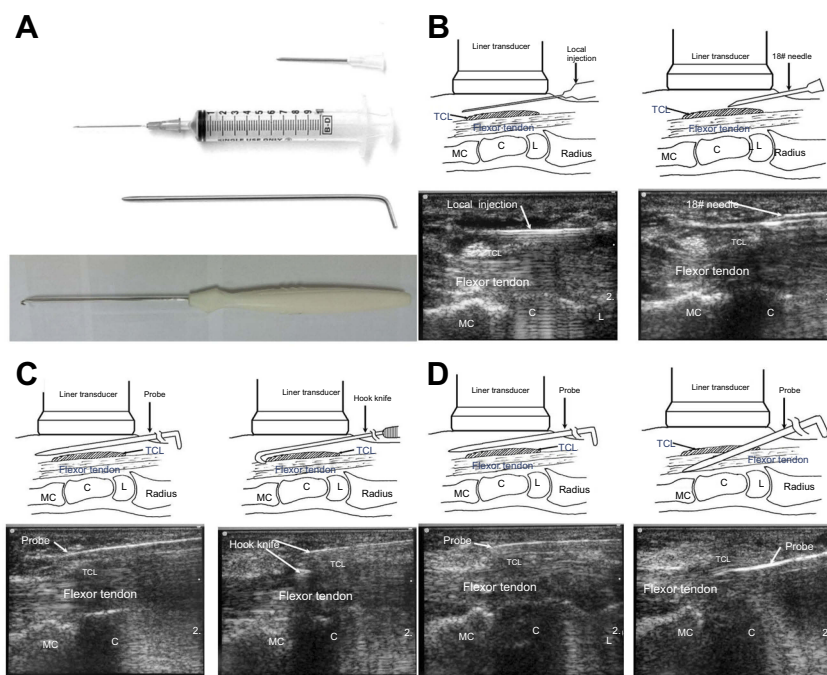


Figure 1 (A) Instruments for ultrasound-guided percutaneous carpal tunnel release: 18-gauge needle, 10 mL syringe with 21-gauge needle, custom-made blunt probe, custom-made hook knife. (B) Insertion of the 21-gauge needle for lidocaine infiltration and then to create an inlet with an 18-gauge needle. (C) The custom-made blunt probe was inserted subcutaneously to create the correct tract, and the hook knife was introduced to cut the TCL backwardly. (D) The probe was inserted to evaluate adequate release.

Abbreviations: C, capitates; L, lunate; MC, metacarpal; TCL, transverse carpal ligament.

(Asimov Engineering, LA, CA, USA). We used the maximum grip strength for statistical analysis. Three-jaw chuck-pinch strengths were measured using a pinch meter (Therapeutic Instruments, Clifton, NJ, USA).

Statistical analysis

All measurements are expressed as means \pm SD. GraphPad (La Jolla, CA, USA) Prism 5.0 software was used to analyze the different graded areas. One-way ANOVA was used to analyze the measurements, and post hoc pair-wise comparisons were performed in cases of significant difference. Statistical significance was set at $P < 0.05$.

Results

This prospective study consisted of 113 consecutive primary ultrasound-guided PCTR cases performed in 84 hemodialysis patients, including 56 women (67%) and 28 men (33%); age range 39–75 years, average age 60.9 ± 9.17 years) during February 2009 to April 2013. The duration of patients on hemodialysis averaged 7.53 ± 3.23 (1.0–15) years. The duration of symptoms averaged 24.3 ± 29.8 (0.5–120) months. Diagnosis was recorded as bilateral CTS in 34 patients (40%, 21 women and 13 men). CTS involved the AVF hand in 75 (89%) patients. Before surgery, 42 (37%) hands

had thenar muscle atrophy, 28 (25%) hands suffered from symptoms all day, and 64 (57%) frequently. A total of 64 (57%) hands had symptoms that intensified during sleep, and 49 (43%) with repetitive movements of the wrist. In sum, 49 (58%) hands had positive Phalen's tests, 63 (75%) positive Tinel's sign, and eight (10%) neither of these. Comorbidities included 61 (73%) patients with trigger finger, 56 (66%) with hypertension, 34 (41%) with diabetes, and 12 (14%) with hyperlipidemia. The 34 recruited diabetic patients had no concomitant evidence of peripheral neuropathy. All patients tolerated the procedure, and division of the TCL was confirmed by ultrasound postoperatively. Complete preoperative data were accessible for all cases. All patients underwent complete follow-up by reexamination or telephone.

Symptom relief, postoperative pain, and tenderness

Subjective sensory disturbances, such as paresthesia and hypoesthesia, improved immediately postoperatively, and such sensory disturbances had dissolved in 82%, 80%, 86%, 89%, 90%, 91%, and 90% of cases at 1 week and 1, 3, 6, 12, 18, and 24 months after the operation, respectively. Evaluation of pain at each follow-up is described in Table 1. Average CTS-SSS was 3.49 and CTS-FSS 3.73 preoperatively. Both

Table 1 Survey scale of symptom improvement after ultrasound-guided percutaneous release

	1 week	1 month	3 months	6 months	12 months	18 months	24 months
Poor	7%	8%	5%	3%	1%	1%	1%
Fair	11%	12%	9%	8%	9%	8%	9%
Good	82%	80%	86%	89%	90%	91%	90%

scores improved continuously throughout the follow-up period. Moderate numbness occurred in 32.7% of patients within 1 week, 21.2% within 1 month, 7% within 3 months, and no patients 18 months after the operation. Moderate pain occurred in 11.5% of patients within 1 week, 8.8% within 1 month, 2.7% within three months, and no patients 12 months after the operation. Ultimately, average CTS-SSS improved to 1.18 and CTS-FSS to 1.00 at 24 months postoperatively (Table 2).

Sensory testing and strength

A total of 91 (91 of 113, 70.5%) hands revealed >6 mm in the two-point discrimination evaluation preoperatively. Postoperative scores (two-point discrimination >6 mm) decreased to 46.7% at 1 week and 4.9% at 24 months. Scores in both static two-point discrimination and Semmes-Weinstein monofilament examinations demonstrated significant improvements after 1 week and 1 month postoperatively and with time. Grip and three-jaw chuck-pinch strength scores averaged 15.17 ± 14.3 and 4.87 ± 2.1 kg preoperatively. Grip power had diminished at 1 week after operation, and then recovered and had significantly increased at 3- and 6-month follow-up examinations. Three-jaw chuck-pinch strength showed a significant increase after 1 month postoperatively. Grip and three-jaw chuck-pinch strength scores had increased to 17.11 ± 10.6 and 9.98 ± 3.6 kg at the final follow-up. All these data imply the effect of the operations (Table 3).

Wound healing and complications

In all patients, the wound healed uneventfully within a few days and became invisible without other cosmetic problems. No patients had neurovascular structure/tendon injury or superficial/deep tendon-sheath infection.

Final outcomes

All 113 hands in this prospective study were assessed by clinical examination or telephone review based on the CTS-SSS and CTS-FSS. The satisfactory level of good symptom improvement of the patients was 90% at the final 24-month follow-up. No patient suffered from moderate hand or wrist pain after 1 year postoperatively

according to the CTS-SSS. Recovery of and improvement in grip power was after 3 and 6 months postoperatively, respectively. Three-jaw chuck pinch had increased significantly after 1 month postoperatively.

Discussion

CTS in long-term hemodialysis patients has incidence varying from 2% to 32%, which is higher than in the general population. When it occurs in renal dialysis, it is also recognized as more serious and unremitting.^{4,5} The etiology of CTS in long-term hemodialysis patients is unclear, and several factors relate to it, including duration of dialysis, age, old age of patient at onset of dialysis, dialysis-membrane characteristics, advanced glycosylation end products, low serum albumin, diabetic nephropathy, β_2 -microglobulin (β_2 MG) clearance, and sex.^{5,16–18} In our study, female hemodialysis patients receiving surgery comprised 67%, which is compatible with a Japanese cohort study,¹⁸ although other studies had different results, with an equal ratio or more of the male population.^{5,19} Dialysis-related β_2 MG amyloidosis deposits are considered to connect with synovial tissue hyperplasia, TCL thickening and flexor tenosynovitis, and all these may cause CTS. There were 61 (73%) patients that also had trigger finger in our study, and this may have been a coeffect of dialysis-related β_2 MG amyloidosis. Some studies have emphasized that AVF has a local effect on the carpal tunnel in CTS,^{20,21} while AVF on arms with CTS show a higher blood-flow rate and double entrapment of the median nerve is associated with polytetrafluoroethylene loop graft.¹⁶ Other studies found no clear relationship between AVF and CTS.^{2,4,22} In the current work, 89% of the hemodialysis patients had CTS at the AVF site. Many systemic diseases contribute to and exacerbate the development of CTS, including rheumatoid arthritis, diabetes mellitus, gout, acromegaly, hypothyroidism, and amyloidosis.^{2,23} In this study, 41% patients had diabetes.

CTS caused by long-term hemodialysis has been recognized as a relatively advanced and incurable disease with acute progression. The rather limited postoperative improvement after CTR and the high recurrence rate would be expected in

Table 2 Results of ultrasound-assisted percutaneous release on outcome measurements after surgery

	Baseline	1 week	1 months	3 months	6 months	12 months	18 months	24 months	P-value
Physical exam/telephone*	113/0	109/4	110/3	90/23	101/12	89/24	63/50	41/72	
CTS-SSS	3.49±0.87 ¹	2.45±0.63 ²	2.13±0.73 ^{2,3}	1.69±0.52 ³	1.58±0.13 ³	1.36±0.31 ⁴	1.25±0.21 ⁴	1.18±0.13 ⁴	0.05
≥3 (numbness), n (%)**	97(85.8%)	37(32.7%)	24(21.2%)	8(7%)	4(4%)	2(1.7%)	0	0	
≥3 (pain), n (%)**	89(78.7%)	13(11.5%)	10(8.8%)	3(2.7%)	2(1.7%)	0	0	0	
CTS-FSS	3.73±1.03 ¹	2.75±1.73 ²	2.14±0.81 ²	1.76±0.33 ²	1.31±0.29 ³	1.17±0.13 ³	1.09±0.18 ³	1.00±0.00 ³	0.05

Notes: *Number of patients assessed using a physical examination and face-to-face interview (physical examination) and a telephone interview (telephone). Both interviews used the same questionnaire. **|= no numbness/pain, 2= mild numbness/pain, 3= moderate numbness/pain, 4= severe numbness/pain, 5= very severe numbness/pain. Superscript numbers indicate statistically significant differences across time points by post hoc test.

Abbreviations: CTS-SSS, Carpal Tunnel Syndrome Symptom Severity Score; CTS-FSS, Carpal Tunnel Syndrome Functional Status Score.

Table 3 Results of ultrasound-assisted percutaneous release on sensations and strengths after surgery

	Baseline	1 week	1 month	3 months	6 months	12 months	18 months	24 months	P-value
Physical exam, n	113	109	110	90	101	89	63	41	
S2PDS (mm)	9.97±0.73 ¹	7.38±1.43 ²	6.47±1.85 ^{2,3}	6.19±1.33 ³	6.06±1.78 ³	6.17±1.73 ³	5.93±0.76 ⁴	5.35±0.62 ⁴	0.05
≥6 mm, % (n)	70.5 (91 of 113)	46.7 (51 of 109)	36.3 (40 of 110)	23.3 (21 of 90)	16.8 (17 of 101)	20.2 (18 of 89)	11.1 (7 of 63)	4.9 (2 of 41)	
SW monofilament (mm)	5.17±0.67 ¹	4.61±0.73 ¹	4.17±0.77 ²	3.96±0.79 ²	3.81±0.75 ²	3.67±0.71 ²	3.27±0.67 ³	3.14±0.87 ³	0.05
Grip power (kg)	15.17±14.3 ¹	5.03±4.6 ²	11.71±10.9 ³	13.35±11.2 ^{1,3}	16.79±9.5 ⁴	17.13±10.8 ⁴	16.01±10.9 ⁴	17.11±10.6 ⁴	0.05
Three-jaw chuck pinch (kg)	4.87±2.1 ¹	3.12±3.4 ¹	6.10±2.7 ²	6.12±3.1 ²	7.14±3.2 ³	8.19±3.1 ⁴	8.03±3.4 ⁴	9.98±3.6 ⁴	0.05

Notes: Data presented as means ± SD unless otherwise indicated. Superscript numbers indicate statistically significant differences across time points by post hoc test.

Abbreviations: S2PDS, static two-point discrimination test; SW, Semmes-Weinstein.

these patients.^{4,16,17} Surgical decompression of the carpal tunnel is suggested if conservative treatment has failed. ECTR, OCTR, and extended OCTR are usually used in these patients. Nonetheless, there are still no randomized trials or well-established guidelines for treating CTS in long-term hemodialysis patients. Theoretically extended or standard OCTR may enable better view and complete decompression of the median nerve. Neurolysis and synovectomy can be done in the same procedure, although the effects remain controversial.^{7,24} ECTR is another alternative, with advantages of less postoperative pain, faster return to work, and fewer wound-related complications. One study revealed that ECTR was effective for idiopathic and long-term hemodialysis patients, but with less satisfactory electrophysiological outcomes in the latter.⁷ A meta-analysis compared OCTR and ECTR and found that the final results were the same.⁸ However, the restricted endoscopic views of the anatomic structures, incomplete decompression, and potential risk of nerve or vessel injury limit the use of ECTR.^{25–28}

Ultrasound-assisted percutaneous release of trigger fingers is now a safe and easy procedure,^{29–31} and ultrasound-guided PCTR has been described in several studies with satisfactory preliminary results without injury to the surrounding structures.^{11,14,32–34} In a pilot open-label uncontrolled trial of 39 patients with ultrasound-guided PCTR, Locoq et al observed scores for pain, formication, and discomfort experienced in the hand had significantly reduced by day 15.³² Chen et al presented results of ultrasound-guided PCTR in 91 CTS cases with 100% sensory disturbances, and showed 98.9% moderate pain disappearance in 12 months postoperatively without intra- or postoperative complication.¹¹ Another study involving 128 CTS patients who received ultrasound-guided PCTR reported significant improvement in symptom severity and functional status scores 6 months postoperatively. Follow-up magnetic resonance imaging showed a complete section of TCL and nerve decompression in all patients.³³ In a randomized clinical trial to compare ultraminimally ultrasound-guided PCTR and blind mini-OCTR, Rojo-Manaute et al found the former provided early functional return and less postoperative morbidity with the same neurological recovery.³⁴ The main advantages of ultrasound-guided PCTR are minimizing surgical damage to nonetiologic anatomic structures and accurately restricting release to the anatomic fibrous layer by small skin incisions, strong resolution, and real-time ultrasound monitoring. The tools of the technique are simple and inexpensive. The procedure has been proved to be safe and effective in cadaveric and clinical studies.^{10,11,32–34} The

limitation of the procedure is that it is technically demanding. Nonetheless, the technical difficulty may be overcome with high-frequency ultrasound, advance cadaver practice, and repeated preoperative skills training.

To our knowledge, there have been no previous clinical reports for ultrasound-guided PCTR in hemodialysis patients. In this study, 90% of the patients had subjectively good results with improved symptoms. All objective parameters (CTS-SSS, CTS-FSS, grip power, three-jaw chuck-pinch strength, Semmes-Weinstein monofilament tests, and two-point discrimination tests) showed progressive and significant improvements during 2 years of follow-up. The difference between subjective and objective satisfaction may be due to preoperative expectations of patients for surgery.

There are some limitations regarding this study. There was no comparative group to compare different surgical strategies. Also, there was no collection of complete follow-up data for the clinical examination (78.7% and 36.3% at 12 and 24 months, respectively), although there were telephone interviews using questionnaires (100%).

Conclusion

Ultrasound-guided PCTR is an effective and safe procedure in hemodialysis patients with CTS. Advantages include a less invasive procedure, no tourniquet needed, limited-infiltration anesthesia, minimal soft-tissue exploration, and relatively short operation time. Our data suggested this technique can reliably relieve clinical symptoms and restore grip and pinch strength. Ultrasound-guided PCTR is a safe and effective procedure in hemodialysis patients with CTS.

Disclosure

The authors report no conflicts of interest in this work.

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