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Running Title: Transplanting Mitochondrial in Spinal Cord Injury

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Abstract

Our previous studies reported that pharmacological maintenance of mitochondrial bioenergetics after experimental spinal cord injury (SCI) provided functional neuroprotection. Recent evidence indicates that endogenous mitochondrial transfer is neuroprotective as well, and so we extended these studies with a novel approach to transplant exogenous mitochondria into the injured rat spinal cord. Using a rat model of L1/L2 contusion SCI, we herein report that transplantation of exogenous mitochondria derived from either cell culture or syngeneic leg muscle maintain acute bioenergetics of the injured spinal cord in a concentration-dependent manner. Moreover, transplanting transgenically-labeled (tGFP) PC-12 derived mitochondria allowed for visualization of their incorporation in both a time-dependent and cell-specific manner at 24 hr, 48 hr and 7 days post injection. tGFP mitochondria colocalized with multiple resident cell types, though they were absent in neurons. Despite their contribution to the maintenance of normal bioenergetics, mitochondrial transplantation did not yield long-term functional neuroprotection as assessed by overall tissue sparing or recovery of motor and sensory functions. These experiments are the first to investigate mitochondrial transplantation as a therapeutic approach to treat spinal cord injury. Our initial bioenergetic results are encouraging, and though they did not translate with improved long-term outcome measures, caveats and technical hurdles are discussed that can be addressed in future studies to potentially increase long-term efficacy of transplantation strategies.

Keywords- BBB, Von Frey Hair, transgenic labeling, mitochondrial respiration
1. Introduction

Contusion spinal cord injury (SCI) can result in lifelong debilitation and has a profound impact on day to day activities of individuals living with SCI. In the event of an injury, the primary injury site comprises necrotic tissue that triggers secondary injury cascades, including glutamate excitotoxicity, decreased ATP production, increased lipid and protein oxidation, and resulting apoptosis and necrosis (see 1 and 2 for comprehensive review of pathophysiology following SCI). Mitochondrial dysfunction is intimately related to detrimental secondary pathophysiology resulting from SCI, and is pivotal to injury progression and disease in other models including traumatic brain injury (TBI), cardiac ischemia, stroke, Parkinson’s disease, and Alzheimer’s disease.3-13 As such, mitochondrial medicine is a developing field of therapeutics aimed at improving mitochondrial health and function for a variety of physiological disease and injury states. Using different methods to target mitochondria dysfunction, multiple groups have reported that this yields neuroprotection, tissue sparing, and functional recovery. For example, in models of nephrotoxicity, TBI, SCI, Parkinson’s Disease (PD), neurodegeneration, and oxidative stress, antioxidants decrease the ROS and RNS damage caused by dysfunctional mitochondria.14-24 Mild uncoupling of the electron transport system (ETS) increases ATP production following neurological insult in vitro, quinolinic acid neurological injury in vivo, and is therapeutic for the recovery from mitochondrial dysfunction after SCI.25-30 Mitochondrial substrate precursors such as acetyl L-carnitine (ALC) and N-acetylcysteine amide (NACA) improve mitochondrial function and behavioral recovery after TBI and SCI, as well as in PD and age-related cognitive function.24, 31-37 In summary, interventions that alleviate mitochondrial dysfunction exert a broad-spectrum therapeutic benefit across a wide range of disorders.

Mitochondrial transplantation is emerging as a potential therapeutic to maintain mitochondrial function after injury, consequently improving long-term functional outcome.38 The mechanisms by which transplanted mitochondria exert beneficial effects may include: increases in ATP production and calcium buffering capacity; decreases in overall ROS production; and generation of a new pool of mtDNA (for review see 39).

Though this therapy is relatively new, mitochondrial transplantation is effective in promoting recovery after ischemic injury to cardiac tissue.7, 40-42 More recently, astrocytes were found to inherently donate their mitochondria to nearby neurons after stroke injury.
in vivo, and astrocytes can be further induced to release mitochondria after injury, both of which can contribute to functional neuroprotection. Indeed, pharmacological blockade of astrocyte donation of mitochondria inhibited mitochondrial transfer and resulted in neuronal cell death, demonstrating the importance of mitochondrial transfer into damaged cells for cell survival. Furthermore, intravenous injections of isolated fluorescently-labeled mitochondria in a mouse model of PD resulted in a decrease in ROS along with increases in ATP production, Complex I activity and behavioral scores compared to vehicle-injected animals. In a rat model of PD, injections of Pep-1 conjugated isolated mitochondria into the medial forebrain bundle increased locomotor scores and dopaminergic neuronal sparing. Additional premise for the current studies is based on our recent report that exogenous transgenically-labeled mitochondria can be incorporated into cells both in vitro as well as in rat spinal cords at 24 and 48 hours post transplantation.

Herein, we investigated the therapeutic effects of mitochondrial transplantation immediately after SCI. We hypothesized that supplementation of the injured spinal cord with a pool of healthy, well-coupled mitochondria will provide a new source of increased energy production, decreased oxidation and dampened effects of glutamate excitotoxicity, resulting in maintenance of tissue bioenergetics. We further posited that exogenous mitochondria will incorporate into a variety of cell types and that the beneficial effects in the short-term would result in long-term benefits such as tissue sparing and improvements in hindlimb functional recovery.

2. Materials and Methods

2.1 Transgenic Labeling of PC-12 Cells

PC-12 cells were transgenically labeled with a mitochondrial-targeting tGFP, as previously described. Briefly, PC-12Adh cells were transfected with a plasmid vector containing tGFP and a mitochondrial targeting sequence for the cytochrome c oxidase complex subunit VIII. Following transfection, cells were kept under constant selection using media containing 3 µg/mL puromycin in complete media (F-12K Medium (ATCC cat # 30-2004 Manassas, VA) with 2.5% fetal bovine serum (Atlanta Biologicals # S1111OH, Atlanta, GA), 15% horse serum (Gibco # 26050-070), and 1.1% penicillin streptomycin (Corning # 30-002-CI, Tewksbury, MA).
2.2 Mitochondrial Isolation from Cell Culture

Purified mitochondria were isolated from transfected tGFP-PC-12 cells as previously described. Briefly, when cells were 95% confluent they were removed manually from culture plates using a cell scraper. Mitochondria were isolated using multiple centrifugation and resuspension steps to yield ficoll-purified mitochondria. Importantly, after ficoll purification a thin non-mitochondrial layer was gently aspirated off the final pellet, ensuring the sample was purified without contamination.

2.3 Mitochondrial Isolation from Soleus Muscle

Immediately following CO2 asphyxiation and decapitation, the soleus muscle was dissected from naïve Sprague-Dawley rats (n=10). The animal was placed on its stomach, with the hindlimb stretched backwards. 70% EtOH was sprayed on the hindlimb, and scissors were used to blunt dissect the skin away from the underlying calf muscle. The skin was then cut away from the muscle on the lateral sides, followed by cutting the Achilles’ tendon. The muscle was pulled up and away from the leg, exposing the underlying soleus muscle. The tendons were cut at both insertion sites, and the soleus muscle was removed and placed in 2mL of isolation buffer (215 mM mannitol, 75 mM sucrose, 0.1% BSA, 20 mM HEPES, and pH adjusted to 7.2 with KOH) containing 1 mM EGTA and 0.025% trypsin. The muscle was chopped into smaller pieces using scissors to increase surface area exposed to the isolation buffer and trypsin. The sample was then mechanically homogenized taking care not to introduce air bubbles. Protease inhibitor was added to halt trypsin activity, followed by centrifugation at 1500 rcf, 4°C, 5min. The supernatant was removed and centrifuged at 13,000 rcf for 10min, 4°C, after which the pellet was resuspended in isolation buffer and purified using ficoll gradient (7.5%/10%) centrifugation at 32,000 rpm for 30 min, 4°C. The pellet was removed and resuspended in 0.6 mL isolation buffer without EGTA, and centrifuged at 10,000 rcf for 10min, 4°C. The resulting pellet contain purified mitochondria that was resuspended in isolation buffer without EGTA for Pierce BCA (Thermo Fisher, cat #23225) protein assays and microinjection into spinal cords.

2.4 Spinal Cord Injury and Microinjection Procedures

Adult (12 weeks old, 225-250g) female naïve Sprague-Dawley rats (Harlan) underwent intraperitoneal injections of anesthesia (80mg/kg ketamine and 10mg/kg xylazine, Butler Animal Health Supply) followed by lacrilube ophthalmic ointment (Medi-Vet) applied to
both eyes before surgery to prevent drying. The surgical site along the dorsal spine was shaved with an electric hair trimmer and cleaned with both 70% ethanol and betadine solutions. Animals then received a T12 laminectomy to expose the L1/L2 spinal level.

Sham animals received a laminectomy only, whereas injured animals received a severe 250 kDyn contusion injury using a force controlled Infinite Horizon impactor computerized injury device (Precision Systems and Instrumentation, LLC, Lexington KY). Those animals that underwent spinal cord injury were split into three cohorts receiving either: 1) vehicle injection, 2) cell-derived tGFP mitochondria, or 3) leg muscle-derived mitochondria; uninjured animals did not receive injections (see Table 1). Injections were made within 30 min after injury into the mediolateral gray matter at 4 circumferential sites of the injured cord separated by 2mm in the rostral caudal direction (see Figure 3) using a glass micropipette needle (World Precision Instruments, Sarasota, FL cat no. 4878) pulled and beveled to a 20-30 µm inner diameter pore opening. Each injection consisted of 750 nL of either vehicle (isolation buffer with 5mM pyruvate, 2.5 mM malate and 10mM succinate) or mitochondria suspended in vehicle (for a total of 50, 100 or 150 µg); hence a total of 3 µL volume was injected per spinal cord. Muscle incisions were closed using silk sutures 3-0 Vicryl (Ethicon, Inc., Somerville, NJ), and skin incisions were closed using wound clips (Stoelting Co., Wood Dale, IL), followed by application of hydrogen peroxide and betadine. Immediately following surgery, animals were single housed in a clean cage on top of a towel on a 37°C heating pad, and monitored until recovery from anesthesia. Yohimbine (1–2 mg/kg s.c., Lloyd Laboratories) was used to counteract the effects of xylazine. Buprenorphine HCl (0.02–0.05 mg/kg s.c, Butler Animal Health Supply) was administered subcutaneously immediately after surgery to alleviate pain, and then every 8 hours for 3 days following operation. The antibiotic cefazolin (33.3 mg/kg, s.c Butler Animal Health Supply) was given twice daily for 5 days post-operatively to minimize incidence of infection. The animals were housed individually until wound clip removal, and cages were lined with absorbent pads without bedding for the duration of buprenorphine use, as pica is often observed in treated rats. After discontinuation of buprenorphine, pine bedding was used. Additional parenteral fluids (10 mL lactated Ringer’s solution, Baxter Healthcare Corporation, Deerfield, IL) were provided as needed for 24–72 h post-surgery to compensate for dehydration brought on by anesthesia. Bladder expression was performed
2 times per day until adequate spontaneous voiding returned to avoid urinary tract infections. Skin wound clips were removed 10–14 days post–injury under isoflurane anesthesia. Animals survived either 1) 24 hours for mitochondrial respiration analyses (n=58), 2) 24h, 48h, or 7 days for histological analyses (n=15), or 3) 6 weeks for long-term behavioral and histological analyses (n=36) (see Table 1).

2.5 Behavioral Analyses

The Basso, Beattie, Bresnahan (BBB) locomotor rating scale was used to analyze the hindlimb functional recovery of animals after contusion SCI. Assessments were performed according to published protocol. Animals underwent contusion injury, followed by BBB assessment at 2, 7, 14, 21, 28, 35, and 42 days post injury. Testing was performed at the same time in the same room on each testing day to reduce variation. The scores for the right and left leg were averaged for each time point. Mechanical tactile sensitivity was measured with von Frey filaments (Stoelting, Inc, Wood Dale, IL) as previously described and modified. The hindpaw was stimulated with an incremental series of eight monofilaments of increasing logarithmic stiffness. A modified up-down method was used to determine the 50% withdrawal threshold. An intermediate force von Frey filament (#4.31, 2.0g force) was first applied to the glabrous skin until the filament was slightly bent. In case of a positive response (withdraw of the paw within 2 seconds), the next filament of lesser force was applied. In the case of a negative response, the next filament greater force was applied. The thresholds for both hindpaws were averaged and reported.

2.6 Mitochondrial Isolation from Spinal Cords

For bioenergetics assays, animals were CO₂ asphyxiated and decapitated 24 hours after injury and injections. A 1cm segment of spinal cord surrounding the injury site and including all of the injection sites was quickly dissected and total mitochondria were isolated as described previously. Briefly, the spinal cord was placed in a homogenization tube with 2mL isolation buffer containing EGTA ((215 mM mannitol, 75 mM sucrose, 0.1% BSA, 20 mM HEPES, and pH adjusted to 7.2 with KOH) containing 1 mM EGTA), and physically homogenized with six quick rotations of the pestle, without introducing air bubbles into the solution. The sample was poured into a 2mL Eppendorf tube and centrifuged at 1400 rcf for 3 min, 4°C. The supernatants containing mitochondria were then poured into a separate Eppendorf tube, and this tube was filled up to 2mL with
isolation buffer containing EGTA. To increase yield, the pellet from the previous spin was resuspended in 500µL isolation buffer containing EGTA and again centrifuged at 1400 rcf for 3 min 4°C. The supernatants containing mitochondria from this spin was again poured into separate tubes and filled to 2mL with isolation buffer containing EGTA. The supernatant tubes were then centrifuged at 13000 rcf for 10 min at 4°C. The resulting supernatant was discarded, and 350 µL isolation buffer containing EGTA was used to resuspend each pellet containing mitochondria. The samples underwent nitrogen bombing at 1200 PSI for 10 min at 4°C before being placed on top of a ficoll gradient (7.5%/10%) and centrifuged at 32000 rpm for 30 min 4°C. The supernatant was removed, leaving a purified mitochondrial pellet. The pellet was resuspended in 450 µL isolation buffer without EGTA and centrifuged 10,000 rcf for 10 min 4°C to remove any remaining EGTA from the sample. The supernatant was removed, and the pellet resuspended in isolation buffer without EGTA to a concentration of ~15µg/µL. Pierce BCA protein assay was performed to determine protein content of the sample.

2.7 Assaying Respiration of Isolated Mitochondria

Mitochondria were assayed for oxygen consumption rate (OCR) immediately after isolation using the Seahorse Bioscience XF® Flux Analyzer at 37°C, as described previously.17, 52 Mitochondrial respiration was assessed using respiration buffer consisting of: 215 mM mannitol, 75 mM sucrose, 2 mM MgCl₂, 2.5 mM inorganic phosphates, 0.1% BSA, 20 mM HEPES, pH 7.2. After isolation, purified mitochondria were added to the respiration buffer in the cell plate, and the cell plate was centrifuged at 2000rpm for 4 min at 4°C to concentrate and adhere mitochondria to the bottom of each well. Then, 5mM pyruvate/2.5 mM malate/1mM ADP, 1µg/µL oligomycin, 3 µM FCCP, and 100nM rotenone/10mM succinate were sequentially added to each well. Oxygen levels were measured after each substrate/inhibitor addition to calculate each state of mitochondrial respiration according to published protocols.52 Briefly, state III respiration is a measure of oxidative phosphorylation that takes place in the presence of pyruvate, malate and ADP. State IV respiration occurs upon oligomycin addition to inhibit the ATP synthase complex, thus uncoupling the electron transport chain from the ATP synthase complex. State V.1 is mitochondrial respiration after addition of FCCP (protonophore) which collapses the inner membrane potential, and State V.2 respiration occurs after inhibition of complex 1 and
addition of succinate to provide electrons through complex 2 of the electron transport chain.

2.8 Spinal Cord Processing for Histological Analyses

Rats survived to their terminal time point after injection (24 hours, 48 hours, 7 days, or 6 weeks). They were overdosed with 0.2 mL Fatal-Plus containing sodium pentobarbital (Vortech Pharma Ltd., Dearborn, MI) followed by transcardial perfusion with 150 mL of 0.1M phosphate buffered saline (PBS), then with 250mL 4% paraformaldehyde (PFA) in PBS. A 3cm segment of spinal cord centered on the L1/L2 injury site was dissected, and post-fixed up to 4 hours with 4% PFA followed by washing overnight in 2M phosphate buffer. The cords were then cryoprotected in 20% sucrose/PBS at 4°C until the cords sank in the solution, then embedded in gum tragacanth (Sigma) blocks and serially cryosectioned coronally at 20 µm. For the 24 hour, 48 hour, and 7 day cohorts, every other section was mounted on a ten slide series so that adjacent spinal cord sections on any given slide were spaced equally 400 µM apart. For the 6 week studies, every 5th section was mounted onto a ten slide series with adjacent sections on a slide separated by 1000 µm.

2.9 Immunohistochemistry and Fluorescent Imaging

Slide-mounted spinal cord sections were washed 3 x 5 minutes in 0.1M PBS, followed by incubation in blocking buffer (0.1M PBS, 1% bovine serum albumin, 0.1% Triton -X 100, and 5% normal goat serum) for 1 hour. Sections were then incubated overnight at 4°C in primary antibody diluted to proper concentration in blocking buffer. The next day, after washing again, the sections were incubated with secondary antibodies overnight at 4°C before another wash followed by 3 hours incubation at room temperature in streptavidin-conjugated fluorochromes diluted to desired concentrations in 0.1M PBS. After final washes they were coverslipped with Vectashield (Vector Labs, cat #H-1000) containing 0.5 m Hoechst dye to label nuclei (Thermo Fisher, cat# 62249). Antibodies used for epifluorescent imaging and co-localization studies included rabbit anti-tGFP (0.13µg/mL, Evrogen # Ab513), mouse anti-RECA1 (5µg/mL, abcam # ab9774), mouse anti-GFAP (2µg/mL, abcam # ab10062), mouse anti-CC1 (0.5µg/mL, Cal Biochem #OP80), mouse anti-NeuN (5µg/mL, Millipore # MAB377), mouse anti-CD11b/c (OX42, 5µg/mL, abcam # ab78457), phosphor-PDGF Receptor β (5uL/mL Cell Signaling mAb #3166), goat
anti-rabbit 488 (4µg/mL, Invitrogen # A11008), goat anti-mouse Biotin (7.5µg/mL, Vector # BA-9200), and Streptavidin Texas red (3.3µg/mL, Vector # SA-5006). Z-stack images were obtained using Nikon Eclipse Ti Confocal microscope and processed with NIS Elements ND2 software (Nikon Instruments).

2.10 Quantification of tGFP Mitochondria Volume and Spread

A set of tissue section mounted slides for each time point was used to calculate the tGFP mitochondria volume and spread over time after injection. Anti-tGFP antibody was used to label injected mitochondria, as described above. Each spinal tissue section was then analyzed using the Nikon confocal microscope for the presence of distinct tGFP labeling. When a discreet tGFP injection bolus was present, Nikon NIS Elements ND2 software was used to measure the area of the bolus. Each of the four injection epicenters per spinal cord was identified by finding the largest area of discreet bolus. Every section, both rostral and caudal of the injection epicenter that contained a discreet tGFP bolus, was measured. After defining the boundaries of the injection bolus, the section thickness and periodicity of the sectioning paradigm were used to calculate the total injection bolus volume of all 4 injections in each cord. The rostral-caudal lengths throughout the spinal cords where punctate tGFP labeling was evident were also calculated, regardless of apparent discreet injection boluses.

2.11 Quantification of Brain Macrophage Response

For the 24 hour, 48 hour, and 7 day animal cohorts, macrophage densities at each injection epicenter was analyzed. Once the injection site was identified, based on tGFP and OX-42 immunolabeling, then a score of 0-3 was assigned for each injection epicenter where 0 indicates normal density in macrophages (as in naïve tissues), a score of 1 indicates activation (change in morphology to an activated state with processes withdrawn), a score of 2 indicates activated brain macrophages densities in the confines of the injection site, and a score of 3 indicates brain macrophage densities that expand beyond the original injection bolus boundaries.

2.12 Time course of Cell-type Co-localization

Cell-type co-localization of tGFP mitochondria was quantified for different cell types in the 24 and 48 hour and 7 day cohorts. Separate sets of slides were labeled for either microglia, endothelial cells, oligodendrocytes, astrocytes, pericytes or neurons, and
co-labeled with antibodies against tGFP. Eight sections total per spinal cord were assessed using the left side injections of each spinal cord including the injection epicenters, one section (400 µm) from the injection epicenter towards the injury, one section (400 µm) from the injection epicenter away from the injury site, and the first section away from the injection epicenter that no longer contained a visible bolus (see Figure 8A for counting schematic). The region of interest (ROI) counted for each section was the left dorsal quadrant of the spinal cord, drawing a vertical line from the central canal to the dorsal surface, and a horizontal line from the central canal to the lateral surface (see Figure 8B for depiction of ROI). Slides were visualized under an Olympus BX51 microscope at 100X magnification (10x eye piece) using a FITC/TRITC combination filter to simultaneously visualize the green and red fluorescent wavelengths. Co-localization was defined in two different ways. The first criterion of positive incorporation was when the tGFP labeling was evident within the confines of the labeled cell membrane, indicating the tGFP mitochondria were within the boundaries of the cell membrane. The second criterion of positive incorporation was when the tGFP labeling was present and overlapped with the cell specific marker within the cytoplasm and/or on cell surface. If either of these criteria were met, then this was considered a positive instance of co-localization. Sections were examined using the DAPI filter to ensure that the nucleus was within the focal plane.

Counts were performed within the ROI in a total of 8 spinal cord sections of each animal for 1) each time a specific cell type was co-localized with tGFP and 2) total cell counts for the specific cell type. Quantification of co-localization was reported as a percentage of tGFP-positive cells of a specific cell type divided by the total number of that specific cell type within the ROI.

2.13 Quantification of Tissue Sparing

Tissue sections were stained for eriochrome cyanine-cresyl violet (ECCV) to differentiate white matter and gray matter as previously described. Mounted sections were imaged using a Nikon microscope (Nikon Corporation, Tokyo, Japan), and Scion Imaging analysis software (Scion Corporation, Frederick, MD, USA) was used to delineate injured versus spared tissue in each section. ROIs were manually drawn around the entire cord, spared gray matter, and injured tissue which allowed extrapolation of the spared white matter area. Eleven sections equally spaced by 1000 µm (6 weeks cohort) or spaced
by 800 µm (24 and 48 hour cohorts) and centered on the injury epicenter were analyzed to calculate the volume of damaged versus spared tissues using the Cavalieri method.\textsuperscript{54, 55}

2.14 Stereology

Analysis was performed to determine the number of neurons in the medial portion of lamina VII and lamina X of the L1/L2 spinal cord in the 6 week injured cohorts. Stereological counts of neurons were performed as we have previously published.\textsuperscript{37} An Olympus BX51 microscope was used to image sections with the Bioquant image analysis program (Nova Prime, V6.70.10, Bioquant Image Analysis Corp., Nashville, TN). Briefly, a region of interest was drawn over lamina VII and X, containing putative central pattern generator (CPG) interneurons. A counting grid was placed over this region, with each counting frame measuring 125 µm X 125 µm. Disectors were set to 50 µm X 50 µm, and section thickness was set at 10µm, with a 2 µm guard zone on both the top and bottom of the tissue to keep from double-counting cells across sections. Bioquant software randomly placed disectors within the ROI, with an average of 13 disectors counted per section. NeuN positive cells were then counted under 40X magnification (10x eyepiece). The criteria for being counted required the neuron cell body to be contained within the disector and the nucleus in focus within the set 10µm counting thickness, as verified by Hoechst staining.\textsuperscript{56}

2.15 Statistical Analyses

Changes in each OCR state and respiratory control ratio (RCR) for acute bioenergetic studies, complex I functional assays, injected tGFP bolus volume and rostral-caudal length, cell-type incorporation, number of co-localization instances, and co-localization percentages within each time point were analyzed using a one-way ANOVA, with Tukey’s multiple comparisons post hoc test. Macrophage density analyses were performed using a Mann-Whitney U test for each time point analyzed. Rostral-caudal co-localization studies, von Frey test, BBB scoring and neuronal stereology counts were analyzed using a repeated measures two-way ANOVA with Tukey’s multiple comparisons when warranted. All analyses were carried out using Graphpad Prism 6 (Graphpad Software, Inc., La Jolla, CA). Significance was set to p < 0.05.
3. Results

3.1 Transplantation Maintains Acute Mitochondrial Bioenergetics

We transplanted mitochondria into the injured spinal cord in a dose-dependent manner. Vehicle or transgenically-labeled tGFP-mitochondria were microinjected into the injured spinal cord at 50, 100, or 150 µg, and 24 hours later the spinal cords were dissected and assessed for mitochondrial bioenergetics from spinal cord containing both endogenous mitochondria and exogenous transplanted mitochondria. The respiration measurements in the following figures represent combined assessments of both host mitochondria as well as exogenous transplanted mitochondria, the latter of which may be either extracellular or incorporated into cells. When assayed for mitochondrial bioenergetics in terms of OCR, there were significant differences in state III \( F(4, 27) = 6.269, P = 0.0011 \) and state V.1 \( F(4, 27) = 11.80, P < 0.0001 \) respiration rates among groups. Post hoc analyses revealed that, compared to significantly impaired rates in the vehicle injected group, the 100µg tGFP-injected group significantly maintained state III (89% of sham) and state V.1 (73% of sham) OCR (Figure 1). There were no significant differences in RCR among all injection groups or shams, indicating that mitochondria in each group were well-coupled.

The use of tGFP-labeled mitochondria allows for visualization and tracking of transplanted mitochondria in situ, which is important for determining cell-type incorporation and time lapse uptake of exogenous mitochondria. However, we have also shown that cultured cells can have mitochondria that are well coupled but respire at lower rates. Therefore, we also tested the injection of muscle-derived mitochondria, both as an alternative cell source with different respiratory capacity and potentially clinically applicable, as autologous transplant may be performed using mitochondria obtained from a muscle biopsy.

Muscle mitochondria were isolated from the soleus muscle of separate naïve rats and then similarly injected into the injured spinal cords. Following transplantation, there was a significant effect in state III respiration \( F(4, 21) = 8.714, P = 0.0003 \); post hoc analyses revealed significant increases with either 50 µg (92% of sham levels) or 100 µg (95% sham levels) mitochondria injection compared to vehicle injected groups (Figure 2). There were also significant effects in state V.1 respiration \( F(4, 21) = 10.73, P < 0.0001 \)
and state V.2 respiration \( [F(4, 21) = 2.977, P = 0.0429] \). Post hoc analyses showed significant decreases in state V.1 respiration after injury, which was maintained near sham levels with 100ug mitochondrial transplantation. All injection groups or shams were equally well-coupled as shown in the RCR values.

The state III OCR results indicate that 100 µg mitochondria transplantation, using either source of mitochondria, maintained oxidative phosphorylation function in the injured spinal cord to ~90% of sham levels. We previously reported that rescue of state III respiration to 67% of sham levels corresponds with long-term tissue sparing and behavioral recovery.\(^\text{37}\) Thus, the 100 µg dosage was used to investigate the effects of mitochondrial transplantation on both cell-type incorporation and long-term recovery.

### 3.2 Characterization of Transplanted tGFP Mitochondria in situ

We characterized the uptake of exogenous tGFP mitochondria into resident cells of the injured spinal cord. Analysis of multiple time points after injury gave a temporal depiction of exogenous mitochondrial incorporation within tissues after 24 hours, 48 hours, and 7 days post injection. Injection sites presented as elliptical boluses at the injection epicenter, with scattered punctate labeling further from the injection site (Figure 3A). Earlier time points showed very discreet injection boluses (Figure 3B, C), in addition to scattered punctate tGFP mitochondria within the cord (Figure 3D). By the 7 day time point, these discreet injection boluses were difficult to distinguish, save for a few instances, and only dispersed punctate tGFP mitochondria were present.

We analyzed the spread of the injected tGFP mitochondria in two ways. The first was a measure of the rostral-caudal distribution of all 4 injection boluses. There were significant differences in bolus length among time points \( [F(2, 9) = 22.46, P= 0.0003, \text{Figure 3E}] \); post hoc analysis revealed a significant decrease in bolus length of the 7 day group compared to 24 and 48 hour time points. The length of the bolus appeared to increase, though not significantly, in the 48 hour group compared to 24 hour group. We further measured the rostral-caudal length of punctate mitochondria, and found that there were no significant differences among transplantation groups \( [F(2, 9) = 2.996, P=0.1006] \) (Figure 3F). The second analysis was a measure of total volume of injectate throughout the spinal cord (Figure 3G). The tGFP mitochondria injection bolus volume was significantly different among time points \( [F(2, 9) = 15.61, P= 0.0012] \); post hoc analysis
indicated that the injection did not change volume at either the 24 hour or 48 hour time points, but decreased volume at the 7 day time point.

All of the tGFP injection sites (4/animal) for the 24 and 48 hour groups were visible using fluorescent microscopy. By 7 days post injection, the injection sites had almost disappeared so that only 3 total injections across all four animals in the group could be visually verified. Some instances of a remaining injection bolus were still evident, and the rostral-caudal spread of punctate tGFP stippling was similar to 24 and 48 hour time points.

We set out to characterize the cell-type specific co-localization with tGFP mitochondria. We looked at multiple different resident spinal cord cell types when performing our analyses including neurons, brain macrophages, endothelial cells, astrocytes, pericytes, and oligodendrocytes (Figure 4).

When examining the different cell types within the injection epicenter bolus, we noted an almost complete absence of neurons while there was a high density of brain macrophages. The concentration of mitochondria or volume of the injections could be damaging the host tissues. When comparing vehicle-injected and tGFP mitochondria-injected spinal cords, there was typically increased brain macrophage activation at the mitochondria injection sites compared to vehicle (Figure 5A). This indicates that the tGFP mitochondria boluses could be causing inflammatory responses, and semi-quantitative analyses of brain macrophage densities within the injection epicenters at different time points (Figure 5B) showed a significant difference between vehicle- and tGFP mitochondria-injected groups at 24 hours [p = 0.0012], 48 hours [p = 0.0002], and 7 days post injection [p = 0.0196].

To discern which cell types incorporated exogenous mitochondria, we quantified incidences of co-localization. Representative images show positive co-localization of various cell types both away from and within the boluses of tGFP injections (Figure 6), defined in two different ways. The first criterion of positive incorporation was tGFP labeling present within the cytoplasm of the cell, as evident by green tGFP labeling within the confines of the red cell membrane marker juxtaposition to the nucleus (DAPI). The second criterion of positive incorporation was when the tGFP labeling overlapped with the cell specific marker, resulting in a yellow signal. If either of these definitions was met, this was counted as a positive instance of co-localization.
We found instances of cell-type specific co-localization with tGFP mitochondria for various cell types including brain macrophages, endothelial cells, pericytes, astrocytes, and oligodendrocytes at 24 hours post transplantation (Figure 7), with most conspicuous labeling in endothelial cells and pericytes. Co-localization was evident for all cell types both within the injection boluses as well as more distally, where sparse punctate tGFP was present. However, no evidence of co-localization was found within neurons either within the bolus or distally.

We quantified the propensity of tGFP mitochondria to be incorporated into different cell types spatially with respect to injection sites and across time. Tissue sections were analyzed per hemi cord as depicted in the schematic (Figure 8A), including the two injection epicenters, rostral and caudal to those, and the first section distally that only had punctate tGFP present. To compare propensities for exogenous mitochondrial uptake, from each section we quantified the percentage of a specific labeled cell type within the dorsolateral cross section ROI (Figure 8B) that were co-localized with tGFP mitochondria.

Comparisons were made to determine which cell type has a higher propensity to incorporate mitochondria. The cell types counted for each of the sections quantified, as described in Figure 8, were summed to give a total number per spinal cord (Figure 9A, white), and instances of co-localization from each of the 8 ROI analyzed were totaled (Figure 9A, black). A one-way ANOVA across the total number of cells counted among each specific type showed significant differences at 24 hours \( F(4, 15) = 72.59, P<0.0001 \), 48 hours \( F(4, 15) = 87.46, P<0.0001 \), and 7 days \( F(4, 15) = 37.32, P<0.0001 \). Post hoc analyses revealed that there were significant differences in the total cell counts among cell types, interestingly the total brain macrophages counted were significantly higher than all other cell types at each time point. Alternatively, a one-way ANOVA showed significant differences among the number of co-localized cells across cell types at 24 hours \( F(4, 15) = 10.62, P=0.0003 \), 48 hours \( F(4, 15) = 45.55, P<0.0001 \), and 7 days \( F(4, 15) = 5.159, P=0.0081 \). Post hoc analyses showed that co-localization numbers were significantly higher in brain macrophages than all other cell types (Figure 9A), further depicted in the percentage of cells that co-localized with tGFP (Figure 9B). A one-way ANOVA revealed significant differences between cell type co-localization percentages at 24 hours \( F(4, 15) = 6.384, P=0.0033 \) and 48 hours \( F(4, 15) = 5.854, P=0.0048 \). Tukey’s multiple
comparisons showed that both brain macrophages and pericytes co-localized with tGFP at significantly higher rates than oligodendrocytes at 24 and 48 hours post injection, which had the lowest co-localization percentages at any time point.

The injection epicenters contained dense boluses of tGFP mitochondria, and based on the high concentrations in this area it may be difficult for mitochondria to have access to various host cells for incorporation. Notably, sections that were further from the injection epicenters showed highly dispersed tGFP mitochondria that may have differential uptake since higher concentrations of mitochondria have been shown to aggregate with less extensive distribution into nearby brain tissues in a rat model of Parkinson’s disease.\textsuperscript{57}

We therefore mapped the propensity of co-localization in relation to the injection epicenters, as described above, and analyzed this across the rostral-caudal distribution of all the sections analyzed for each spinal cord (Figure 10).

Two-way ANOVAs were performed within time points for each cell type and showed differences in cell-type co-localization for macrophages \([F (2, 9) = 8.123, P = 0.0096]\), endothelial cells \([F (2, 9) = 10.69, P = 0.0042]\), oligodendrocytes \([F (2, 9) = 9.575, P = 0.0059]\), astrocytes \([F (2, 9) = 22.54, P = 0.0003]\), and pericytes \([F (2, 9) = 928.3, P < 0.0001]\). We found that co-localization percentages depended upon their location in relation to the injection epicenters, especially at the 24 hour time point when macrophages had higher incorporation percentages one section rostral of each injection site (400 \(\mu\)m) compared to the most caudal section analyzed. Endothelial cells and astrocytes had higher co-localization percentages closer to the injection epicenters compared to the distal sections, while pericytes tended to have more stable co-localization across sections. Oligodendrocytes had very low incorporation percentages, with the least incorporation distal of the injection epicenters. This indicates spatial-dependent incorporation in addition to cell-type dependent incorporation.

We also performed morphometric analyses to examine possible changes in lesion volume or tissue sparing at acute time points after injury (Figure 11). Overall, there were no apparent changes in lesion volume, gray matter sparing, or white matter sparing.

### 3.3 Behavioral Recovery and Tissue Sparing after Mitochondrial Transplantation

We tested the effects of mitochondrial transplantation after SCI on hindlimb functional recovery and tissue sparing. Anesthetized animals underwent a severe
contusion injury at the L1/L2 spinal level, followed immediately by 4 injections for a total of 100 µg tGFP-labeled mitochondria into the injury penumbra (see Figure 3A for injection schematic).

The starting numbers per experimental group were n=12. Through the course of the 6-week long study, we observed an attrition of animals typical of SCI studies. Three animals died from anesthesia during surgical procedures, three animals were removed from the study and euthanized due to extensive autophagia (self-cannibalism or chewing and biting of tissue) of the hindlimbs, and one animal was removed from the study due to its high 2-day BBB score (more than 2 SD from the mean, indicating a bad contusion injury). This resulted in final groups of Vehicle injection n= 10, tGFP mitochondria injection n=11, muscle mitochondria injection n=8.

Animals were tested for mechanical sensitivity of hindpaws with the von Frey hair test and for hindlimb motor recovery with the overground locomotor BBB scale. Within 3 weeks of injury, we observed the development of mechanical hypersensitivity in all groups (Figure 12A). Mitochondria transplant produced an extended duration of hypersensitivity to at least 6 weeks post injury, as compared to all other groups. A two-way ANOVA revealed a main effect of time \([F (2, 52) = 17.64, P<0.0001]\) but not treatment \([F (2, 26) = 1.370, P = 0.2718]\). Post hoc analyses showed each treatment group had significantly decreased scores at 3 weeks compared to their within group baseline levels, indicating potentially increased hypersensitivity. As indicated earlier, three animals were removed for the study due to extensive autophagia- after experiments were completed and data analyzed, the study was unblinded and revealed that these animals had all been in the muscle mitochondria transplanted group.

A two-way repeated measures ANOVA revealed no effect of treatment on BBB scores \([F (2, 26) = 0.01894, P = 0.9812]\) (Figure 12B). All animals showed some functional recovery consistent with our previous studies. \(^{37}\) The injured rats in the current studies plateaued with an average BBB score of ~8, which is indicative of hindlimb sweeping and plantar placement without weight bearing. \(^{48}\) This level of recovery correlates to historical data using this injury severity and spinal level, indicating there was no overt damage/effect of injections volumes. It is important to note that while we did not see improvements in functional recovery with mitochondrial transplantation, there was no further impairment
of function, which had been a concern, based on possible increased inflammatory responses.

Following terminal behavioral analyses, animals were euthanized and spinal cords were processed for tissue sparing analyses that showed similar histopathology among treatment groups, supporting behavioral findings (Figure 13). There were no significant differences among groups for lesion volume (Figure 13A). Mitochondria injections were targeted to the medial lateral gray matter, allowing for possible mitochondrial integration into both gray matter and white matter, but there were no differences among treatment groups for either gray matter or white matter tissue sparing (Figure 13B and C, respectively).

Our previous documentation of improved functional recovery after treatment with an alternative mitochondria biofuel (ALC) was highly correlated with significant sparing of putative CPG interneurons rostral to the SCI (Patel et al., 2012). Despite no behavioral differences, we investigated similarly whether mitochondrial transplantation spared interneurons by performing stereology after injury and transplantation (Figure 14). We defined the ROI for neuronal counts to include lamina X and the central portion of lamina VII (Figure 14A).

The number of neurons estimated in each section was extrapolated using a previously published equation. The coefficient of error (CE: vehicle - 0.116, tGFP mitochondria - 0.1126, muscle mitochondria - 0.1456) and coefficient of variance (CV: vehicle - 0.7007, tGFP mitochondria - 0.6506, muscle mitochondria - 0.5145) were calculated for each group to establish intra- and inter-animal variation, respectively. There were significant differences in neuronal sparing among groups [F (2, 286) = 3.658, P=0.0270], and post hoc analyses showed significantly more neurons at 1 mm caudal of the injury epicenter in the tGFP mitochondria-injected groups compared to both vehicle- and muscle mitochondria-injected groups.

4. Discussion

The current study is the first to assess mitochondrial transplantation into the injured rat spinal cord and its effects on bioenergetics at 24 hours post injury/transplantation, as well as long-term behavior and histology. We found that injections of both tGFP labeled culture-derived and muscle-derived mitochondria
significantly maintained state III respiration at the 100 µg dosage compared to vehicle injections. We have previously shown that pharmacologically maintaining state III respiration near sham levels after L1/L2 SCI results in functional neuroprotection 37, accordingly both sources of isolated mitochondria were injected at 100 µg to evaluate the effects of mitochondrial transplantation on host cellular uptake acutely, as well as long-term histological and behavioral outcome measures.

4.1 Mitochondrial Incorporation into Host Cells

We first assessed the time course of exogenous tGFP mitochondrial incorporation into host cells. Discreet tGFP mitochondria boluses were evident at 24 and 48 hours, though they decreased significantly in volume and rostral-caudal spread by 7 days post injection. There are likely various reasons for the scarcity of tGFP label at 7 days (and 6 weeks) after injection, notably phagocytosis especially at the injection site bolus where the mitochondria density may be too high to be incorporated into the surrounding host cells. 57 The tGFP mitochondria within the extracellular space likely diffuse easily through tissues away from the injection site over time, especially in the white matter where there is greater longitudinal diffusion. Moreover, when tGFP mitochondria successfully incorporate into cells, they may fuse and divide with host cell mitochondria leading to a wider dispersal of tGFP fluorescent within cells. Conversely, others have reported evidence of fluorescently-labeled mitochondria at chronic time points. Using direct mitochondrial transplantation and an antibody against human mitochondria, the presence of exogenous human mitochondria in a porcine model of cardiac ischemia was shown 4 weeks after injection. 59 This method of labeling, however, was not dependent on the retention of a fluorescent tag that was present at the time of injection, ie the tGFP tag, and instead used antibodies to label a different species-derived mitochondria in situ. Alternatively, GFP-labeled mitochondria conjugated with the peptide carrier protein, Pep-1, and directly injected into rat brains in a Parkinson’s disease model were observed in the medial forebrain bundle 12 weeks after injection. 57

4.2 Differences in Cell-type Incorporation

We also evaluated the propensity of tGFP mitochondria incorporation into various cell types and revealed that brain macrophages and pericytes had the highest propensity to co-localize with exogenous mitochondria. We cannot discern, however, whether the
tGFP mitochondria were functionally incorporated within macrophages or if they had been degraded and phagocytosed. Alternatively, pericytes are found within the vascular basement membrane and are closely associated with endothelial cells with many similarities to macrophages, including mesodermal origin, possessing of lysosomes, similar cell surface markers, and prostaglandin secretion (for review on CNS pericytes see 60–62).

Further, these cells have contractile properties and can regulate capillary diameter, making them influential in vascular perfusion of the spinal cord, especially after injury.63 These traits may explain their high propensity to co-localize with exogenous mitochondria, though as with the macrophage co-localization we cannot definitively say at this time if the mitochondria were phagocytosed or incorporated functionally into the cells.

Oligodendrocytes were least co-localized with tGFP mitochondria, though this could possibly be due to the location of the injection sites more targeted to the gray matter. While tGFP mitochondria co-localized with variety of cell types, it is important to note that they were conspicuously absent in neurons. In fact, there was conspicuous neuronal loss in the dorsal gray matter where injection boluses were located (Figure 4) which may be due to multiple factors. High concentrations of mitochondria can result in aggregation of the mitochondria, further making it difficult to incorporate into cells. These aggregations could physically injure neuronal cells as well as cause a heightened inflammatory response.

This, in addition to the volume of fluid injected could disrupt the tissue. We have noted that injection volume of vehicle alone activated brain macrophages. The white matter of the spinal cord is more amenable to fluid diffusion along axons- in gray matter where cell bodies are more compactly arranged, injected fluids could cause more physical stress even with the slow injection rates. Therefore, a balance must be attained that allows for low enough concentration of mitochondria that there is minimal aggregation, but also low total volume injected to avoid tissue damage. Additionally, a delivery method in which the mitochondria are somehow encapsulated may be a defining feature necessary for incorporation into neurons, as was used in previous studies utilizing mitochondria transplant in stroke43. It is important to note, however, that if other cell types in the injury penumbra can be spared following mitochondrial transplantation, then the inhospitable environment may be more favorable for neuroprotection. For instance, salvaging
pericytes and endothelial cells could increase perfusion of the spinal cord, thus being overall protective indirectly of neuronal protection.

### 4.3 Long-Term Outcome Measures

Despite significant effects on injured tissue bioenergetics acutely, we found that the same transplantation paradigm used in long-term studies did not result in reciprocal restoration of hindlimb locomotion or sensory tactile thresholds, nor tissue sparing. As expected, contusion SCI significantly decreased BBB locomotor scores for each group, with characteristic spontaneous functional recovery that plateaued five weeks post-injury consistent with severe injury at the L1/L2 spinal level. Transplantation of mitochondria from either source did not increase or hamper this recovery. An important consideration is the origin of the donor mitochondria. The current experiments utilized mitochondria isolated from cultured PC-12 cells or soleus muscle tissue. Mitochondrial respiration and function can vary dependent upon the source. Here, we used a mitochondrial origin similar to the recipient species (rat), and we compared two different tissue source types. Future studies and characterizations comparing donor tissues with different mitochondrial respiratory activities may establish sources that are more beneficial in our trauma/transplantation paradigm. Determining the compatibility of injected mitochondria with host cell mitochondrial networks may reveal that autologous transplantation has different effects than syngeneic or xenogeneic sources.

Contusion SCI also decreased mechanical sensitivity thresholds in each group, with some recovery towards baseline. However, mitochondrial transplantation increased, rather than decreased, the duration of this hypersensitivity, indicating possibly enhanced central neuropathic pain. Morphometric analyses in relation to the injury epicenter were consistent with the BBB outcome measures, showing no significant group differences in the rostral-caudal extent of lesion volume or spared gray and white matter volumes. Interestingly, however, cell counts showed significantly higher numbers of interneurons in the dorsal gray commissure 6 weeks after tGFP mitochondria injections compared to vehicle or muscle-derived mitochondria, though this was only evident 1 - 2mm caudal to the injury epicenter. Nevertheless, while these differences were significant, it did not result in changes in locomotor recovery. This is in diametric contrast to our documentation that sparing of dorsal gray commissure interneurons rostral to the injury epicenter
following treatment with the alternative mitochondrial biofuel, ALC, correlated with significant functional improvement in hindlimb function and tissue sparing, in the same injury model.37, 65

Our behavioral (BBB) findings are in agreement with Magnuson et al. (2005) showing that kainic acid depletion of hindlimb CPG neurons with caudal gray matter sparing after contusion SCI does not result in improved BBB scores. This indicates that critical interneurons comprising the CPG located at the L1/L2 spinal level in rats were not spared with mitochondrial treatment, notably those rostral to the injury epicenter that appear critical for functional improvements. These neurons mediate rhythmic output to motor neurons of the lumbar enlargement, so had we found sparing of interneurons cells rostral to the injury then we may have seen BBB improvements, as in previous reports.37, 65

4.4 Addressing Potential Inflammatory Responses

Our findings show the feasibility of mitochondrial transplantation into injured spinal cord tissue and ability to maintain acute bioenergetics. There are multiple caveats to consider as to why this did not correspond to improved long-term functional outcome measures. We noted that the influx of brain macrophages into the injection sites appeared enhanced in the mitochondrial transplantation group, although this did not alter overall tissue sparing at all acute and long-term time points. The mitochondria we assessed included both endogenous host mitochondria as well as injected mitochondria, so those from accumulating brain macrophages were included in, and may have contributed to results of our bioenergetics assays. The volume of vehicle or tGFP mitochondria injections likely disturb the surrounding spinal cord tissue causing nominal injury and subsequent influx of macrophages. Further, mitochondria are a source of damage associated molecular patters (DAMPS) 66 that can elicit stronger immune responses compared to vehicle alone. Thus, the consequent accumulation of brain macrophages after mitochondrial injections may have increased the overall number of respiration competent cells in injury sites evaluated. While the noted focal cellular immune responses did not alter acute and long-term histopathology or long-term over ground locomotion, mitochondrial transplantation prolonged the duration of hindlimb hypersensitivity after SCI. Hypersensitivity in the muscle-derived mitochondria transplant group may have contributed to noted attrition
due to autophagia, but whether enhanced local inflammatory responses contributed is uncertain.

On the contrary, previously published studies investigating mitochondrial transplantation that tested for inflammatory markers did not find increases. Masuzawa et al (2013) found that after direct autologous mitochondria injection into cardiac tissue there was significantly decreased expression of tumor necrosis factor alpha, interleukin 6, and high-sensitivity C-reactive protein.40 Further, they did not find anti-mitochondrial antibodies present in blood serum samples 28 days after injections. However, there could be differences when allogeneic mitochondria are transplanted, as in our study, compared to a syngeneic or autologous mitochondrial transplant; and into the injured spinal cord.

4.5 Possible Caveats and Future Directions

While there was no apparent integration of transplanted mitochondria into resident neurons, their incorporation into other cell types should not be discounted. The spinal cord is a heterozygous mixture of different cell types, each with important characteristics for the overall health and function of the spinal cord. Macrophages can be either pro-inflammatory or anti-inflammatory, and are responsible for phagocytosing debris and foreign substances as well as having a part in mediating immune response.67 Astrocytes are important support cells for neurons, help maintain the blood-brain barrier, and carry out functions in ion homeostasis of the extracellular fluids.68 Oligodendrocytes play an imperative role in insulating axons to increase signal transmission efficiency.69 Endothelial cells supply blood and oxygen to tissues, and pericytes wrap around endothelial cells for contraction of vessels and regulation of blood flow.70 Each of these cell types have important roles in maintaining homeostasis within spinal cord tissues, and salvaging any of these cell types with healthy mitochondrial delivery could be beneficial for functional recovery and neuronal cell survival after SCI.

There is the unconventional possibility that transplanted mitochondria exert their effects extracellularly. The McCully lab noted functional benefits of cardiac tissue mechanics after ischemia/reperfusion injury was followed by direct mitochondrial injections.7,40 Similar to our studies, they noted a low incorporation rate into host cells and theorized that their results were attributed to extracellular functions of the transplanted mitochondria. If mitochondria can function extracellularly then it can be reasoned that this
would be short-lived before immune responses and debris-clearing mechanisms are enacted to remove them, and may thus explain our short-term benefits without corresponding long-term correlates.

Our recent studies in vitro show that co-incubation alone was sufficient for tGFP mitochondria uptake into cultured cells. Similarly, we show here that direct injection of isolated mitochondria into spinal cord tissue in vivo results in cellular uptake, though there is a relatively low incorporation rate (around 6% or lower for any specific cell type analyzed). Increasing incorporation efficiency after SCI may be a critical step to making such transplantation approaches successful in achieving long-term recovery goals. Multiple mechanisms of exogenous mitochondrial incorporation have been proposed, although there is currently no consensus whether it is endocytosis-mediated or much more complicated.

Mitochondrial uptake is reported following co-incubation with cells in vitro or into host cells after direct injection in vivo. Conjugation of mitochondria with Pep-1, a cell penetrating peptide that increases incorporation, further promotes mitochondrial uptake, and connexins or gap junction-mediated communication appears necessary for cells to take in extracellular mitochondria, perhaps via actin-dependent processes. Since mitochondria are theorized to be bacterial in origin, and various bacteria up to 5 µm in size can enter cells and replicate, we posit here that similar mechanisms may be used to incorporate isolated mitochondria 500 nm in size. Bacteria can have surface proteins that bind to receptors located on host cells, subsequently initiating zipper-like actin-mediated phagocytosis. As mitochondria are bacterial in origin, and have retained their own set of mitochondrial DNA that encodes for proteins, it is possible then that mitochondria may have the same or similar surface proteins that can bind to cellular receptors. If mitochondria do indeed utilize the zipper-like actin-mediated phagocytosis method of cellular entry, this theory would support others’ findings that mitochondrial incorporation is actin-dependent. The zipper-like mechanism of bacterial incorporation can utilize different receptors such as integrins for binding and signaling (for review of bacterial invasion methods see). Internalization of exogenous mitochondria-containing particles in vivo is reported to be dependent upon integrin-mediated Src/Syk signaling, further implicating the zipper-like method of internalization as the means for
exogenous mitochondrial incorporation *in vivo*. Further in depth comparisons are needed to determine if mitochondria contain surface proteins that can be responsible for binding and infiltration of host cells.

While our findings of acute maintenance of cellular respiration is promising, perhaps assessing acute bioenergetics is not the optimal criteria for predicting long-term functional recovery in mitochondrial transplant paradigms after SCI. Accordingly, further investigations of other biomarkers and/or outcome measures of recovery at the more acute time points within the first week after injury appear necessary. Herein we show the feasibility of mitochondrial transplantation after SCI, though important caveats such as incorporation mechanism and mitochondrial viability and functional integration remain unanswered. Consequently, this seminal study advances the burgeoning field of mitochondrial transplantation and lays the foundation to increase incorporation efficiency and cell-type targeting that may be key in improving long-term functional neuroprotection in models of SCI.

**Acknowledgements**

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**Conflict of Interest**

The authors do not have any conflicts of interest to disclose.
References-


Table 1. Separation of animal cohorts by outcome measures at different time points. It should be noted that in addition to these rats, 10 naïve rats were used as donors for leg muscle-derived mitochondria, and 7 rats from long-term studies were excluded for various reasons (see Section 3.3).

<table>
<thead>
<tr>
<th>Outcome Measure</th>
<th>Group</th>
<th>Dosage of Mitochondria (ug)</th>
<th>Terminal Time Point</th>
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<tr>
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<td>N/A</td>
<td>24 hr</td>
<td>11</td>
</tr>
<tr>
<td></td>
<td>Vehicle</td>
<td>N/A</td>
<td>24 hr</td>
<td>11</td>
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<tr>
<td></td>
<td>tGFP Mito</td>
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<td>24 hr</td>
<td>6</td>
</tr>
<tr>
<td></td>
<td></td>
<td>100</td>
<td>24 hr</td>
<td>7</td>
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<tr>
<td></td>
<td></td>
<td>150</td>
<td>24 hr</td>
<td>6</td>
</tr>
<tr>
<td></td>
<td>Muscle Mito</td>
<td>50</td>
<td>24 hr</td>
<td>5</td>
</tr>
<tr>
<td></td>
<td></td>
<td>100</td>
<td>24 hr</td>
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<td>6</td>
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<tr>
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<td>1</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>48 hr</td>
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<td></td>
<td></td>
<td>7 days</td>
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<tr>
<td></td>
<td>tGFP Mito</td>
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<td>24 hr</td>
<td>4</td>
</tr>
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<td></td>
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<td>48 hr</td>
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<td></td>
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<td>7 days</td>
<td>4</td>
</tr>
<tr>
<td>Long Term Studies- Behavior, Tissue Sparing, Neuronal Cell Counts</td>
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<td>tGFP Mito</td>
<td>100</td>
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<td>11</td>
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<tr>
<td></td>
<td>Muscle Mito</td>
<td>100</td>
<td>6 weeks</td>
<td>8</td>
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Figure 1. tGFP mitochondrial transplantation maintains respiration rates. SCI caused significant decreases in mitochondrial respiration at state III, IV, and V.1, which were spared partially following mitochondrial transplantation. State III and V.1 OCR were maintained near sham levels after 100μg tGFP mitochondrial transplantation. Bars are means ± SEM. *p<0.05 vs Sham; #p<0.05 vs Vehicle. (One-way ANOVA for each state, Tukey’s multiple comparison) n=6-7/group in triplicate.
Figure 2. Muscle-derived mitochondrial transplantation maintains respiration rates.

State III, V.1 and V.2 OCR were maintained near sham levels after 100 µg muscle-derived mitochondrial transplantation. Bars are means ± SEM. *p<0.05 vs Sham; #p<0.05 vs Vehicle. (One-way ANOVA for each state, Tukey’s multiple comparison). n=4-6/group in triplicate.
Figure 3. Quantification of transplanted tGFP mitochondria distribution.

A. Schematics depict location of mitochondrial injections into the spinal cord. Cross section shows a portion of the injection bolus is boxed in red, and representation of the punctate area analyzed is boxed in yellow. Longitudinal horizontal section shows four peri-contusion injection sites (black stars) with injection boluses as solid green ellipsoids. Punctate tGFP labeling is represented by the more shaded green halos. B. Representative image of bilateral tGFP injections. For volumetric measurements, the discreet bolus was circled (yellow) to give an area for each section which was used to calculate the subvolume of each section measured. C. High magnification image of tGFP located within the bolus of (B) as delineated in red box in (A). D. Representative high magnification image of punctate tGFP-labeled mitochondria in (B) that were not part of the discreet bolus, as delineated in yellow box in (A). E. Rostral-caudal length of tGFP bolus was significantly decreased at 7 days compared to 24 hour and 48 hour groups. F. Length in the rostral-caudal direction that any visible evidence of punctate mitochondria was apparent. G. tGFP injection bolus volumes significantly decreased at 7 days post injection. Scale Bars- B= 200µm, C and D =10µm. Bars are means ± SEM. *p<0.05 vs 24 and 48 hrs. (One-way ANOVA, Tukey’s multiple comparisons) n=4/group.
Figure 4. Resident cell types of the spinal cord proximal to tGFP mitochondrial injection bolus. The top panel shows an injection bolus into the dorsal horn of the spinal cord after 24 hours. Adjacent slide series (separated by 40µM) were stained with specific cell type markers, and the white dotted lines outline the dorsal horn gray matter while green lines delineate margins of the injection bolus. Within the injection bolus, there appeared to be a loss of neurons and oligodendrocytes. While there were few astrocytes within the bolus, they were prevalent in the area surrounding the injection. There was a noticeably increased density of brain macrophages in the area of the injection bolus. Additionally, there were endothelial cells and pericytes seen throughout the injection bolus. tGFP = exogenous tGFP mitochondria, NeuN = neurons, OX42 = brain macrophages, RECA = endothelial cells, GFAP = astrocytes, PDGFR-B = pericytes, CC1 = oligodendrocytes, DAPI = cell nucleus. Scale bars = 100 µm.
Figure 5. Brain macrophage activation at the mitochondrial injection bolus.

A. Microgliosis was compared among naïve, injured/vehicle injected, and injured/tGFP mitochondria injected spinal cords 24 hours after injury. There is a conspicuous high density of brain macrophages within the tGFP injection site compared to vehicle. B. At the injection epicenters, brain macrophage density was given a score of 0-3, where 0 reflects normal quiescent morphologies (A naïve) and 3 reflects overt microglial activation and accumulation of brain macrophages. There was a significant increase in the macrophage response when tGFP mitochondria were injected compared to vehicle injections at all time points. Error bars depict median ± SEM. *p<0.05 vs vehicle; (Mann-Whitney U test for each time point) n=4 vehicle injections per time point, n=16 tGFP injections per time point.

OX42 = brain macrophages, DAPI = cell nucleus. Scale bars= 200 µm.
Figure 6. Representative images of cell type-specific tGFP co-localization.

Co-localization with different cell types at 24 hours post injury in captured Z-stack images, with the bottom panel of each image representing the x plane and the right panel representing the y plane. ‘Within Bolus’ (left column) indicates images that were taken in the area of dense tGFP mitochondria, while ‘Away from Bolus’ (right column) indicates images that were taken in areas with more sparse, punctate tGFP mitochondria. Co-localization was evident within macrophages (A, B), endothelial cells (C, D), pericytes (E, F), astrocytes (G, H), and oligodendrocytes (I, J). OX42 = brain macrophages, RECA = endothelial cells, PDGFR-B= pericytes, GFAP = astrocytes, CC1 = oligodendrocytes, tGFP = exogenous tGFP mitochondria, DAPI = cell nucleus. Scale bars= 10um.
Figure 7. High magnification images showing positive co-localization. Images show instances of cell-type specific co-localization of tGFP mitochondria with either macrophages, endothelial cells, or pericytes. White arrows point to instances of tGFP mitochondria co-localization within the cells. Tissues were processed 24 hours post tGFP mitochondria transplantation. OX42 = brain macrophages, RECA = endothelial cells, PDGFR-B = pericytes, tGFP = exogenous tGFP mitochondria, DAPI = cell nucleus. Scale bar = 10um.
Figure 8. Counting schematic for cell-type co-localization with tGFP mitochondria.  A.

Eight sections were analyzed on the left side of each injected spinal cord encompassing the contusion site, including both injection epicenters (red boxes), sections 400 µm rostral and 400 µm caudal to the injection epicenters (blue boxes), and the first section distal to the injection epicenter beyond the bolus where only punctate tGFP mitochondria were present (dashed black boxes). Injection epicenters are denoted by black stars, injection boluses are depicted as solid green ovals, and punctate tGFP stippling is represented by faded light green ovals. Note that the caudal injection sites are closer to the injury due to anatomical constraints of T12 laminectomy to effectively injure the L1/L2 spinal level.  B.

For each section analyzed, the ROI for cell counts was defined by drawing a horizontal line from the central canal to the lateral side and a vertical line from the central canal to the dorsal surface (red). Both the total number of labeled specific cell types and number of instances of tGFP co-localization with a specific cell type were counted within the ROI.
Figure 9. Cell-type co-localization of tGFP mitochondria across time. A. Total numbers of a given cell type were counted (white bars), as well as the number of cells co-localized with tGFP (black bars) for all ROI analyzed per spinal cord. B. Cell type co-localization was calculated as a percentage of total cells that had tGFP. Brain macrophages and pericytes had significantly higher co-localization percentages compared to oligodendrocytes at 24 and 48 hours. There were no differences in percent co-localization among cell types at 7 days. OX42 = brain macrophages, RECA = endothelial cells, PDGFR-B = pericytes, GFAP = astrocytes, CC1 = oligodendrocytes, tGFP = exogenous tGFP mitochondria. *p<0.05 vs OX42, # p<0.05 vs RECA, @ p<0.05 vs PDGFR-B, $ p<0.05 vs GFAP; (One-way ANOVA, Tukey’s multiple comparisons) n=4/group. Bars are means ± SEM.
Figure 10. Rostral-caudal cell-type incorporation of tGFP mitochondria over time. Co-localization was quantified within ROIs across eight different sections (numbered 1-8) in relation to injection epicenters. Top panel schematic of counted sections corresponds to the rostral-caudal counts of co-localization percentages at both injection epicenters (red boxes), sections 400 µm rostral and 400 µm caudal to the injection epicenters (blue boxes), and the first section where only punctate tGFP mitochondria were present (dashed black boxes). There were differences in co-localization percentages among location sites when analyzed within cell types. *p<0.05 vs section 8, #p<0.05 vs section 1, @p<0.05 vs section 3, $ p<0.05 vs section 5; (Two-way repeated measures ANOVA, Tukey’s multiple comparisons) n=4/group. Bars= mean ± SEM.
Figure 11. Histological assessments of tissue sparing at acute time points. Lesion volume (A), gray matter sparing (B), and white matter sparing (C) showed no apparent differences. Representative images show no apparent differences in tissue sparing at the injury epicenter among groups (D). n=1 vehicle, n=4 mito. Bars= mean ± SEM, Scale = 1mm for histological photos.
This paper has been peer-reviewed and accepted for publication, but has yet to undergo copyediting and proof correction. The final published version may differ from this proof.

**Figure 12. Hindlimb sensitivity and functional recovery following SCI and mitochondrial transplantation.**

A. Mechanical hypersensitivity following SCI and mitochondrial transplantation. Mitochondrial transplantation extended the duration of mechanical hypersensitivity after SCI. SCI similarly decreased mechanical threshold in all groups at 3 wk after SCI. At six weeks post injury, however, mechanical thresholds in the vehicle-injected and tGFP mitochondria-injected animals returned towards baseline levels, while those in the muscle mitochondria transplanted group remained significantly lower.

B. BBB scores for all animals were significantly decreased after SCI, with gradual increases over time after injury, though there were no significant differences among treatment groups. DPI = days post injury. Bars are means ± SEM. $ p<0.05$ vehicle vs baseline vehicle; *$ p<0.05$ tGFP mito vs baseline tGFP mito; #$ p<0.05$ muscle mito vs baseline muscle mito (Two-way repeated measures ANOVA, Tukey’s multiple comparisons) n=8-11/group.
**Figure 13. Tissue sparing analyses 6 weeks after injury and mitochondrial transplantation.** At 6 weeks post injury and injection, morphometric analyses were performed to find no differences among groups for total lesion volume (A), or for gray matter (B) or white matter (C) sparing at different spinal cord levels in relation to the injury epicenter. Bars are means ± SEM. (Two-way ANOVA) n=8-11/group.
Figure 14. Neuronal counts in the upper lumbar spinal cord after injury. A. ECCV stained section of the spinal cord showing the ROI analyzed including Lamina X and part of Lamina VII, which include putative central pattern generator neurons. CC= central canal. B. Representative images showing neurons in the ROI (dashed boxes) at 4mm rostral and -1mm caudal of the injury epicenter. There appeared to be more NeuN staining within the ROI of the mitochondria injected animals at 1mm caudal to the injury compared to either vehicle or muscle mitochondria injected groups. C. Stereological counts were performed on 11 equally spaced spinal cord sections, centered on the injury epicenter (epi). Each point on the graph is the estimated number of NeuN positive cells within subvolumes of spinal cord separated by 1mm length. NeuN positive cells were significantly higher in tGFP mitochondria injected groups compared to the other cohorts at 1mm caudal of injury epicenter, and also significantly higher compared to muscle mitochondria groups at 2 mm caudal of the injury epicenter. Bars are means ± SEM. * p<0.05 tGFP mito vs muscle mito; # p<0.05 tGFP mito vs vehicle, (2-way ANOVA, Tukey’s multiple comparisons) n=8-11/group.