The functional properties of synapses made by regenerated axons across spinal cord lesion sites in lamprey

Abstract

While the anatomical properties of regenerated axons across spinal cord lesion sites have been studied extensively, little is known of how the functional properties of regenerated synapses compared to those in unlesioned animals. This study aims to compare the properties of synapses made by regenerated axons with unlesioned axons using the lamprey, a model system for spinal injury research, in which functional locomotor recovery after spinal cord lesions is associated with axonal regeneration across the lesion site. Regenerated synapses below the lesion site did not differ from synapses from unlesioned axons with respect to the amplitude and duration of single excitatory postsynaptic potentials. They also showed the same activity-dependent depression over spike trains. However, regenerated synapses did differ from unlesioned synapses as the estimated number of synaptic vesicles was greater and there was evidence for increased postsynaptic quantal amplitude. For axons above the lesion site, the amplitude and duration of single synaptic inputs also did not differ significantly from unlesioned animals. However, in this case, there was evidence of a reduction in release probability and inputs facilitated rather than depressed over spike trains. Synaptic inputs from single regenerated axons below the lesion site thus do not increase in amplitude to compensate for the reduced number of descending axons after functional recovery. However, the postsynaptic input was maintained at the unlesioned level using different synaptic properties. Conversely, the facilitation from the same initial amplitude above the lesion site made the synaptic input over spike trains functionally stronger. This may help to increase proprioceptive activity across the lesion site to compensate for the lesion-induced reduction in supraspinal inputs. The animal experiments were approved by the Animal Ethics Committee of Cambridge University.

Key Words: electrophysiology; lamprey; plasticity; regeneration; reticulospinal axon; spinal cord; spinal injury; synapse

Introduction

Regeneration remains the dominant approach in attempts to restore function after spinal cord injury (Ramer et al., 2014). The anatomical properties of regenerated axons (their number and projections) have been studied extensively in various systems, but their physiological properties have received relatively little attention. Given that these properties will determine the functional effect of any regenerated inputs it is important that they are considered, not least because this may help to explain the variable relationship between regeneration and recovery (Steward et al., 2012).

The small size of regenerated axons makes them difficult to study in the mammalian spinal cord. This study has used the lamprey, a model system for studying axonal regeneration and functional recovery after spinal cord lesions. Analyses in lamprey typically focus on the larger Müller reticulospinal axons as these allow stable intracellular recordings to be made from identified axons (Wood and Cohen, 1981; Hall et al., 1989; Mchale et al., 1995; Oliphint et al., 2010; Zhang et al., 2011). These axons regenerate to make functional synaptic connections below lesion sites (Mackler and Selzer, 1987).

While regeneration is associated with functional recovery in the lamprey, regeneration is never complete (it ranges from 0–70%; McClellan, 1994) and regenerated axons project to ectopic locations (Wood and Cohen, 1981): regeneration thus does not ‘repair’ the spinal cord to its original state. Given that not all of the axons regenerate, for regeneration alone to equal recovery either means that in the unlesioned animal there are more axons than needed (i.e. there is a redundancy), or that there are other changes that allow the reduced number of axons to allow normal function despite the reduction in axon number after recovery. The latter seems more likely given that redundant descending inputs would be costly to develop and maintain, and given the evidence from the lamprey to the human spinal cord of functional changes below spinal cord lesion sites (Grasso et al., 2004; D'Amico et al., 2014; Parker, 2017).

Morphological analyses have shown that regenerated Müller axons make fewer synapses below the lesion site and that these synapses contain fewer vesicles and have smaller active zones than axons from unlesioned animals (Oliphint et al., 2010). Despite this, Müller axons evoke postsynaptic inputs that match those in unlesioned animals (Cooke and Parker, 2009). While this is unexpected given the morphological changes shown by Oliphint et al. (2010), there are compensations that can allow single axons to evoke the same postsynaptic input (Parker, 2017). While the large uniquely identifiable Müller axons are convenient targets for analyses, lesioning the medial column where they project does not abolish locomotion in functionally recovered animals. Lateral column axons, which are smaller and more numerous, seem to be more important for locomotor recovery (McClellan, 1990; Chen et al., 2017).

From the Contents

Introduction 2272
Materials and Methods 2273
Results 2273
Discussion 2275

How to cite this article: Parker D (2022) The functional properties of synapses made by regenerated axons across spinal cord lesion sites in lamprey. Neural Regen Res 17(10):2272-2277.
This analysis has focused on lateral tract axons, with the specific aim of determining how the properties of synapses made by regenerated axons compare to those made by axons in unlesioned animals, and the mechanisms of corresponding plasticity. The results suggest that regenerated synaptic inputs to motor neurons below the lesion site match those in unlesioned animals, but that the input is evoked using different release properties. However, inputs to motor neurons from axons above the lesion site are altered to become functionally stronger than those in unlesioned animals.

Materials and Methods

Animals and lesioning

Juvenile male and female adult lampreys (Petromyzon marinus) between 100–130 mm (n = 84) were purchased from commercial suppliers (Acme Lamprey, Hamburg, UK; Salmon-Netz, Hamburg, Germany). Animals were maintained in aquarium at room temperature. To lesion the spinal cord, animals were anesthetized by immersion in MS-222 (Sigma, Welwyn Garden City, UK; 300 mg/mL, pH adjusted to 7.4). The spinal cord was exposed by making a dorsal incision approximately 1 cm below the last gill and was then transected using iris scissors. The incision site was repaired with Vet Bond Tissue adhesive (3M Animal Care Products). Following transection, animals were kept at 21°C for 8–10 weeks (Cohen et al., 1999). Animals were examined after 8–10 weeks as axons were typically examined 2–5 months post-lesion (Paech and Parker, 1997). There are changes in electrophysiological properties at later time points (Becker and Parker, 2019), but later time points were not examined here. All experiments were conducted under the license of the UK Home Office (Animals Scientific Procedures Act 1986; Project license PPI; 80/2417 (03/2011)) and with the approval of the Animal Ethics Committee of Cambridge University.

Electrophysiological recordings

Video and electromyogram recordings were made from randomly assigned control animals and animals 8–10 weeks after lesioning to assess the degree of recovery. The animals used here were assessed to have either recovered normal locomotion (good recovery) or failed to show any recovery (poor recovery); see Hoffman and Parker, 2011 for details. The spinal cord was then isolated for intracellular recordings by anesthetizing animals in MS-222 and removing the spinal cord and notochord. The spinal cord was isolated in the notochord and placed ventral side up in a Sylgard-lined Perspex experimental chamber where it was superfused with Ringer containing (in mM): 138 NaCl, 2.1 KCl, 1.8 CaCl₂, 1.2 MgCl₂, 4 glucose, 2 HEPES, 0.5 L-glutamine. The pH was adjusted to 7.4 with NaOH, and the osmolarity was 300 osm. A single hippocampal bipolar electrode was inserted into the spinal cord. Paired recordings were made from axons in the lateral tract and motor neurons using thin-walled micropipettes filled with 3 M potassium acetate and 1 M potassium chloride. The lateral tract in the lamprey spinal cord also contains axons from the giant interneurons which relay sensory input to the brainstem and reticulospinal neurons. Giant interneuron axons are located below the lateral tract, and the spinal cord is divided into the retrocerebral complex, which is the ventral part of the spinal cord and the neuropil in the ventral part of the spinal cord. The electrode was identified by its reliability and latency. Motor neurons were recorded by identifying antidromic extracellular spikes in response to electrical stimulation of ventral root following current injection into the somata. Axons were identified by recording antidromic extracellular spikes on the rostral end of the spinal cord following their stimulation, and by the absence of a slow afterhyperpolarization following the action potential. Motor and reticulo-spinal neurons were identified by their reliability and latency (i.e. the input did not fail or the latency change over the spike train) following presynaptic stimulation at 20 Hz (Bryan and Pentreath, 1976) to minimize potential differences due to the location of cells in different regions of the spinal cord. Experiments were performed in the rostral trunk region of the lamprey spinal cord (Fig. 1A–C). However, changes in the EPSP levels the connection was classified as unchanged. Paired-pulse plasticity was expressed as EPSP/EPSP, and plasticity over different regions of the spike train as EPSPθ/EPSP. Single, low frequency-evoked EPSPs were used to measure EPSP rise times and half-widths.

Statistical analysis

Values are presented as the mean ± SEM. Statistical significance was examined using either a paired or independent samples t-tests, or one-way analysis of variance (ANOVA), and differences in the proportion of effects by the chi-square test. When a one-way analysis of variance was used, a Tukey’s test was used for post-hoc analysis of differences between group means. All statistical tests refer to the number of connections examined (each connection reflects a new axon and motor neuron). Linear regression was also used to determine the correlations between various factors. n refers to the number of paired recordings. Numbers are given for testing in a single spinal cord. Statistical tests were only performed when the sample size allowed a power of at least 0.8: where a P-value was not reported the sample size was below that needed for this power.

Results

Basic synaptic properties

In lesioned animals, the EPSP amplitude, rise time, and half-width above (n = 19 paired recordings) and below (n = 15 paired recordings) the lesion site did not differ significantly from unlesioned values (n = 62 paired recordings, P > 0.05). However, paired-pulse plasticity was stronger than those in unlesioned animals below the lesion site than for EPSPs in unlesioned animals or below the lesion site (P < 0.05; Figure 1B), an effect that can be associated with a lower release probability (Zucker and Regehr, 2002). EPSPs can also show an electrical component or be exclusively chemical, so a reduction in the overall column Müller axons suggest a reduction of electrical connections after injury (Wood and Cohen, 1981), but the proportion of electrical versus non-electrical synapses (number of connections with an electrical component/total number of connections) here did not differ significantly between unlesioned and lesioned animals above and below the lesion site (data not shown; P > 0.05). There were no changes in axonal action potential properties that could influence transmitter release (e.g. spatial dispersion; Parker et al., 1997; data not shown). These changes have been reported in regenerated axons in lamprey, but they have returned to unlesioned values at the time the analyses here were performed (McClellan et al., 2008).

Activity-dependant plasticity

Facilitation was the usual paired-pulse effect (i.e. the 2nd EPSP compared to the 1st) to 20 Hz stimulation in both lesioned and unlesioned animals. Plasticity over spike trains was assessed from the effect over the 11th and 20th EPSPs in response to 20 Hz stimulation (effects typically plateau over this part of the spike train; Figure 1D). In unlesioned axons, the average effect was a depression to approximately 60% of the initial EPSP amplitude. However, individual connections varied (Brodin et al., 1994). In unlesioned animals, 30 (of 62) connections depressed, 14 connections facilitated, 2 connections were biphasic, and 16 were unchanged (i.e. showed no activity-dependent plasticity over the spike train). In lesioned axons, below the lesion site (n = 10 paired recordings) the lesion site was again the usual effect (10 connections depressed, 2 facilitated, and 3 were unchanged). In contrast, above the lesion site facilitation was the most common effect over spike trains (Figure 1D). 12 connections facilitated, 4 connections depressed, and 3 were unchanged. When the proportions of the different types of activity-dependant plasticity did not differ in unlesioned and below lesion spinal cords, above the lesion site facilitating connections were significantly more common (P < 0.05).

When connections were separated by the type of activity-dependant plasticity, the degree of facilitation did not differ in unlesioned and above lesion spinal cords: facilitation developed to approximately 1500% over Train1–21, in all cases (Figure 1E and G), and did not differ significantly in unlesioned and above lesion axons (P > 0.05; the number of facilitating connections below the lesion site was too small to compare statistically). Depression also did not differ in unlesioned and above and below lesion spinal cords (Figure 1F and G), the magnitude not differing significantly in unlesioned and below lesion axons (P > 0.05; the number of depressing connections above the lesion site was too small to compare statistically).

In unlesioned animals there was no significant correlation between the initial EPSP amplitude and the paired-pulse (PP) ratio (P = 0.45), or the change in the paired-pulse (PP) ratio (P = 0.62; Figure 2A) in all cases (Figure 1E and G), and did not differ significantly in unlesioned and above lesion axons (P > 0.05; the number of facilitating connections below the lesion site was too small to compare statistically). In unlesioned and above lesion spinal cords, the EPSP amplitude thus did not predict the type of activity-dependant plasticity at these connections (a similar effect occurs at other excitatory synapses in the lamprey spinal cord). However, below the lesion site, while there was no significant correlation with the initial excitatory synapse EPSP amplitude above a paired-pulse ratio of 0.16, there were significant negative relationships with the Train1–21 (r = 0.23; data not shown) and Train1–21 (r = 0.29; Figure 2A) in all cases (Figure 1E and G), and did not differ significantly in unlesioned and above lesion axons (P > 0.05; the number of depressing connections above the lesion site was too small to compare statistically).
Larger initial EPSPs thus evoked greater depression, an effect that can be associated with release probability-dependent depression due to depletion or greater postsynaptic desensitization (Zucker and Regehr, 2002). The number of vesicles was estimated from the model of Wang and Zucker (1998):
\[ N_{min}(\tau(V_r, \tau_0)/q(V_r, V_{min})) \]
\( V_r \) is the initial EPSP amplitude, \( \tau \) the inverse rate constant of EPSP decay (expressed as the number of presynaptic spikes needed for the EPSP to fall to 1/e of the initial value), \( q \) the mean quantal amplitude (assumed to be 0.1 mV), and \( V_{min} \), the EPSP amplitude at the plateau level of depression. Because this analysis depends on the rate of depression (Wang and Zucker, 1998; Schneggenburger et al., 1999; Millar et al., 2002), it could only be applied to unlesioned and below lesion axons as the number of depressions was too small above the lesion site. The analysis also required that the input decreased to 1/e of the initial value over the spike train. This often does not happen even over longer spike trains (> 50 EPSPs), probably due to the concomitant development of activity-dependent replenishment of the releasable pool over the spike train (Parker, 2000). As a result, the extrapolated exponential depression calculated from the initial depressing EPSPs of the train was used to determine \( \tau \) and \( V_r \) to allow approximation of the vesicle pool. Any influence of replenishment would bias the rate of rundown and give an over-estimation of the initial vesicle pool. This would apply equally to unlesioned and lesioned synapses unless there was a difference in the rate of replenishment between unlesioned and lesioned synapses (Parker, 2000), an effect that would be functionally equivalent to an increase or decrease in vesicle numbers if the rate of replenishment was faster or slower, respectively. In unlesioned animals, the mean number of vesicles was within the range of direct vesicle counts of 100–500 vesicles in the middle section of the vesicle pool from Müller axons in lamprey (Gustafsson et al., 2002). However, below the lesion site the estimated mean number of vesicles was almost 50% greater (P < 0.05; Figure 3A and B).

Presynaptic or postsynaptic parameters were assessed using the coefficient of variation (CV') to compare changes in mean and variance of the initial and final EPSPs in spike trains (Brock et al., 2020). The analysis assumes that parallel changes in synaptic amplitude and CV' reflect presynaptic effects, while synaptic changes without a change in CV' reflect postsynaptic effects (Faber and Korn, 1991). As the method is sensitive to noise (Brock et al., 2020), only connections with no spontaneous synaptic activity were analyzed. In unlesioned animals (Figure 4A), 3 of 6 facilitating connections had CV' changes associated with presynaptic effects, 3 with postsynaptic or mixed pre and postsynaptic effects: for depressing connections 17 had CV' changes consistent with a presynaptic effect, and 11 with postsynaptic/mixed effects. Above the lesion site, 4 of 8 facilitating connections had changes consistent with a presynaptic effect, 4 with mixed or postsynaptic effects (Figure 4B). Below the lesion site (Figure 4C), 4 connections showed CV' changes consistent with presynaptic effects and 2 with mixed or postsynaptic effects. The proportions of presynaptic and postsynaptic/mixed effects did not differ in the unlesioned or lesioned spinal cord (P > 0.05).

As the CV' method relies on certain conditions and assumptions to identify a pre or postsynaptic locus of change (Faber and Korn, 1991; Brock et al., 2020), release properties were also examined using the variance-mean method (Clements and Silver, 2000). This examines presynaptic potentials in normal, high, and low calcium Ringer. Unless the release probability is low (< 0.3), the relationship of the EPSP variance and mean in different calcium Ringers results in a parabolic relationship from which release parameters can be estimated (Clements and Silver, 2000 for details): \( N_{min} \), the number of release sites is determined from the width of the parabola; \( p \), the probability of release from the curvature; and \( q \), the postsynaptic quantal amplitude from the initial slope (Figure 5A).

The analysis again only includes connections that were stable throughout the experiment and in which EPSP amplitudes could be measured unequivocally (i.e. without contamination by an electrical component to the EPSP or spontaneous synaptic inputs). This was a significant issue, as the lateral tract axons are relatively small and recordings stable enough to allow assumption of the different calcium Ringer solutions to be examined were not common, and low calcium Ringer often increased background synaptic inputs, presumably through a reduction of surface screening (Piccolino and Pignatelli, 1996). This meant that the n number of fully analyzed connections is small (this is not uncommon for variance-mean analyses; see for example, Mitchell and Silver,
Differences in synaptic properties in animals that recovered well or poorly. The absence of parabolic relationship between the variance and mean (n = 4; Figure 5A). This is indicative of a low release probability (< 0.3; Clements and Silver, 2000), and is consistent with the higher PP ratio and the development of facilitation above the lesion site (Figure 1D). The absence of the parabolic relationship prevented estimation of N_{st}, and P, but the slope of the relationship was not increased compared to unlesioned animals, suggesting against a change in Q above the lesion site (Clements and Silver, 2000).


discussion

This analysis has examined the properties of regenerated axons in the lamprey. Synaptic inputs from lateral column reticulospinal axons below the lesion site matched those from unlesioned axons in terms of their amplitude and activity-dependent plasticity. However, despite the same output being generated by these inputs in unlesioned animals, the properties of connections by axons above the lesion site differed from those in the unlesioned spinal cord as they showed facilitation that developed from the same initial EPSP amplitude and would thus be functionally stronger.

Regeneration remains the dominant approach in spinal cord injury research. Analyses focus on the anatomical properties of regenerated axons (e.g. their number, location, and how far they regenerate below the lesion site), and the correlation of regeneration with recovery. However, it is not enough just for axons to regenerate; functional recovery requires that they make appropriate connections with targets below the lesion site. Differences in the release properties that determine the features of these connections could thus contribute to the variable influence of regeneration on recovery seen in experimental model systems like the lamprey (Parker, 2017) and clinical trials of regeneration (Steward et al., 2012).

It is difficult to examine synaptic properties directly to compare effects in unlesioned and lesioned spinal cords. A range of approaches was used here, including direct measurements of basic synaptic properties, the estimation of vesicle numbers, comparison of presynaptic versus postsynaptic mechanisms (CV^2), and comparison of release properties (variance-mean analysis). While these are commonly used approaches, they have limitations and determining the mechanisms of release at central synapses is still difficult (Lanore and Silver, 2016; Pulido and Marty, 2017). Changes in postsynaptic response can be assessed simply and directly by comparing the response to exogenously applied glutamate in functionally isolated cells (e.g. in the presence of TTX; Parker et al., 1997), but this only works for comparisons of the same cell under different conditions (e.g. before and after application of a neuromodulator), not for comparing effects in different spinal cords. The variance-mean approach makes fewer assumptions than conventional quantal mechanisms (Clements and Silver, 2000), including the CV^2 analysis (Faber and Korn, 1991), and has been applied at a range of central synapses (Lanore and Silver, 2016). However, the analysis needs relatively long-term stable recordings from relatively small presynaptic axons to allow the changes in Ringer calcium levels and evoked EPSPs that are not contaminated by spontaneous background inputs, requirements that limit the sample size. However, the results of this analysis should be consistent with each other, providing support for the general conclusions.

The same amplitude and activity-dependent properties of regenerated axons to motor neurons below the lesion site as unlesioned animals suggest against an increase in the synaptic strength of individual axons to compensate for the overall reduction of the descending input after recovery from a spinal cord lesion (fully recovered animals showed a 30% reduction in the number of axons; Cohen et al., 1988; McClellan, 1990). However, the same synaptic effect is evoked by different synaptic mechanisms. The different
relationship between the initial EPSP amplitude and the paired-pulse ratio in below lesion synapses suggests a change in presynaptic release (that larger initial EPSPs evoked greater depression may reflect a larger quantal content with a concomitant increase in vesicle depletion and depression or greater probability of release: see Bevan and Parker, 2004), while the variance-mean analysis suggests an increase in the postsynaptic response (quantal amplitude, q) below the lesion site, an effect that is consistent with previously identified sub-lesional functional changes (Parker, 2017). This effect could scale up regenerated inputs to allow them to match the properties of unlesioned synapses despite the reduced number of functional sites suggested by the reduced N_p, from the variance-mean analysis, and the potential for a reduced number of synaptic contacts from morphological analyses of regenerated Müller axons below the lesion site (Oliphant et al., 2010). The increase in the estimated vesicle numbers below the lesion site could also help strengthen each contact. The combination of these presynaptic and postsynaptic mechanisms below the lesion site may account for the lack of a difference in the proportions of putative presynaptic and postsynaptic in the CV-2 analysis.

Why should different release properties be used to generate the same synaptic output below the lesion site? The maintenance of unlesioned properties could obviously reflect the need for inputs of a certain magnitude to effectively activate sub-lesion networks. The difficulties of recovering the original development of these axons and their synaptic contacts in a mature nervous system may account for the smaller and limited number of synaptic contacts below the lesion site (Oliphant et al., 2010). Each contact made by a single axon could be made stronger by increasing the transmitter release probability, but the presynaptic vesicle depletion and depression was also more efficient this could lead to a release probability-dependent increase in depression that would weaken inputs over spike trains (Zucker and Regehr, 2002). Instead of strengthening the connection, an increase in release probability alone would make the input over the spike train, from being initially stronger to being subsequently weaker than unlesioned inputs, depending on how many inputs were evoked. Maintaining high levels of release and the associated need for increased transmitter clearance and replenishment would also invoke a significant energy cost (Howarth et al., 2012), which may be an issue for growing regenerating axons given the compromised state of the lesioned spinal cord. The scaling of the synapse through a postsynaptic increase in quantal amplitude and presynaptic increase in vesicle numbers suggested here could maintain the pre-lesion synaptic amplitude at the unlesioned value despite each axon potentially making fewer synaptic contacts, without the energetic costs of upregulating transmitter release and replenishment mechanisms. The various changes in cellular and synaptic properties seen below the lesion site in various model systems from lamprey to mammals could make the spinal cord below the lesion site more excitable (D’Amico et al., 2014; Parker, 2017). These postsynaptic changes may better compensate for the reduced descending input below lesion sites than a presynaptic modification of the regenerated axons, as postsynaptic cells will be able to modify their response to synaptic inputs depending on the integrated descending input that they receive (Parker and Grillner, 2000). The variance-mean analysis and paired-pulse ratio suggest that the release probability is reduced at synapses between lateral tract axons and locomotor neurons in the lamprey (Bevan and Parker, 2004): this is consistent with the increased PP ratio and facilitation over spike trains in this region. However, the reduction of release probability was not associated with a significant reduction of the initial EPSP amplitude in the spike trains, which would be expected of a reduction in the presynaptic probability, suggesting that other factors are also altered. This could reflect a concomitant increase in q (see Bevan and Parker, 2004 for an example), but the variance-mean and CV^2 analyses suggest against this. It could alternatively reflect an increase in vesicle numbers (Bevan and Parker, 2004): this will need anatomical analyses as the method used here for estimating vesicle numbers is based on the rate of depression and thus cannot be applied to facilitating connections. Irrespective of the mechanism, facilitation without a significant reduction of the initial EPSP amplitude would make the summed input above the lesion site functionally stronger than connections to unlesioned spinal cord or firing of the lesion site, suggesting that other factors are also altered. The compensation that may allow stronger propriospinal activity across the lesion site (Courtine et al., 2008), Regeneration is never complete in lamprey, varying between 0–70% of the unlesioned value (Cohen et al., 1988; McClellan, 1994). That the same functional output can be generated despite a reduction in descending input either suggests that some of the descending input is redundant, which seems unlikely given the resources that would have to be dedicated to developing and maintaining these connections, or that there is some compensation for the reduction. Each regenerated axon could have increased its postsynaptic effect in proportion to the reduction of the total descending input to maintain the same overall sub-lesion synaptic input, but functionally the overall synaptic connections occurred above, not below the lesion site where a compensatory increase would be expected. However, increasing the amplitude of regenerated inputs below the lesion site to maintain the total descending synaptic at the unlesioned value would not necessarily be an effective compensation as it would reflect the specific functional roles of inputs to different spinal cord regions. If only 50% of the axons regenerated, doubling the amplitude of each would lead to the same summed descending input, but doubling inputs to some regions would not necessarily compensate for zero inputs to other regions.

While regeneration remains a major focus of spinal injury research and overcoming the inhibition of regeneration in mammals has been a great success, regeneration alone will not be enough for the recovery of function after spinal cord injury. The functional efficacy of regenerated inputs is not only dependent on the properties of the synaptic connections they make but also on the properties of the neurons and circuits that they connect to (Ullström et al., 1999). There are changes in these properties after spinal cord injury in systems from lampreys to humans (Grasso et al., 2004; Parker, 2017). Understanding the functional consequences of regeneration and to improve functional recovery for spinal cord injury patients will require a focus on how regenerated interact with altered spinal cord properties above and below the lesion site.

Limitations

The analysis of release properties is difficult to determine, and the measures used here are subject to several caveats. A number of different approaches have been used, and while none is ideal, the results from these analyses generally support each other. In addition, the analysis only considers properties at one-time point after the lesion, and while animals typically recover locomotor function at this time, other properties show changes at later times, which suggests that the effects seen here are not necessarily the final changes.

Conclusion

Regenerated synapses below lesion sites in the lamprey show changes in their release properties. These are presumably important to the successful recovery of locomotor function after lesioning. In addition, synapses above the lesion site also show functional changes, which may be a compensation for the reduced descending input below the lesion site.

Author contributions: Study design, experiment implementation, data analysis, and manuscript writing: DP. The author read and approved the final version of the manuscript.

Conflicts of interest: The author declares no conflict of interest.

Author statement: This paper has been posted as a preprint on bioRxiv with doi:10.1101/2021.06.21.449247, which is available from: https://www.biorxiv.org/content/10.1101/2021.06.21.449247v1.full.

Open access statement: This is an open access journal, and articles are distributed under the terms of the Creative Commons AttributionNonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

References

Becker M, Parker D (2015) Changes in functional properties and 5-HT modulation above and below a spinal transection in lamprey. Front Neural Circuits 8:148.

Becker M, Parker D (2019) Time course of functional changes in locomotor and sensory systems after spinal cord lesions in lamprey. J Neurophysiol 121:2323-2335.

Berry M, Pentreath V (1976) Criteria for distinguishing between monosynaptic and polysynaptic transmission. Brain Res 105:1-20.

Bevan S, Parker D (2004) Metaplastic facilitation and ultrastructural changes in synaptic properties are associated with long-term modulation of the lamprey locomotor network. J Neurosci 24:9458-9468.

Brock JA, Thomeaue A, Watanabe A, Li SSY, Sjöström PJ (2020) A practical guide to using CV analysis for determining the locus of synaptic plasticity. Front Synaptic Neurosci 12:11.

Brodin L, Shupliakov O, Pieribone V, Hellgren J, Hill R (1994) The reticulospinal glutamate synapse in lamprey: plasticity and presynaptic variability. J Neurophysiol 72:592-604.

Brock JA, Thomeaue A, Watanabe A, Li SSY, Sjöström PJ (2020) A practical guide to using CV analysis for determining the locus of synaptic plasticity. Front Synaptic Neurosci 12:11.

Chen J, Larmore C, Shifman MI (2017) The expression of chemorepulsive guidance receptors and the regenerative abilities of spinal-projecting neurons after spinal cord injury. Neuroscience 341:95-111.

Clements JD, Silver RA (2000) Unveiling synaptic plasticity: a new graphical and analytical approach. Trends Neurosci 23:105-113.

Cohen AH, Mackler SA, Selzer ME (1988) Behavioral recovery following spinal transection: functional regeneration in the lamprey CNS. Trends Neurosci 11:227-231.

BEVERLY J, McCARTHY F (2016) Criteria for distinguishing between monosynaptic and polysynaptic transmission. Brain Res 105:1-20. This is an open access journal, and articles are distributed under the terms of the Creative Commons AttributionNonCommercial-ShareAlike 4.0 License, which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.
Cohen AH, Kiemel T, Pate V, Blinder J, Guan L (1999) Temperature can alter the function outcome of spinal cord regeneration in larval lampreys. Neuroscience 90:957-965.

Cooke RM, Parker D (2009) Locomotor recovery after spinal cord lesions in the lamprey is associated with functional and ultrastructural changes below lesion sites. J Neurotrauma 26:597-612.

Courtine G, Song B, Roy RR, Zhong H, Herrmann JE, Ao Y, Qi J, Edgerton VR, Sofroniew MV (2008) Recovery of supraspinal control of stepping via indirect propriospinal relay connections after spinal cord injury. Nat Med 14:69-74.

D’Amico JM, Condille EG, Martins KJ, Bennett DJ, Gorassini MA (2014) Recovery of neuronal and network excitability after spinal cord injury and implications for spasticity. Front Integr Neurosci 8:36.

Faber DS, Korn H (1991) Applicability of the coefficient of variation method for analyzing synaptic plasticity. Biophys J 60:1288-1294.

Grasso R, Ivanenko Y, Zago M, Molinari Y, Scivoletto G, Castellano V, Macellari V, Lacquaniti F (2004) Distributed plasticity of locomotor pattern generators in spinal cord injured patients. Brain 127:1019-1034.

Gustafsson JS, Birinyi AS, Crum J, Ellisman M, Brodin L, Shupliakov O (2002) Ultrastructural organization of lamprey reticulospinal synapses in three dimensions. J Comp Neurol 450:167-182.

Hall G, Poulos A, Cohen M (1989) Sprouts emerging from the dendrites of axotomized lamprey central neurons have axon-like ultrastructure. J Neurosci 9:588-599.

Hoffman N, Parker D (2011) Interactive and individual effects of sensory potentiation and region-specific changes in excitability after spinal cord injury. Neuroscience 199:563-576.

Howarth C, Gleeson P, Attwell D (2012) Updated energy budgets for neural computation in the neocortex and cerebellum. J Cereb Blood Flow Metab 32:1222-1232.

Kazama H, Wilson RI (2008) Homeostatic matching and nonlinear amplification at identified central synapses. Neuron 58:401-413.

Lanore F, Silver R (2016) Extracting quantal properties of transmission at central synapses. In: Advanced patch-clamp analysis for neuroscientists (Korngreen A, ed), pp 193-211. New York: Springer.

Lawrence JJ, Haario H, Stone EF (2015) Presynaptic cholinergic neuromodulation alters the temporal dynamics of short-term depression at parvalbumin-positive basket cell synapses from juvenile CA1 mouse hippocampus. J Neurophysiol 113:2408-2419.

Mackler SA, Selzer ME (1987) Specificity of synaptic regeneration in the spinal cord of the larval sea lamprey. J Physiol 388:183-198.

Malagon G, Miki T, Liao I, Neher E, Marty A (2016) Counting vesicular release events reveals binomial release statistics at single glutamatergic synapses. J Neurosci 36:4010-4025.

McClellan AD (1990) Locomotor recovery in spinal-transsected lamprey: Role of functional regeneration of descending axons from brainstem locomotor command neurons. Neuroscience 37:781-798.

McClellan AD (1994) Time course of locomotor recovery and functional regeneration in spinal cord-transsected lamprey: in vitro preparations. J Neurphysiol 72:847-860.

McClellan AD, Kovalenko MO, Benes JA, Schulz DJ (2008) Spinal cord injury induces changes in electrophysiological properties and ion channel expression of reticulospinal neurons in larval lamprey. J Neurosci 28:650-659.

McHale MK, Hall GF, Cohen MJ (1995) Early cytoskeletal changes following injury of giant spinal axons in the lamprey. J Comp Neurol 353:25-37.

Miliar AG, Bradacs H, Charlton MP, Atwood HL (2002) Inverse relationship between release probability and readily releasable vesicles in depressing and facilitating synapses. J Neurosci 22:9661-9667.

Mitchell SJ, Silver RA (2000) Glutamate spillover suppresses inhibition by activating presynaptic mGluRs. Nature 404:498-502.

Oliphant PA, Alieva N, Folders AE, TyeT Ed, Lau BY, Pariseau JS, Cohen AH, Morgan JR (2010) Regenerated synapses in lamprey spinal cord are sparse and small even after functional recovery from injury. J Comp Neurol 518:2854-2872.

Parker D (2000) Activity and calcium-dependent mechanisms maintain reliable interneuron synaptic transmission in a rhythmic neural network. J Neurosci 20:1754-1766.

Parker D (2017) The lesioned spinal cord is a “new” spinal cord: evidence from functional changes after spinal injury in lamprey. Frontiers in Neural Circuits 11:84.

Parker D, Grillner S (2000) Neuronal mechanisms of synaptic and network plasticity in the lamprey spinal cord. Prog Brain Res 125:381-398.

Parker D, Svensson E, Grillner S (1997) Substance P modulates sensory action potentials in the lamprey via a protein kinase C-mediated reduction of a 4-aminopyridine-sensitive potassium conductance. Eur J Neurosci 9:2064-2076.

Piccolino M, Pignatelli A (1996) Calcium-independent synaptic transmission: artifact or fact? Trends Neurosci 19:120-125.

Pulido C, Marty A (2017) Quantal fluctuations in central mammalian synapses: functional role of vesicular docking sites. Physiol Rev 97:1403-1430.

Ramer LM, Ramer MS, Bradbury EJ (2014) Restoring function after spinal cord injury: towards clinical translation of experimental strategies. Lancet Neurol 13:1241-1256.

Rovainen CM (1979) Neurobiology of lampreys. Physiol Rev 59:1007-1077.

Schneeganscher R, Meyer AC, Neher E (1999) Released fraction and total size of a pool of immediately available transmitter quanta at a calyx synapse. Neuron 23:399-409.

Steward O, Popovich PG, Dietrich WD, Kleitman N (2012) Replication and reproducibility in spinal cord injury research. Exp Neurol 233:597-605.

Ullstrom M, Parker D, Svensson E, Grillner S (1999) Neuropeptide-mediated facilitation and inhibition of sensory inputs and spinal cord reflexes in the lamprey. J Neurophysiol 81:1730-1740.

Wang C, Zucker RS (1998) Regulation of synaptic vesicle recycling by calcium and serotonin. Neuron 21:155-167.

Wood MR, Cohen MJ (1981) Synaptic regeneration and glial reactions in the transected spinal cord of the lamprey. J Neurocytol 10:57-79.

Zhang G, Jin L, Selzer ME (2011) Assembly properties of lamprey neuromodulin subunits and their expression after spinal cord transection. J Comp Neurol 519:3657-3671.

Zucker, RS, Regehr WG (2002) Short-term synaptic plasticity. Annu Rev Physiol 64:355-405.