



ELSEVIER

Contents lists available at [ScienceDirect](http://ScienceDirect.com)

Journal of Neuroscience Methods

journal homepage: www.elsevier.com/locate/jneumeth

Basic Neuroscience

Evaluation of optimal electrode configurations for epidural spinal cord stimulation in cervical spinal cord injured rats

Monzurul Alam^a, Guillermo Garcia-Alias^b, Prithvi K. Shah^e, Yury Gerasimenko^{b,f,g}, Hui Zhong^b, Roland R. Roy^{b,c}, V. Reggie Edgerton^{b,c,d,*}^a Department of Neurosurgery, University of California, Los Angeles, CA 90095, United States^b Department of Integrative Biology and Physiology, University of California, Los Angeles, CA 90095, United States^c Brain Research Institute, University of California, Los Angeles, CA 90095, United States^d Department of Neurobiology, University of California, Los Angeles, CA 90095, United States^e Health & Technology and Neurobiology, Stony Brook University, Stony Brook, New York, United States^f Pavlov Institute of Physiology, St. Petersburg 199034, Russia^g Institute of Fundamental Medicine and Biology, Kazan Federal University, Kazan 420006, Russia

H I G H L I G H T S

- Dorsal funiculi were injured at C4 spinal cord in adult female rats.
- Epidural stimulation electrodes were implanted chronically at C6 and C8.
- Potentials evoked in forelimb muscles by cervical spinal stimulation were examined.
- Evoked potentials were dependent on electrode configuration and current intensity.
- Evoked potentials are useful biomarkers to identify optimal stimulation parameters.

A R T I C L E I N F O

Article history:

Received 5 January 2015

Received in revised form 7 March 2015

Accepted 9 March 2015

Available online 16 March 2015

Keywords:

Epidural stimulation

Cervical spinal cord injury

Dorsal funiculi crush

Motor evoked potentials

A B S T R A C T

Background: Epidural spinal cord stimulation is a promising technique for modulating the level of excitability and reactivation of dormant spinal neuronal circuits after spinal cord injury (SCI). We examined the ability of chronically implanted epidural stimulation electrodes within the cervical spinal cord to (1) directly elicit spinal motor evoked potentials (sMEPs) in forelimb muscles and (2) determine whether these sMEPs can serve as a biomarker of forelimb motor function after SCI.

New method: We implanted EMG electrodes in forelimb muscles and epidural stimulation electrodes at C6 and C8 in adult rats. After recovering from a dorsal funiculi crush (C4), rats were tested with different stimulation configurations and current intensities to elicit sMEPs and determined forelimb grip strength. **Results:** sMEPs were evoked in all muscles tested and their characteristics were dependent on electrode configurations and current intensities. C6(–) stimulation elicited more robust sMEPs than stimulation at C8(–). Stimulating C6 and C8 simultaneously produced better muscle recruitment and higher grip strengths than stimulation at one site.

Comparison with existing method(s): Classical method to select the most optimal stimulation configuration is to empirically test each combination individually for every subject and relate to functional improvements. This approach is impractical, requiring extensively long experimental time to determine the more effective stimulation parameters. Our proposed method is fast and physiologically sound.

Conclusions: Results suggest that sMEPs from forelimb muscles can be useful biomarkers for identifying optimal parameters for epidural stimulation of the cervical spinal cord after SCI.

© 2015 Elsevier B.V. All rights reserved.

1. Introduction

Epidural electrical stimulation of the spinal cord, i.e., electrical enabling motor control (eEmc), is a promising therapy for the rehabilitation of sensorimotor function after spinal cord injury (SCI) (Edgerton and Roy, 2012; Alam and He, 2014; Dietz and Fouad,

* Corresponding author at: Department of Integrative Biology and Physiology, University of California, Los Angeles, 621 Charles E. Young Drive LS 1804, Los Angeles, CA 90095-1527, United States. Tel.: +1 310 825 1910; fax: +1 310 267 2071.
E-mail address: vre@ucla.edu (V.R. Edgerton).