Selective stimulation of the spinal cord surface using a stretchable microelectrode array

Kathleen Williams Meacham, Liang Guo, Stephen P. DeWeerth and Shawn Hochman

INTRODUCTION

Recovery of motor function lost to disease or injury is an objective shared by approximately 255,000 individuals in the United States living with spinal cord injury (NCSISC, 2008). This goal is also shared by 1.5 million Americans living with motor-control impairing neurological disorders (National Institute of Neurological Disorders and Stroke, 2009) as well as the five million individuals experiencing temporary or permanent loss of function due to stroke (Rosamond et al., 2008). This remaining spinal circuitry to evoke coordinated movements similar to volitional gait (Courtine et al., 2009). Spinal cord epidural stimulation (ES) approaches to reactivate movement have utilized some of the axons of sensory afferents as well as spinal neurons in the dorsolateral funiculus (DLF; Gerasimenko et al., 2008). Spinal neurons can also be activated via intraspinal microstimulation (ISMS), which involves insertion of stimulating microwires into gray matter laminae of the lumbar-sacral enlargement. This method has been used to recruit hindlimb activity in a selective, graded, and controllable manner (Tresch and Bizzi, 1999; Saigal et al., 2004). ISMS of the spinal cord has been shown to directly and indirectly activate motoneurons via intraspinal axons (Tresch and Bizzi, 1999) as well as the axons of sensory afferents whose collaterals extend many spinal segments (Gaunt et al., 2006).

By electrically stimulating the spinal cord, it is possible to activate functional populations of neurons that modulate motor and sensory function. One method for accessing these neurons is via their associated axons, which project as functionally segregated longitudinal columns of white-matter funiculi (i.e., spinal tracts). To stimulate spinal tracts without penetrating the cord, we have recently developed technology that enables close-proximity, multi-electrode contact with the spinal cord surface. Our stretchable microelectrode arrays (sMEAs) are fabricated using an elastomer polydimethylsiloxane substrate and can be wrapped circumferentially around the spinal cord to optimize electrode contact. Here, sMEAs were used to stimulate the surfaces of rat spinal cords maintained in vitro, and their ability to selectively activate axonal surface tracts was compared to rigid bipolar tungsten microelectrodes pressed firmly onto the cord surface. Along dorsal column tracts, the axonal response to sMEA stimulation was compared to that evoked by rigid microelectrodes through measurement of their evoked axonal compound action potentials (CAPs). Paired t-tests failed to reveal significant differences between the sMEAs and the rigid microelectrode's stimulus resolution, or in their ranges of evoked CAP conduction velocities. Additionally, dual-site stimulation using sMEA electrodes recruited spatially distinct populations of spinal axons. Site-specific stimulation of the ventrolateral funiculus – a tract capable of evoking locomotor-like activity – recruited ventral root efferent activity that spanned several spinal segments. These findings indicate that the sMEA stimulates the spinal cord surface with selectivity similar to that of rigid microelectrodes, while possessing potential advantages concerning circumferential contact and mechanical compatibility with the cord surface.

Keywords: multielectrode array, neuroprosthesis, brain computer interface, neurophysiology, spinal cord injury, PDMS, neurorehabilitation, spinal cord

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Hochman, Shawn, Department of Physiology, Emory University School of Medicine, 615 Michael Street, Atlanta, GA 30322, USA
E-mail: shochman@physio.emory.edu

Edited by:
Mario I. Romero-Ortega, University of Texas Southwestern Medical Center, USA
Reviewed by:
Xavier Navarro, Universitat Autonoma de Barcelona, Spain
Jeffrey Petruska, University of Louisville, USA
*Correspondence: Shawn Hochman, Department of Physiology, Emory University School of Medicine, 615 Michael Street, Atlanta, GA 30322, USA
E-mail: shochman@physio.emory.edu

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For both epidural and intraspinal stimulation, the axons of neurons are activated at lower thresholds than their associated cell bodies (Ranck, 1975; Gustafsson and Jankowska, 1976; Gaunt et al., 2006). Conveniently, the axons of most spinal neurons have collaterals located in longitudinal white matter tracts surrounding the central gray matter (i.e., spinal tracts), and have topographic relationships to distinct functional regions in gray matter laminae (Hochman, 2007). Direct contact and stimulation of these spinal tracts, therefore, has the potential to recruit functional subpopulations of spinal neurons. The ability of spinal cord surface stimulation to elicit coordinated motor neuron activity patterns, including fictive locomotion, has been demonstrated in the in vitro isolated spinal cords of neonatal rats (Iwahara et al., 1991; Magnuson and Trinder, 1997; Todd et al., 2009). In these studies, glass or metal electrodes were pressed directly onto the spinal cord surface at either the lower-thoracic ventrolateral funiculus (VLF; Magnuson and Trinder, 1997; Antonino-Green et al., 2002), the lumbosacral enlargement’s dorsal column (Iwahara et al., 1991), or the sacral spinal dorsal column (Todd et al., 2009).

Regionally precise stimulation of these and other spinal cord surface locations has the potential to enable a more controlled study and regulation of evoked neural activity. Selective surface stimulation can be facilitated by electrodes placed in very close contact to the interfaced neural tissue (Rutten, 2002; Rodgers et al., 2017). In addition, high spatial resolution MEAs have been shown to facilitate stimulus patterns that use spatiotemporal adaptation as well as fatigue-minimizing interleaving (McDonnall et al., 2004).

Safety constraints must be determined and followed when placing close-proximity, high-resolution MEAs on neural tissue; for example, the charge/phase and charge density should be below pre-determined thresholds found to be harmful to the interfaced biology (Merrill et al., 2005). Additionally, mechanical damage by either the electrodes or their substrate must be minimized. As the nervous system is comprised of soft tissue, arrays of rigid micro-electrodes have the potential to produce significant mechanical damage. Tissue/electrode property mismatches — including differences in mechanical compliance, damping, and density — are a particularly significant issue for long-term implantable devices. Consequences range from reduced interface integrity to irreversible loss of neuronal function due to scarring. Such damage has been evaluated and quantified for several types of MEAs (Biran et al., 2005; Leventhal et al., 2006; McConnell et al., 2009), and techniques for its reduction or prevention continue to be explored (Grill et al., 2009).

With the above-mentioned constraints in mind, we have developed an elastomer-substrate MEA technology for conformal contact with the spinal cord surface (Guo et al., 2007; Meacham et al., 2008; Guo and DeWeerth, 2009b; Figure 1A). This stretchable MEA (sMEA) is comprised of gold traces and electrodes that are photo-patterned between layers of thin-film polydimethylsiloxane (PDMS). PDMS is a gas-permeable elastomer with well-characterized biocompatibility (Belanger and Marois, 2001; Peterson et al., 2003; Guven et al., 2006) and expanding use as a microfluidic and cell-interfacing substrate. In addition, its elasticity is orders of magnitude greater than more traditional flexible electrode array materials, including parylene and polyimide (Yang et al., 1998; Armani et al., 1999; Rousche et al., 2001; Table 1). To further isolate the space between electrode and interfaced tissue, each electrode is surrounded by a raised, conical isolation well (Figure 1B; Guo et al., 2010). The impedance profiles of these electrodes compare favorably to those of rigid microelectrodes used for stimulation and recording of neurons (Guo et al., 2010), and iterations of this design have been shown to maintain electrical connectivity during both stretching (Meacham et al., 2008) and bending (Guo and DeWeerth, 2010).

In the following experiments, we hypothesize that the sMEA can stimulate spinal cord surface tracts with axonal selectivity that is the same as that evoked by a rigid, bipolar microelectrode pressed

![Figure 1](image-url)
Table 1 | Comparison of Young’s Moduli for MEA (multi-electrode array) substrates and spinal cord.

| Material       | Young’s modulus (stiffness) |
|----------------|-----------------------------|
| Parylene       | 4–4.5 GPa (Yang et al., 1998) |
| Polyimide      | 2.3–2.8 GPa (Rouse et al., 2001) |
| PDMS           | 0.05–1.79 MPa (Armanii et al., 1999; Brown et al., 2005) |
| Spinal Cord    | 0.1–1.4 MPa (Mazuchowski and Thibault, 2003) |

directly onto the cord. To test our hypothesis, we wrap sMEAs around isolated young rat spinal cords maintained in vitro, with electrodes overlying the dorsal surface for bipolar stimulation. Stimulus selectivity is quantified by measuring the orthogonal (lateral) spread of activated white matter tracts, as represented by evoked tract compound action potentials (CAPs).

Rigid microelectrodes pressed firmly upon – but not penetrating – the cord are used as a comparison because of their well-established use in experimental neurophysiology protocols requiring selective surface stimulation (Loeb et al., 1995). We also compare sMEA vs. rigid microelectrode’s ability to recruit axons of increasingly slower conduction velocities as a function of electrode proximity and stimulus magnitude.

We then evaluate the ability of dual-site sMEA stimulation to activate distinct axonal tracts, and the ability of single-pulse sMEA stimulation to evoke motoneuron activity in ventral roots associated with hindlimb flexor and extensor movements (Kichn and Kjaerulf, 1996).

MATERIALS AND METHODS
The spinal cords of 20 Sprague-Dawley young rats (postnatal days 10–14) were isolated and maintained in vitro for sMEA and rigid electrode interfacing and evaluation. All animal procedures were performed in accordance with policies of the Association for the Assessment and Accreditation of Laboratory Animal Care International and approved by Emory University’s Institutional Animal Care and Use Committee. Each rat was administered 10% wt./vol. urethane (2.0 mg/kg injected intraperitoneally) and, following confirmation of anesthesia, submerged in an ice slurry for 5 min to decrease body temperature. Following subsequent decapitation and evisceration, the cervical-to-sacral spinal cord was isolated along with ventral and dorsal roots and placed in an oxygenated (95% O2, 5% CO2) bath of artificial cerebrospinal fluid (aCSF, in mM: NaCl 128; KCl 1.9; D-glucose 10; MgSO4 1.3; CaCl2 2.4; KH2PO4 1.2; and NaHCO3 26). Further isolation, data collection, and analysis procedures were specific to experiments and are described in the following sections.

PREPARATION FOR MEA-SPINAL CORD INTERFACING
The fabrication steps for the conical-well sMEA technology have been reported previously (Guo et al., 2010). Briefly, our fabrication process involves lift-off patterning of gold features onto a PDMS substrate cured on a glass slide, lithographically defining sacrificial posts where electrode and contact pad openings are to be made in assisting the formation of conical wells, covering the sample with another thinner PDMS layer for encapsulation, and then removing the sacrificial posts to expose the electrodes and contact pads.

The sMEA substrate thickness is approximately 80 μm, with the isolation wells surrounding each electrode possessing an additional height of 40 μm. The exposed electrode surfaces are each approximately 100 μm in diameter, and minimum inter-electrode spacing (between electrodes in the same row) is 230 μm (Guo et al., 2010; Figure 1). Fabrication, measurement, and imaging of the sMEAs were undertaken using the Microelectronics Research Center core facility at Georgia Institute of Technology. A five-electrode configuration was chosen for these experiments; in related work, eight-electrode version of the sMEA with integrated packaging has also been implemented successfully (Guo and DeWeerth, 2009a).

Prior to in vitro testing, each sMEA was treated briefly with oxygen plasma and stored in deionized water for preservation of hydrophilicity (Guo et al., 2010). Contacts on the MEA were adapted to leads for the MCS STG-2008 stimulator (MultiChannel Systems) using 32 AWG wires (Belden, Inc.). These wires were bonded to each contact using conductive epoxy (CircuitWorks CW2400) and sealed with a thin layer of PDMS (Sylgard 184, Dow Corning). PDMS tabs were cut out of the sMEA substrate on either side of the five-electrode exposures prior to interfacing. These tabs facilitated conformal attachment of the sMEA to a single segment of the spinal cord, and enabled a snug, customized fit as the wrapped around tab could be threaded through an oval-cut opening in the opposite side (Figure 2). The sMEA was then wrapped around a spinal thoracic segment with the exposures facing inward.

ELECTROPHYSIOLOGY: WHOLE-CORD EXPERIMENTS
The first series of experiments assessed the degree to which adjacent axonal tracts were activated by single-site surface stimulation by the sMEA, as compared to that evoked by a rigid tungsten microelectrode placed directly onto the same longitudinal tract (Figure 2). Immediately following isolation, in vitro cords were secured using insect pins (1.0 mm, Fine Science Tools) in a Sylgard-coated Petri dish containing cold (4°C), oxygenated aCSF. Dura was carefully and thoroughly removed, and the cord was given an hour to equilibrate at room temperature before further experimentation.

To stimulate the cord, sMEA electrodes were placed directly on the center of the dorsal column at thoracic levels 6, 7, or 8. For comparison, a conventional tungsten bipolar electrode (conical-shaped exposed tip with base diameter 5 μm, exposed tip height 25 μm, 50–100 μm inter-electrode distance; Harvard Apparatus, Inc.) was pressed firmly onto – but not penetrating – the cord surface immediately caudal to the stimulating sMEA electrode. For all experiments, the tungsten electrode was placed at the same, fixed (approximately 45%) angle to the spinal cord to ensure consistent amount of electrode contact. To maximize the probability that this rigid control electrode position was activating the same longitudinal tract as the sMEA electrode, both electrode types were positioned to evoke a maximal CAP response on the same caudal tract, as confirmed with a surface recording electrode (Figure 2).

The dorsal column was chosen because its axon composition is well-characterized and because it possesses architecture comprised of mostly parallel tracts (Willis and Coggeshall, 2004). The
dorsal column is primarily responsible for conveying ascending proprioceptive, fine touch, and vibration information from primary afferents and a class of ascending tract neurons. It also contains Aδ and unmyelinated C visceral afferents (Willis et al., 1999). It has been shown that, in the adult rat dorsal column, at least 2/3 of the axons in the dorsal column are myelinated Aα, Aβ, or propriospinal fibers with diameters of 1–10 µm, and that the remaining 1/3 of axons are Aβ or C/unmyelinated propriospinal fibers with diameters of 0.2–1 µm (Chung and Coggeshall, 1983; Patterson et al., 1989). In the juvenile rat, however, the composition of the dorsal column has a higher ratio of unmyelinated-to-myelinated fibers (Chung and Coggeshall, 1987). To our knowledge, activation threshold in the juvenile rat dorsal column has not been investigated previously. However, the activation thresholds for these afferents within the peripheral nerve have been estimated in rats in this age range in vitro where it has been shown that they are recruited in the order of Aαβ, Aδ, and C with constant current stimulation intensities of 50 µA/50 µs, 500 µA/50 µs 500 µA/500 µs, respectively (Thompson et al., 1990). Associated conduction velocities for these afferent fiber groups in peripheral nerve have been reported as 2.4–6.7 (Aαβ fibers), 0.6–4.3 and (Aδ fibers), and 0.3–0.5 m/s (unmyelinated C fibers; Thompson et al., 1990).

For all white matter tract selectivity studies, the delivered stimuli were single, charge-balanced, 500 µs duration current pulses using adjacent electrodes (160 µm inter-electrode distance for sMEA, 50–100 µm inter-electrode distance for rigid tungsten bipolar electrode). A current (vs. voltage) delivery modality was employed based on evidence that regulated current waveforms guarantee generation of electric fields that are independent of electrode polarization (Guo and DeWeerth, 2009a). A charge-balanced square wave pulse was used to minimize possible charge injection damage to tissue (Merrill et al., 2005). The minimum current value at which a CAP was visible on any distal white matter tract was defined as the threshold stimulus value (T). Amplitude multiples of these threshold values (1.0–2.0T, in 0.2T increments) were used to stimulate the cord to evoke increasingly stronger evoked CAPs, and this series of stimuli were repeated as the recording electrode was moved incrementally across the surface of the cord.

To measure activation of axonal tracts, a single glass recording suction electrode (40–50 µm internal diameter; described in more detail below and shown in Figure 2) was used to record evoked CAPs in 50 µm lateral increments across the cord surface (MO-10 One-axis Oil Hydraulic Micromanipulator, Narishige International USA). This electrode was placed 10–15 mm caudal to the stimulation site (typically spinal lumbar level L2) to allow...
for temporal separation of stimulus artifact and the evoked CAP. A reference ground electrode was placed in the bath at the caudal end of the spinal cord. For a subset of these experiments, an additional, adjacent pair of sMEA electrodes was also used to stimulate the cord. This was done to determine the resolution of white matter tract selectivity achievable between adjacent sMEA electrodes (bipolar configuration).

**ELECTROPHYSIOLOGY: HEMISECTED SPINAL CORD**

For the second series of experiments, which evaluated motoneuron responses to sMEA surface stimulation, cord hemisections were performed to ensure sufficient surface area exposure for complete oxygenation of all spinal gray matter. Mid-sagittal hemisection was accomplished using 0.10 mm diameter insect pins (light brushing of the cord with the sharp end) and was performed immediately after isolation and placement of the cord into a bath of ice-cold, oxygenated, high sucrose-containing aCSF (in mM: sucrose 250; KCl 2.5; NaHCO3 26; NaH2PO4 1.25; d-glucose 25; MgCl2 3; CaCl2 1). Following hemisection, the spinal cord was placed in room temperature, oxygenated aCSF and allowed to equilibrate for at least 1 h before commencement of sMEA testing. Throughout the experiments that followed, the viability of spinal neurons was monitored intermittently by evaluating the magnitude of the reflex at the spinal cord lumbar level 5 (L5) ventral root evoked by stimulation of dorsal root L5.

Following equilibration of the hemisected cord, the sMEA was wrapped around the cord's lateral surface, and its electrodes (bipolar configuration) were used to stimulate the surface of 12th thoracic segment along the DLF, lateral funiculus (LF), and VLF. These spinal tract regions contain a combination of ascending, descending, and propriospinal axonal tracts of importance to movement coordination as well as sensory transmission (Davidoff, 1983; Loy et al., 2002; Noga et al., 2003). Single, monophasic current pulses were applied at each of three electrode pair sites, using minimum current values required to elicit a visible evoked response on either the recorded ventral roots or on that tract's recorded surface CAP. For recording, glass suction electrodes were placed on the ventral root at lumbar level 2 (L2), the ventral root at lumbar level 5 (L5), as well as on the surface of the spinal cord at the 6th lumbar level DLF and VLF. Previous work has shown that recordings from ventral roots L2 and L5 are predominantly associated with hindlimb flexor (L2) and extensor (L5) motor output, respectively (Kiehn and Kjaerulff, 1996). Ventral root recording was accomplished using bipolar glass suction electrodes (90–150 μm internal diameter glass, silver chlorided wire differential recording); surface recording electrodes had smaller internal diameter (40–50 μm).

**DATA COLLECTION AND ANALYSIS**

All extracellular responses were amplified by 10,000 and recorded using pClamp data acquisition software (Axon Instruments, new part of MDS Analytical Technologies). For a given stimulus amplitude and electrode type, 7–10 trials of spinal cord responses were recorded. The strength of the evoked responses for both CAPs and ventral root responses was quantified via MATLAB routines that rectified and integrated the signal for the time window of 100 ms post-stimulus, starting immediately after the end of a given stimulus artifact. To adjust for baseline noise, a 100-ms window of rectified, integrated pre-stimulus recording was subtracted from each value.

For selectivity comparisons across animals, values were also amplitude-normalized such that the maximal response was equal to 1. Visual verifications of response onset and offset values were performed for each trial used in analysis, and the consistency of response across trials was also verified visually. The time to onset and duration of a given response was calculated using a MATLAB detection algorithm, which used a combination of threshold detection and time windowing to identify the onset and offset of a given response. Values for delay and duration of evoked response were subsequently used, in combination with distance measurements, to calculate axonal fiber conduction velocities. When appropriate, paired Student’s t-tests with two-tailed distribution were used to statistically compare spinal cord responses to each of the electrode methodologies.

**RESULTS**

In the evaluation of in vitro stimulation efficacy of the sMEA, five different characteristics were assessed: charge density required for threshold activation of CAPs, spatial selectivity, axonal conduction-velocity selectivity, dual-site stimulus precision, and ability to recruit motoneurons. Spatial selectivity was defined as the ability of single-site stimulation to activate axons with a highly localized pattern comparable to conventional rigid stimulating electrodes.

Axonal conduction-velocity selectivity was defined as the sMEA’s ability to stimulate axonal fibers of incrementally lower conduction velocities as a function of stimulating electrode proximity and current amplitude. Dual-site stimulus precision was defined as the degree to which adjacent electrodes could discriminate between activation of parallel tracts of axons. Ability to recruit motoneurons was defined as the sMEA’s ability to evoke motor output via surface spinal tract stimulation, as measured by ventral root responses associated with hindlimb flexor and extensor activity (Kiehn and Kjaerulff, 1996). The following sections describe the results of these analyses, and demonstrate that the sMEA can be used to stimulate spinal tracts in a precise and controllable manner.

**CHARGE AND CURRENT DENSITY COMPARISON**

The minimum current required to evoke a detectable CAP on a recording electrode placed on the same longitudinal axonal tract (central–dorsal column) was evaluated for the sMEA and compared to those required for the rigid tungsten electrode. The threshold current required for the sMEA was found to be significantly greater than that required for the rigid electrode (153 ± 109 vs. 47 ± 24 μA, 500 μs biphasic pulses; paired Student’s t-test: p = < 0.01, n = 16). However, the surface area of the sMEA electrode was an order of magnitude greater than that of the rigid tungsten microelectrode (7,854 μm² vs. maximum of 400 μm², assuming total electrode tip contact with the cord surface). As a result, the charge density required by the sMEA to elicit a minimum CAP (972 ± 696 μC/cm²) was significantly less than that required for the rigid tungsten microelectrode (5,910 ± 3039 μC/cm²; paired Student’s t-test: p = < 0.01).
SPATIAL STIMULUS RESOLUTION
The spatial resolution of single-site sMEA stimulation was measured by evoking CAPs on surface spinal tracts and measuring the progressive bilateral reductions in CAP amplitude with orthogonal (lateral) distance from the site of maximal activation (Figure 2). For the 13 spinal cords tested, the evoked CAPs were recorded at orthogonally incremented locations from the site of maximal activation, approximately eight spinal segments caudal to the stimulation site. The sMEA and rigid tungsten electrodes were used to stimulate the cord surface at incrementing multiples of threshold (T–2T, in 0.2T increments).

Compound action potential magnitude die-off was plotted as a function of lateral distance from site of maximal spinal tract activation in order to compare the stimulus resolution properties of both electrode types (Figure 3A). Paired t-tests failed to reveal significant differences in the sMEAs vs. rigid microelectrode’s evoked CAPs’ stimulus resolution. For both array and rigid electrode, increasing the stimulus amplitude increased the magnitude of CAP responses in a graded manner. These gradations appeared to be more evenly spaced in the case of rigid microelectrode stimulation, which may indicate a greater consistency in their electrode-to-axon distance due to rigid microelectrode flattening of the interfaced cord.

To determine overall resolution of lateral axonal tract activation, the CAP responses evoked by sMEA and rigid tungsten electrode stimulation at 1–2T ranges (0.2T increments) were magnitude-normalized such that all maximal responses were equal to 1 (Figure 3B). The magnitudes of evoked CAPs for the sMEA and the rigid electrode both decreased to below 50% within approximately 100 μm of either side of the site of maximal spinal tract activation.

AXONAL THRESHOLD SELECTIVITY
We further tested the sMEAs ability to activate spinal axons in a controlled manner by assaying recruitment of higher-threshold axons as a function of electrode proximity (n = 13). Slower-conducting (smaller diameter) axons are known to be recruited at incrementally higher thresholds than faster-conducting (larger diameter) axons (Hursh, 1939); in this way, axonal conduction-velocity selectivity can help characterize this technology’s ability to stimulate white matter axons in inverse order of their axon diameter.

Minimum conduction velocities (m/s) for the evoked CAPs were calculated using the measured distance between stimulating and recording electrodes and the latency of the slowest evoked responses at offset (Figure 4A). Because of the reduced temperature’s slowing effect on axonal conduction-velocity, the conduction velocities evoked by T and 2T stimulation using our present in vitro setup were not directly comparable to ranges reported in vivo at body temperature for Aβ fibers (16–36 m/s).

FIGURE 3 | Spatial stimulus selectivity of sMEA vs. tungsten microelectrode. The magnitude of evoked compound action potentials (CAPs; μV × ms) on the central-dorsal column surface was measured and then plotted over lateral distance from site of maximal CAP response (i.e., the longitudinal location of the stimulating electrodes; n = 13). CAP response magnitudes were calculated using the rectified, integrated, and baseline-subtracted responses for a fixed time window following the stimulus artifact. These values were then averaged over 10 trials for a given recording site on the dorsal column. Standard error bars are shown. (A) Comparison of CAP magnitude over lateral distance from site of maximal activation with sMEA (top) and tungsten electrodes (bottom) at multiple stimulus intensities. (B) Magnitude-normalized comparison of lateral spread of axonal tract activation for sMEA and tungsten electrodes. The magnitudes of evoked CAPs for the MEA electrode pair and the rigid electrode each decreased to below 50% within approximately 100 μm of either side of the site of maximal activation.
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FIGURE 4 | Recruitment of slower-conducting axonal fibers: sMEA vs. tungsten microelectrode. Minimum conduction velocities were calculated for dorsal column CAPs evoked by single-site sMEA and tungsten microelectrode stimulation at threshold (T; n = 13). (A) Example recordings of sMEA-evoked CAPs. The upper CAP was recorded on the longitudinal site of maximal recorded response (i.e., site of maximal activation) and the lower CAP was recorded on a white matter tract 100 μm lateral to the site of maximal activation. Arrows indicate onset of stimulus artifact, and the vertical lines surrounding the responses indicate the onset and offset of the evoked CAP. (B) Minimum conduction velocities for the CAPs evoked by threshold current central–dorsal column stimulation, sMEA vs. tungsten microelectrode.

nor for nociception-encoding Aδ and C fibers (4–16 and 0.5–2 m/s, respectively, as measured in the rat; Suh et al., 1984; Chung and Coggeshall, 1987).

While studies using rat tail motor neuron axons have shown that lower temperatures slow axon conduction in a linear manner that would place axonal conduction-velocity ranges in our experiments within range of Aβ fibers (Miyoshi and Goto, 1973), such extrapolations could not be applied definitively to the tested spinal tracts. This was due not just to the differences in temperature, but also the significant difference in the ratio of unmyelinated-to-myelinated fibers in the juvenile vs. adult rat dorsal column (Chung and Coggeshall, 1987).

Results comparing minimum conduction velocities of evoked CAPs demonstrated that both electrode types were capable of graded recruitment of slower-conducting fibers as a function of lateral proximity to the stimulus site (Figure 4B). Two-tailed, paired t-tests comparing these results showed that there was no statistically significant difference between the sMEA and tungsten electrode concerning ability to recruit slower axons as a function of electrode proximity.

At stimulus values of twice threshold current, the sMEA and tungsten electrode both evoked significantly slower axons (paired t-test, p = 0.040 for sMEA and p = 0.070 for tungsten electrode, minimum conduction velocities of 0.270 ± 0.070 and 0.254 ± 0.070 m/s, respectively. These results demonstrate that sMEA and tungsten electrodes are similarly capable of modulating recruitment of slower-conducting fibers based on electrode position and stimulus amplitude.

DUAL-SITE STIMULUS PRECISION

For a subset of experiments (n = 3 cords), an additional pair of electrodes on the sMEA was used to stimulate a second site on the dorsal column surface. For each of the sites (approximately 400 μm distance between electrode pairs), stimuli were applied independently at single-pulse threshold current levels (biphasic pulses, 500 μs duration). As in the single-site
selectivity experiments, evoked CAPs were recorded at orthogonally incremented locations approximately eight segments caudal to the site of maximal activation. For all three cords on which the sMEA electrodes were placed on the central–dorsal column, distinct peaks in CAP response magnitude were found (Figure 5). The spacing of these peaks was approximately 400 μm in two out of the three cords tested, and less than 400 μm for the third cord. These results showed that adjacent electrodes on the sMEA were capable of selectively recruiting distinct spinal tracts along the central–dorsal column of the in vitro spinal cord.

**MOTONEURON RECRUITMENT**

Experiments were conducted to determine if sMEA-based spinal tract surface stimulation was capable of activating motoneuron population responses associated with hindlimb flexor and extensor movement. Electrodes on the sMEA were placed on the lateral aspect of the T12 segment (DLF, LF, and VLF tracts). These regions are known to carry axons that subserve movement coordination (Davidoff, 1983; Jordan and Schmidt, 2002), and pulse-train stimulation of the in vitro VLF has been shown to elicit locomotor-like activity (Magnuson and Trinder, 1997; Antonino-Green et al., 2002). Adjacent electrodes on the sMEA were configured for bipolar stimulation of the three surface locations such that the distance between DLF and LF stimulation was approximately 400 μm and the distance between LF and VLF stimulation was approximately 200 μm (Figure 6A). Single-pulse current stimuli were delivered at minimum values required to evoke a visible ventral root response, which was measured at the second and fifth lumbar levels. In addition, CAPs evoked on the surface of the VLF and DLF were recorded at the sixth lumbar level to monitor axon tract recruitment. LF and VLF, but not DLF stimulation, elicited combined ventral root L2 and L5 responses (Figure 6B, n = 2). Ventral root L5 responses were of greater magnitude and longer delay-to-onset when compared with ventral root L2 responses.

To further determine whether surface stimulation of the spinal cord using sMEA electrodes was capable of accessing hindlimb flexor and extensor related motoneuron populations, the sMEA was used to stimulate the T12 VLF using a train of pulses (Figure 6C, n = 1). A pair of sMEA electrodes (bipolar current stimulation configuration) was placed on the VLF at thoracic level.
FIGURE 6 | Hindlimb flexor- and extensor related responses to sMEA electrode stimulation at three distinct surface sites. Adjacent sMEA electrode pairs were used to deliver separate stimuli to the T12 DLF, LF, and VLF (n = 2, eight stimulus trials per site, representative traces from one trial per site shown). (A) Electrodes were configured for bipolar stimulation of the three surface locations. (B) Response recordings were made at ventral roots L2 and L5 (as well as at the DLF and VLF L6 surfaces, in order to determine the ability of the sMEA to activate distinct motor patterns. Numerals (i,ii) indicate separate spinal cord preparations. (C) A sMEA electrode pair was placed on the VLF at thoracic level 12 (T12) and responses were recorded at ventral root L2, ventral root L5, the L6 DLF surface, and the L6 VLF surface. At right, a longer-duration tracing shows ventral root L5’s prolonged activity.

A total of two sMEAs were used for the 20+ in vitro, above-mentioned experiments. These sMEAs were found to be mechanically durable and re-usable without signs of delamination or decomposition. For the several months over which these sMEAs were repeatedly manipulated, they were constantly soaking in either a CSF or deionized water (in between experiments). The temporal consistency of the sMEAs’ electrical properties was not formally tested; however, the required currents for evoking CAPs on in vitro cords displayed consistent ranges across months of experiments: for sMEA #1, the beginning and ending experiments required 300 and 200 µA (total range: 200–300 µA with a 20-µA outlier); for sMEA #2, the beginning and ending experiments required 50 and 150 µA (total range: 50–250 µA). A single, charge-balanced pulse of 500 µs duration was used to find each threshold current value.

DISCUSSION
The sMEA’s ability to precisely stimulate spinal surface axons was evaluated by comparing its activation profile to that of rigid (tungsten) bipolar microelectrodes pressed directly onto the cord. The regional selectivity of the sMEA was similar to that of the rigid electrode; however, the rigid electrode displayed a slightly more even gradation. While the reasons for this difference are undetermined, they may be related to the way in which the rigid microelectrode...
as the presented experiments did not incorporate pharmacologic threshold stimuli were used to recruit only the largest and most (Gustafsson and Jankowska, 1976; Gaunt et al., 2006) and only low are activated at lower thresholds than their associated cell bodies activated by threshold stimulation at the surface. This likely, as axons decays is that gray matter deep to the dorsal column was not activated dorsal column axons propagate their signals caudally with a focused current delivery relies on the assumption that the activation. An MEA electrode stimulus resolution of 200 μm is roughly equivalent to anywhere from 27 to 270 adult rat dorsal column axon widths, assuming that at least 2/3 of the axons in the dorsal column are myelinated Aα, Aβ, or propriospinal fibers with diameters of 1–10 μm, and that the remaining 1/3 of axons are Aδ, C, or unmyelinated propriospinal fibers with diameters of 0.2–1 μm (Russell, 1980; Chung and Coggeshall, 1983; Patterson et al., 1989).

Our conclusion that the observed CAP decays were the result of a focused current delivery relies on the assumption that the activated dorsal column axons propagate their signals caudally with negligible loss or (postsynaptic) addition of activity. This assumption is supported by the fact that the dorsal column (aside from the deepest white matter) contains mostly primary afferents and postsynaptic dorsal column tract cells, which can be activated both antidromically and orthodromically (Willis and Coggeshall, 2004). As the presented experiments did not incorporate pharmacologic block of synaptic transmission, the contribution of synthetically recruited spiking of postsynaptic dorsal column tract cells cannot be excluded. Nonetheless, such recruitment following a single stimulus is unlikely to produce a suprathreshold synaptic response in these neurons (Jankowska et al., 1979).

Another assumption made in interpreting the observed CAP decays is that gray matter deep to the dorsal column was not activated by threshold stimulation at the surface. This likely, as axons are activated at lower thresholds than their associated cell bodies (Gustafsson and Jankowska, 1976; Gaunt et al., 2006) and only low threshold stimuli were used to recruit only the largest and most easily recruited axons.

A comparison of minimum CAP conduction velocities evoked by the sMEA and tungsten microelectrode stimulation demonstrated a progressively reduced recruitment of higher-threshold (slower-conducting, smaller diameter) fibers as a function of lateral distance to the stimulating electrode site. This suggests that both the sMEA and rigid tungsten electrodes are in adequate proximity to the cord surface to recruit axons in a manner dependent primarily on axonal activation properties. A significant limitation to our analysis of recruited axonal tracts was loss of faster components of the evoked responses in stimulus artifact. In future studies, stimulus artifact elimination technologies (Blum et al., 2007) can be used to circumvent this limitation and also enable closer-proximity response recordings.

The sMEA was used to stimulate the lower thoracic (T12) dorsalateral, lateral, and ventrolateral funiculi, which are regions that have been shown to carry axons that subserve movement coordination (Chung and Coggeshall, 1983; Jordan and Schmidt, 2002). Threshold pulse stimulation of the LF and VLF, but not the DLF, was capable of eliciting combined ventral root L2 and L5 responses. Pulse-train stimulation of the T12 VLF evoked transient response bursts at ventral root L2 and the VLF surface, and produced long-lasting excitability at ventral roots L2, L5, and surface DLF and VLF. It remains undetermined how much of this post-stimulus excitability was noise- and/or damage-related, and how much correlates with meaningful motor activity.

There are fundamental limitations to the correlation of evoked ventral root potentials recorded from in vitro, hemisected spinal cords to coordinated motor output. While the in vitro experiments performed here were appropriate for initial assessment of our sMEA’s capabilities, a more intact preparation is required to determine whether sMEA stimulation of the spinal cord surface is capable of activating motor activity.

As with any device designed for implantation, biocompatibility characterization of the sMEA is an essential parallel step for in vivo development of neural prosthetics (Green et al., 2010; McCreery et al., 2004). In addition, further experiments must be performed to evaluate the sMEA’s potential for in vivo spinal cord mechanical damage.
One proposed sMEA configuration for these initial in vivo implantation experiments is illustrated in Figure 7. Using a dorsal laminectomy approach, the sMEA could be wrapped around the circumference of the cord segment in between root pairs. This approach would test the sMEA’s potential ability to access multiple tracts on the anterolateral surface, which is presently a speculated advantage of this technology. The damage incurred by initial implantation – as well as long-term implantation both with and without FES – could then be evaluated.

To support the proposed, next-step implantation experiments, interconnect technology has been developed for the sMEA that provides stable electro-mechanical interfacing with external stimulators (Guo and DeWeerth, 2009a). Stabilization of the sMEA and wiring can be accomplished using spinal processes as anchoring points.

CONCLUSION
We have constructed specialized, elastomer-substrate sMEA technology that enables close-proximity, multi-point electrical interfacing to the spinal cord surface. The stimulus precision of the elastomer-substrate sMEA, when compared to rigid (tungsten) bipolar electrodes pressed directly onto the spinal cord surface, was similar in both regional selectivity and gradation. This gradation was demonstrated both by incrementally increased current amplitudes as well as by relative (lateral) distance of the recruited axons from the stimulus site.

The regional selectivity of the sMEA was quantified by measuring the axonal activation magnitude as a function of lateral (orthogonal) distance of the axons to the stimulus site, when stimulated at threshold values. The magnitudes of evoked CAPs for the sMEA electrode pair and the rigid electrode both decreased to below 50% within approximately 100 μm of either lateral (orthogonal) distance side of the site of maximal activation. When using the sMEA to stimulate central–dorsal column surface sites 200 mm apart, the sMEA recruited non-overlapping parallel tracts.

Compared to the tungsten electrode, the sMEA demonstrated a lower minimum charge density required to activate dorsal column surface tracts. While these results are preliminary, they serve as important starting points for demonstrating the relative utility of the sMEA for axonal stimulation protocols.

The potential for the sMEA to incur harm to its interfaced neural tissue – via charge density, mechanical shear/compression, or otherwise – remains largely undetermined, and will require further investigation. If in vivo testing proves that the implanted sMEA is capable of minimal-damage spinal tract stimulation, and if this in vivo stimulation provides similar precision to that of the presented in vitro studies, this sMEA might be a powerful enabling technology for multiple neuro-interfacing protocols.

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