

Targeted Muscle Reinnervation in Partial Hand Amputations

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Summary: Targeted muscle reinnervation (TMR) surgery has been shown to aid in prevention and treatment of neuropathic pain. Technical and anatomical descriptions of TMR surgery for upper extremity amputees (including transradial, transhumeral, and forequarter amputations) have been reported, yet such descriptions of TMR surgery for partial hand amputations are currently lacking. Herein we outline the technique of different types of partial hand amputation TMR surgeries to serve as a reference and guide. A retrospective review was performed by our multi-institutional team to identify clinical cases where partial hand TMR surgeries were performed. Patient demographics, characteristics, amputation subtype, nerve transfer, pain score, pain outcome, and functional outcome data were collected and analyzed. From January 2018 to September 2019, 13 patients underwent partial hand TMR procedures. Eight cases resulted from trauma, and 6 were secondary to oncologic procedures. The amputations consisted of 8 ray, 2 trans-metacarpal, 2 radial-sided hand, and 1 index finger amputation with recurrent painful neuromas. Twelve patients were weaned off narcotics completely and only 3 remained on a neuromodulator for ongoing pain control. Technical considerations for partial hand TMR surgery have been outlined, with early pilot data showing beneficial pain control outcomes. (*Plast Reconstr Surg Glob Open* 2021;9:e3542; doi: 10.1097/GOX.0000000000003542; Published online 28 May 2021.)

INTRODUCTION

Targeted muscle reinnervation (TMR) is a nerve transfer technique where proximal stumps of the transected major peripheral nerves are transferred to recipient redundant motor nerves within adjacent muscles of the amputated limb.¹⁻⁵ TMR purposefully redirects the major peripheral nerve into an intact nerve and accompanying neuromuscular junction that is acutely transected, thus providing a target for the regenerating major peripheral nerve to grow into rather than forming a painful neuroma.¹⁻⁵

TMR provides additional benefits by increasing electromyography signal generators for advanced myoelectric prostheses.^{1,6,7} Following reinnervation, the targeted muscles then serve as biological amplifiers for motor nerve signals, providing increased electromyography signals and focused, intuitive control of prosthetic devices.^{1,6-12} TMR facilitates advanced prosthetic device use and desensitizes the amputation stump by decreasing symptomatic neuroma development. Subsequent studies demonstrated that TMR can also effectively reduce phantom limb pain (PLP) or phantom limb sensation.^{3-5,8}

In this study, we describe the application of TMR to 4 different, but common types of partial hand amputations, and outline the technique for others to employ in their practices. This technique was applied to manage partial hand amputations in the clinical setting at multiple academic and community medical centers.

METHODS AND MATERIALS

Multiple partial hand amputations, including ray, radial- and ulnar-sided partial hand, and metacarpal/

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Table 1. Patient Demographics and Characteristics

	No. Patients (%), N = 13
Demographics	
Men	11 (85%)
Mean age, y (range)	45.1 (18–72)
Mean follow-up, mo (range)	19.8 (5–35)
Hand involved	
Right	8
Left	5
Partial hand amputation type	
Ray amputation	8
Thumb amputation	1
Radial hand	2
Trans-metacarpal	2
Type of TMR performed	
Primary TMR	10
Secondary TMR	3

trans-metacarpal amputations, were performed. Coaptation of transected proximal major nerves from the amputation procedure was performed to end motor nerve targets in an end-to-end fashion following TMR principles.^{1,2}

Clinical Correlate Cases

After institutional review board approval at our respective centers, the authors reviewed patients who had undergone partial hand amputations where TMR was performed. Thirteen cases were identified between January 2018 and September 2019. Six cases were completed for oncologic reasons, 6 cases were related to trauma, and 1 case was for recurrent neuropathic neuromas. Primary TMR was

Table 2. TMR Clinical Case Descriptions, with Nerve Transfer and Outcome Descriptions

Category	Case	Amputation Sub-type	TMR Transfers Performed	Primary versus Secondary TMR	VAS Pain Scores	Pain Outcomes and Medication Usage	Functional Outcome
Oncologic	1	3 rd Ray amputation right hand	Proper digital-to-VIMMN x2	Primary—neuroma prevention	0/10	Neuroma pain: none PLP: none Narcotics: none	Prosthetic not indicated
	2	3 rd Ray amputation right hand	Proper digital-to-VIMMN x2	Primary—neuroma prevention	0/10	Neuroma pain: none PLP: none Narcotics: none	Prosthetic not indicated
	3	4 th Ray amputation left hand	Proper digital-to-VIMMN x2	Primary—neuroma prevention	1/10	Neuroma pain: none PLP: mild Narcotics: none	Prosthetic not indicated
	4	Thumb amputation/radial partial left hand	(1) Thumb PDN-to-adductor pollicis MN; (2) Thumb PDN-to-FPB-MN (3) RSN to AIN MB	Primary—neuroma prevention	0/10	Neuroma pain: none PLP: none Narcotics: none	Tolerates prosthetic wear
	5	3 rd Ray amputation left hand	Proper digital-to-VIMMN x2	Primary—neuroma prevention	2/10	Neuroma pain: none PLP: mild Narcotics: none +neuromodulator	Prosthetic not indicated
	6	4 th Ray amputation right hand	Proper digital-to-VIMMN x2	Primary—neuroma prevention	0/10	Neuroma pain: none PLP: none Narcotics: none	Prosthetic not indicated
Trauma	7	3 rd Ray amputation right hand	Proper digital-to-VIMMN x2	Primary—neuroma prevention	1/10	Neuroma pain: none PLP: Mild Narcotics: none	Prosthetic not indicated
	8	3 th Ray amputation left hand	Proper digital-to-VIMMN x2	Primary—neuroma prevention	1/10	Neuroma pain: none PLP: mild Narcotics: none	Prosthetic not indicated
	9	Trans-metacarpal right hand	(1) RSN-to-AIN MB; (2) Proper digital MN-to-2 nd & 3 rd VIMMNs; (3) Proper digital Uln-to-4 th VIMMN	Secondary—neuroma treatment	<i>Pre-TMR:</i> 6-7/10 <i>Post-TMR:</i> 2/10	Neuroma pain: none PLP: mild Narcotics: +opioid +neuromodulator	Tolerates prosthetic
	10	Trans-metacarpal right hand	(1) RSN-to-MN-Thenar MB; (2) MN-to-2 nd VIMMN; (3) Uln-to-3 rd DIMMN	Primary—neuroma prevention	0/10	Neuroma pain: none PLP: None Narcotics: none	Declined prosthesis
	11	Radial-sided partial left hand	Thumb PDN-to-index LMMN	Secondary—neuroma treatment	<i>Pre-TMR:</i> 7-8/10 <i>Post-TMR:</i> 1/10	Neuroma pain: none PLP: none Narcotics: +preoperative and none postoperative	Tolerates prosthetic
Recurrent neuroma	12	Radial-sided partial right hand	(1) Thumb PDN-to-adductor pollicis MN; (2) Thumb PDN-to-FPB-MN Proper digital-to-LMMN	Primary—neuroma prevention	0/10	Neuroma pain: none PLP: none Narcotics: none	Tolerates prosthetic
	13	Index finger ray amputation of right hand	Proper digital-to-LMMN	Secondary—neuroma treatment	<i>Pre-TMR:</i> 7-8/10 <i>Post-TMR:</i> 2/10	Neuroma pain: none PLP: none Narcotics: +preoperative and none postoperative neuromodulator: +preoperative and none postoperative	Prosthetic not indicated

performed in 10 cases, and secondary TMR in 3 cases. Ten patients were adults and 3 were children, with an average age of 45 at the time of surgery. The distribution of cases consisted of 8 ray amputations, 2 trans-metacarpal level amputations, 2 radial sided partial hand amputations, and 1 index finger amputation with recurrent painful digital neuromas. Follow-up ranged from 5 to 35 months, with mean follow up of 19.8 months (See Table 1). Of these patients, 12/13 were able to be weaned off of narcotics and only 3/13 remained on a neuromodulator (eg, Gabapentin or Lyrica). No postoperative neuroma pain was reported in any of the cases. Postoperative PLP was reported as “none” in 8/13 cases and “mild” in 5/13 cases. A prosthetic device was not indicated in 8/13 cases, and tolerated in 4/4 cases where a prosthetic device was utilized. Although indicated, 1 patient declined the use of a prosthetic (Table 2). (See Video [online], which displays middle finger ray amputation.)

OPERATIVE TECHNIQUE

The first step is meticulous dissection and appropriate identification of the major peripheral nerves and the recipient target motor nerves. Nerve stimulators are useful in identifying target recipient motor nerve branches

intraoperatively. Typically, we use an amperage of 0.5–2 mA and voltage of 50–100 and coordinate preoperatively with our Anesthesia colleagues to avoid long acting paralytic and muscle relaxant agents.

The nerve stumps must be resected sharply back to healthy fascicles. The nerve transfer coaptation site must be tension free; if this is not possible, nerve grafting or other coaptation aids may be indicated. The nerve transfers should be performed with the joints positioned under maximal stress to ensure the nerve coaptations are guaranteed tension-free in all positions. When nerve grafts are required, we prefer spare parts from the amputated extremity if available; otherwise, we consider processed nerve allograft or traditional autologous nerve.

For nerve preparations, we recommend transecting recipient target motor nerves *proximally*, and the transferred nerve stumps *distally* with neuroorrhaphies performed using a hand-sewn interrupted epineurial technique using 8-0 nylon sutures. To limit regeneration times, shorten the distal target motor nerve stump as much as possible, while preserving a tension-free nerve coaptation. The neuroorrhaphies are performed within 1–2 millimeters of where the recipient motor nerves enter the recipient muscles. Performing the coaptation close to the motor nerve

Table 3. Nerve Transfers Available for Partial Hand Amputations

Proximal Major Peripheral Nerve and Associated Distal Branches	Ulnar Nerve Branches	Median Nerve	Radial Sensory Nerve
Target motor nerves available for receipt of major peripheral nerve branch transfer	Volar/dorsal interossei Lumbricals 4/5 AIN to PQ FCU Any other local motor branch	Volar/dorsal interossei Lumbricals 2/3 AIN to PQ Thenar motor branch Any other local motor branch	1st dorsal interossei Thenar motor branch AIN to PQ Any other local motor branch

Table 4. Specific Nerve Transfer Options for Each of the Various Partial Hand Amputation Subtypes

Partial Hand Amputation Subtype	Nerve Transfer Options		
	Ulnar Nerve	Median Nerve	Radial Sensory Nerve
Ray amputation	<i>Utilize motor nerve target of residual muscles adjacent to resected ray:</i> <ul style="list-style-type: none"> • Volar interosseus muscle • Dorsal interosseus muscle • Lumbrical 		
Radial hand amputation: <i>Transfer sensory nerve stumps to ulnar hand (away from prosthesis docking site)</i>	N/A	Volar interossei 2/3 Dorsal interossei 3/4 Lumbricals 4/5	Dorsal interossei 3/4 AIN to PQ
Ulnar hand amputation: <i>Transfer sensory nerve stumps to radial hand (away from prosthesis docking site)</i>	Volar interossei 1/2 Dorsal interossei 1/2 Lumbricals 2/3	N/A	N/A
Trans-metacarpal ± thumb amputation	Volar interossei 2/3 Dorsal interossei 3/4 Lumbricals 4/5	Volar interossei 1/2 Dorsal interossei 1/2 Lumbricals 2/3	1 st dorsal interosseus thenar motor branch AIN to PQ
Trans-carpal amputation	Volar interossei 2/3 Dorsal interossei 3/4 Lumbricals 4/5 FCU AIN to PQ	Volar interossei 1/2 Dorsal interossei 1/2 Lumbricals 2/3 Thenar motor branch AIN to PQ	1 st dorsal interosseus thenar motor branch AIN to PQ
Starfish procedure**	Interossei dissected out, pedicled on the palmar neurovascular arch, and transposed to the dorsal hand		

**From Gaston et al.¹³

entrance to the muscle belly permits the shortest possible reinnervation time and more rapid onset of reinnervation.

Nerve-nerve coaptation size mismatch frequently occurs when performing TMR in partial hand amputations, but does not seem to adversely affect the outcome.¹⁴ Following nerve transfer, the coaptation site is reinforced and wrapped using a cuff of local denervated muscle carefully raised from around the interface of the native target motor nerve and its corresponding muscle units.¹⁴ This denervated muscle

cuff is gently wrapped around the nerve coaptation site and secured with interrupted 4-0 Vicryl suture.

Although tension-free TMR to a short recipient motor nerve with a muscle cuff is ideal, this is not always possible. If the motor branch to the target muscle is left long and the coaptation site too proximal for a muscle cuff to reach, nerve conduits, protectors or wraps can assist with coaptation size mismatches. Refer to [Tables 3](#) and [4](#) for recommended nerve transfer options.

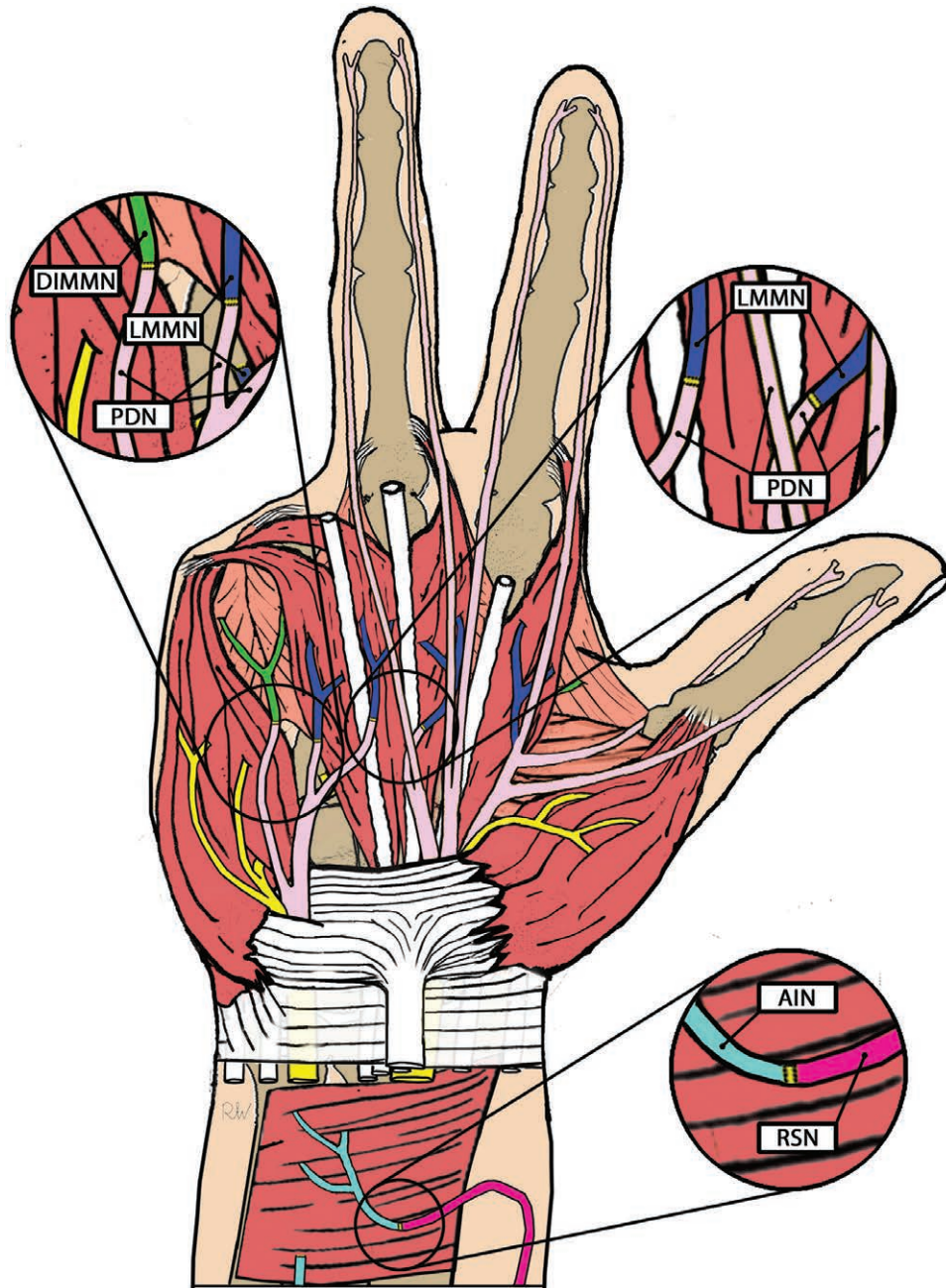


Fig. 1. Partial hand amputation TMR schematic diagram illustrating various TMR donor/target combinations in the setting of partial hand amputation. LMMN, lumbrical muscle motor nerve; DIMMN, dorsal interosseus muscle motor nerve; PDN, proper digital nerve; AIN, anterior interosseus nerve; RSN, radial sensory nerve.

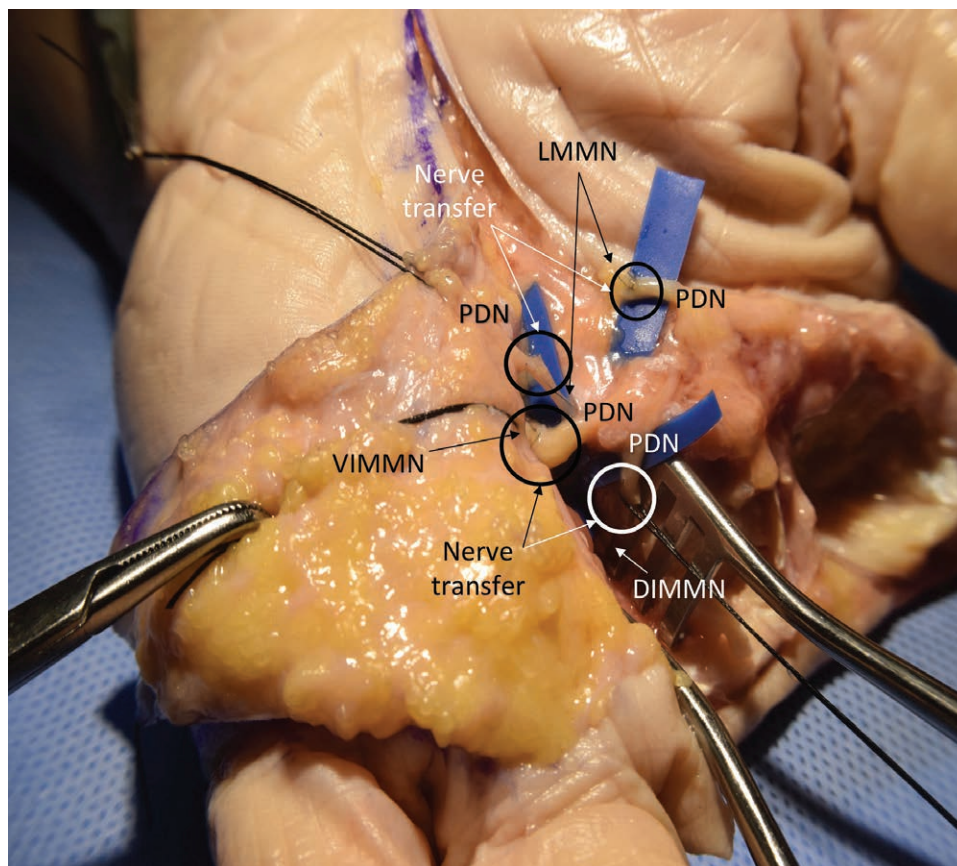


Fig. 2. Radial hand amputation, TMR cadaver dissection, following completion of 4 TMR nerve transfers. LMMN, lumbro-motor nerve; VIMMN, volar interosseus muscle motor nerve; PDN, proper digital nerve; DIMMN, dorsal interosseus muscle motor nerve.

DISCUSSION

It is important that these procedures are performed without a paralytic or local nerve blockade. Adequate soft tissue coverage must be maintained over the sites of neuro-rhaphy. Consideration should be given to utilization of spare parts for grafts. If a regional flap is to be performed, extended dissections into the recipient muscles targeted for reinnervation should be spared (ie, the TMR targets).

Postoperative therapy includes early mobilization of the remaining extremity, desensitization using neuromodulators such as gabapentin or pregabalin, mirror therapy, and exercises designed to trigger the nerve transfers. Patients must be preoperatively counseled on expectations and outcomes, including the importance of early therapy and neuromodulation. With well-performed TMR, minimal retraining is required for successful incorporation of a future myoelectric prosthesis.

CONCLUSIONS

We have described the specific steps and techniques necessary to perform TMR surgery for multiple partial hand amputation injuries. As reported in the literature, TMR yields improved pain control, decreased likelihood of PLP and phantom limb sensation, and a decrease in symptomatic neuroma formation.

Following TMR leads to greater ability to actively participate in aggressive postoperative therapy, improved tolerance of the prosthetic during rehabilitation and during the activities of daily living, and offers patients improved bioprosthetic function and potential.

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