The Successful Treatment of Chronic Pain Using Microcurrent Point Stimulation Applied to Scars

Abstract

Objectives: Although microcurrent is widely used for chronic pain and stress management, as well as scar or neural therapy as a popular approach for the treatment of pain, there remains considerable controversy as to their combined therapeutic value in chronic pain management. We aimed to determine the effect and magnitude that DC microcurrent therapy has when applied to physical scars and its effects on a wide variety of non-specific chronic pain syndromes.

Design: This was a cohort study analysis of treatment outcomes pre, post and 48-hour follow-up after Microcurrent Point Stimulation (MPS) was applied to physical scars on 51 patients with history of non-specific pain.

Interventions: An MPS Scar Release protocol was applied bi-laterally to physical scars. Evaluations entailed a baseline Visual Analogue Score (VAS) pain scale assessment, which was repeated after an electro-therapy treatment and 48 hours later. All 51 patients received one Microcurrent Point Stimulation Scar Release session.

Outcome Measures: The VAS response of the 51 patient sample with chronic pain reflected a statistically significant reduction of 3.706 points or 59% reduction in mean pain levels post MPS Scar Release application, when compared to initial pain levels [95% CI (3.033, 4.379; p=0.0001]. When VAS was measured at 48-hour follow-up, there was another statistically significant reduction of 0.902 points or 34% reduction in mean pain levels post treatment [95% CI (0.406, 1.398; p=0.001]. Together, MPS Scar Release protocol produced a statistically significant reduction of 4.608 points or 73% reduction in mean pain levels post treatment, when compared to initial pain levels [95% CI (3.940, 5.275); p=0.0001].

Conclusion: The positive results in this study could have applications to patients who have physical scars and are impacted by chronic pain syndromes.

Keywords: Scars; Microcurrent point stimulation; chronic pain

Abbreviations : ANS: Autonomic Nervous System; AC: Alternating Current; DC: Direct current; VAS: Visual Analogue Scale

Introduction

Chronic pain affects millions of people every year and the effects of pain result in tremendous health care costs, in terms of rehabilitation and lost worker productivity, plus the emotional and financial burden it places on patients and their families. According to a recent Institute of Medicine Report: Relieving Pain in America: A Blueprint for Transforming Prevention, Care, Education, and Research, pain is a significant public health problem that costs society at least \$560-\$635 billion annually, an amount equal to about \$2,000 for every living person in the U.S. This includes the total incremental cost of health care due to pain that ranges from \$261-\$300 billion to \$297-\$336 billion related to lost productivity (based on days of work missed, hours of work lost, and lower wages) [1,2]. In addition, there is currently a massive concern with the enormous use/abuse of analgesics and opioids throughout the USA [3-5]. If pain control can be achieved

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through other means as exemplified in this report, this could then impact favourably on this problem. Scars and trauma have long been recognized in neural therapy as a source of chronic pain as a result of Autonomic Nervous System (ANS) (in particular sympathetic nervous system) upregulation [6-10]. It is theorized that damaged local cells lose their normal membrane potential, transmitting abnormal electric signals throughout the rest of the body via the autonomic nervous system, acting as physical agonists to sympathetic upregulation (stress) and pain [11].

Traditionally, the modality of choice for electro-therapy has been Alternating Current (AC) [12-14]. However, there are two known types of electrical currents, AC and Direct current (DC). AC moves bi-directionally and is applied in the miliamperage range (10-3 amperes), usually called TENS or electro-acupuncture (EA) [15-17]. DC is uni-directional and when applied in the microamp or millionth of amp (10-6 -amperes) range to acupuncture or trigger points, is called Microcurrent Point stimulation (MPS) [18-19]. Microcurrent therapies involve applying weak direct currents (80μ A - 1mA), and are now being increasingly recognized as an adjunct for pain relief and autonomic nervous system regulation [20-24]. It is theorized that electro-acupuncture and microcurrent electro-currents have different modulating affects on the autonomic nervous system and pain outcomes [25]. There is no consensus in the literature identifying the best practice measures for microcurrent applied to scars for the treatment of chronic pain. Although sufficient evidence supports the application of microcurrent and neural therapy for chronic pain, there is limited evidence in literature to support the application of electro-therapies to scars to reduce chronic pains. The purpose of this pilot study was to assess the impact of Microcurrent Point Stimulation Scar Release protocol applied to physical scars on the pain levels in a random sample of chronic pain patients, after single application.

Patients, Materials and Methodology

This study entailed the use of MPS in 51 patients (42 females, 9 males; mean age 47 years, SD 12.81) with chronic non-specific pains with a mean pain duration average of 7.61 years (SD 1.34) (Table 1) presenting to us for therapy of their problem. The location of the scars are shown in Table 2 and the pain location sites in Table 3. Inclusion criteria were simple: patients who were currently suffering from chronic pain for greater than 3 months,

with a recorded >4 VAS Pain Scale score and have visible physical scar(s). Physical scars were defined as surgical or trauma induced. The diagnoses of pain, location, severity, sex, previous interventions or surgeries were not considered exclusion criteria. Informed consent was obtained to partake in treatment and the study assessments. Patient pain scores were recorded immediate pre treatment and twice post treatment: immediately after application, and again 48 hours later. Microcurrent Point Stimulation was simultaneously applied bi-laterally to scars using [26] two Dolphin Neurostim (Center for Pain & Stress Research Ltd, Ontario, Canada) devices. This is an FDA-approved device which apply low frequency, concentrated, microcurrent stimulation for the relief of chronic pain and stress [24-25]. MPS application time was 30 seconds per point at approximate one-half inch intervals along the length of the scars. Polarity of application is important, as on one side of the scar, the device is set to negative pole (-) and on the other side of scar, the second device is set to a positivenegative pole (+/-). The intent of this methodology is to push a negatively charged current back and forth through a positively charged (oriented) scar tissue. For the purpose of this study, only physical scars were treated, with the average treatment duration time of 30 minutes per patient.

Table 1: Descriptive statistics.

Descriptive Statistics							
	N	Minimum	Maximum	Mean	Std. Deviation		
Duration of Pain	51	90 days	36 years	7.61 years	1.339 years		
Age (years)	51	21	72	47.34	12.812		
Pain Before Treatment (0-10)	51	4	10	6.33	1.912		
Pain after Treatment (0-10)	51	0	10	2.63	1.913		
Follow up Pain - 2 Days after Treatment(0-10)	51	0	10	1.73	1.930		

Table 2: Scar Location and percentages in the 51 patients.

Scar Location	Total Number=51	Percentage 100%
Abdomen	31	60.70%
Knee	8	15.6%
Hernia (abdomen)	3	5.88%
Ankle	3	5.88%
Breast	2	3.9%
Neck	2	3.9%
Wrist/Hand	2	3.9%
Total	51	100%

Visual Analogue Scale (VAS) was used to evaluate the patient's pain. The VAS is an 11-point scale from 0-10 with 0 being no pain and 10 being the most intense pain imaginable [27-30]. The patient verbally selects a value that is most in line with the intensity of the pain that they have experienced in the last 24 hours or is often reported as a rating during a specific movement pattern or functional task. The VAS has good sensitivity and excellent test-retest reliability [31].

Table 3: Pain Location in the 51 patients with percentages.

Pain Location	Total Number=51	Percentage 100%
Back	20	39.2%
Neck	9	17.6%
Shoulder	8	15.6%
Scapula (shoulder)	1	1.96%
Hip	4	7.85%
Finger/Hand	3	5.88%
Abdomen	3	5.88%
Arm	1	1.96%
Knee	1	1.96%
Total	51	100%

The aim of this cohort preliminary study was to evaluate whether

a) Microcurrent Point Stimulation Scar Release protocol, when applied to SCARS, can modulate or improve VAS pain scale in patients suffering with chronic pain.

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b) Microcurrent Point Stimulation Scar Release protocol, when applied to scars, is a valid option for the nonpharmacological pain management of chronic pain conditions.

Results

Outcome Measures

The VAS response of the 51 patient sample with chronic pain reflected a statistically significant reduction in mean post pain levels of 3.706 points or 59% reduction in mean pain levels post MPS application to physical scars, when compared to initial pain levels [95% CI (3.033, 4.379; p=0.0001]. When VAS was measured at the 48 hour follow-up, there was a further statistically significant reduction of 0.902 points or 34% reduction in mean pain levels post treatment [95% CI (0.406, 1.398; p=0.001], for a total pain reduction of 4.608 points or 73% reduction in mean pain levels post MPS treatment, when compared to initial pain levels [95% CI (3.940, 5.275); p=0.0001] (Figure 1). There was no correlation between pain location and site of the physical scars (Tables 2 & 3).



Discussion

Chronic pain often equates to stress, both of which can make our daily lives miserable, and can lead to significantly impaired physical health and high societal costs [1,2]. For some time now, chronic pain has been difficult to diagnose and treat for many health care professionals. When the millions of physical scars produced annually throughout North America [32,33] are combined with the day-to-day accumulated patient traumas, the data represents a significant pre-existing pool of stress and pain patients within the general population [32,33]. It may help to explain the causation of symptoms for millions of chronic pain sufferers. In addition, the long-term use of opioids is now approaching epidemic levels in the USA, with few viable solutions for treatment in the foreseeable future [3-5]. Treatments like the kind described in this report could have a favourable impact on this problem.

The data from this study clearly shows that the application of Microcurrent Point Stimulation to physical scars had a marked improvement in pain outcomes when compared to baseline measurements in chronic pain patients. The improved outcomes were even more impressive given the patient sample for pain duration (mean 7.61 years) and the intensity (mean 6.33/10) improved after a single MPS scar release application. Increased pain relief between post application and results 48 hours later was also noted as an improved outcome, suggesting internal functional changes may have occurred. It is suggested in the literature that DC microcurrent mimics human biocellular communications, enhancing autonomic nervous system regulation and the production of beta-endorphins, resulting in a body-wide therapeutic benefits [21,35]. These biochemical processes may provide a plausible explanation for the improved pain modulation over time after concentrated DC microcurrent is applied, and is an area where future research is required. We have previously reported, in several published studies, reduction in pain and cortisol with improvements in autonomic nervous system functionality in patients using MPS [18-21]. The consistency of chronic pain outcome improvements through the application of MPS to physical scars suggests there may be a strong relationship between chronic pain symptomology and physical scars throughout the body.

Conclusion

Chronic pain can limit quality of life, restrict work and social engagement, and is often blamed for the development of drug dependency of various forms. This study showed MPS Therapy applied to physical scars provided statistically significant reduction in initial pain levels with a further reduction after a 48-hour follow-up. These significant changes help validate the potential application of MPS to SCARS as an viable option to treating patients with non-specific soft tissue chronic pain. However, long term further investigation is warranted with a larger focus group to confirm these results and to assess their duration.

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Statistical analyses were done by 3rd party freelance statistician using SPSS software, a widely used program for statistical analysis in social and medical science.

Author Disclosure Statement

All the authors whose names are listed in this study have a educational association with the sponsoring company that may create the appearance of a conflict of interest in connection with the submitted manuscript.

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