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Short term voluntary exercise alters GDNF protein expression in rat spinal cord

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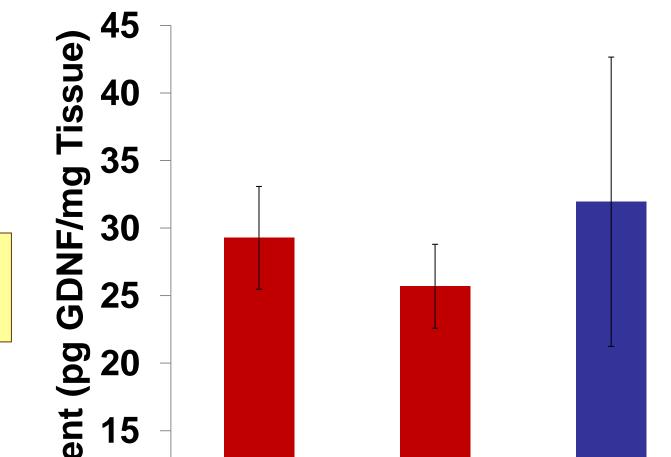
Α	bsti	ract

Glial cell line-derived neurotrophic factor (GDNF) is considered to be one of the most potent neurotrophic factors for motor neurons of the spinal cord. However, little is known about the response of GDNF in the spinal cord following exercise from healthy individuals. **PURPOSE:** Previous studies from our laboratory have shown that long term voluntary exercise (6 weeks and 6 months) has no effect on GDNF protein content in rat spinal cord. Therefore, the aim of the current study was to examine changes in GDNF protein content in the spinal cord following 2 weeks of voluntary exercise from adult and aged rats. METHODS: Male Sprague Dawley rats aged 6 months and 2 years were examined. The exercised animals were housed with continuous access to voluntary running wheels for 2 weeks. Age-matched sedentary control animals had no exposure to the voluntary wheels. GDNF protein content of the spinal cord was measured using an enzyme-linked immunosorbent assay (ELISA) and western blot analysis. Immunohistochemical analysis was performed to localize GDNF in nerve cells in the spinal cord. **RESULTS:** The 6-month-old animals that had undergone 2 weeks of voluntary exercise ran an average distance of 25,828.7 m whereas the 2-year-old animals ran an average distance of 1683.2 m. Total GDNF protein content in the spinal cord of 6 month old animals was significantly decreased following 2 weeks of voluntary exercise (1293.4 188.9 pg GDNF) as compared to sedentary control animals (2298.6 374.5 pg GDNF). Two weeks of voluntary exercise for the spinal cord of the 2 year old animals was approaching significant increase of total GDNF (p = 0.08) when compared to agematched sedentary control animals (1797.8 370.1 pg GDNF and 792.0 222.2 pg GDNF, respectively). **CONCLUSION:** Short term voluntary exercise may increase GDNF protein content in the spinal cord of aged individuals and may act as a measure to prevent motor neuron loss commonly associated with senescence. This work was supported by a grant from the Faculty Research and Crative Activities Award, Western Michigan University, NIH grant 1 R15 AG022908-01A2, NSF grant DBI 0552517 and MSU-KCMS

Introduction

One major problem with aging individuals is the continuous loss of somatic motor neurons that innervate skeletal muscles. Increased amounts of physical activity have been shown to prevent the loss of motor neurons in aging individuals (Kanda and Hashizume, 1998). Glial cell line-derived neurotrophic factor (GDNF) is one of the most potent trophic substances for motor neurons that innervate skeletal muscles. GDNF rescues somatic motor neurons from natural occurring cell death (Oppenheim et al., 2000), rescues motor neurons from axotomy-induced cell death (Oppenheim et al., 1995), and protects motor neurons from chronic degeneration (Corse et al., 1999). If GDNF protein expression is controlled by physical activity, then increasing levels of physical activity may reverse agerelated changes of the motor neuron structure and function.

6 weeks and 6 months of voluntary exercise does not change GDNF protein content in the Spinal Cord.



2 year old Control Spinal Cord

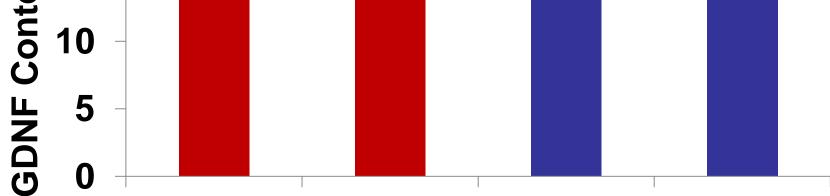
Summary

•2 weeks of voluntary exercise increases GDNF protein content in spinal cord of 6-month-old and 2year-old rats •Exercise may increase the size of motor neuron cell bodies in spinal cord •Exercise may change the isoform of GDNF protein found in the spinal cord •Short term exercise may increase GDNF protein content more than long term exercise

Hypothesis: Voluntary exercise will increase GDNF protein content of the spinal cord.

Aims

•Does GDNF protein content (bd) 3000 decrease in the spinal cord in aging animals? **(**) 2500 Does short term voluntary Tota 0005 exercise **GDNF** increase protein content in the spinal



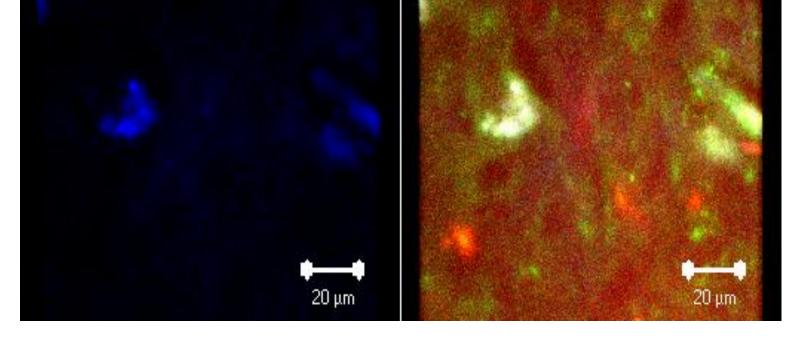
1 Year 1.5 Year **1.5 Year** 1 Year Exercise **Control Exercise** Control

Figure1. GDNF protein content in spinal cord measured via ELISA. Spinal cord was taken from both control animals aged 1 year and exercised animals with 6 weeks of voluntary exercise, and from control animals aged 1.5 years and exercised animals with 6 months of voluntary exercise. Data represent the mean SEM ($p \le 0.05$).

2 weeks of voluntary exercise increases **GDNF** protein content in the Spinal Cord of 6 month old animals.

*

4000 3500

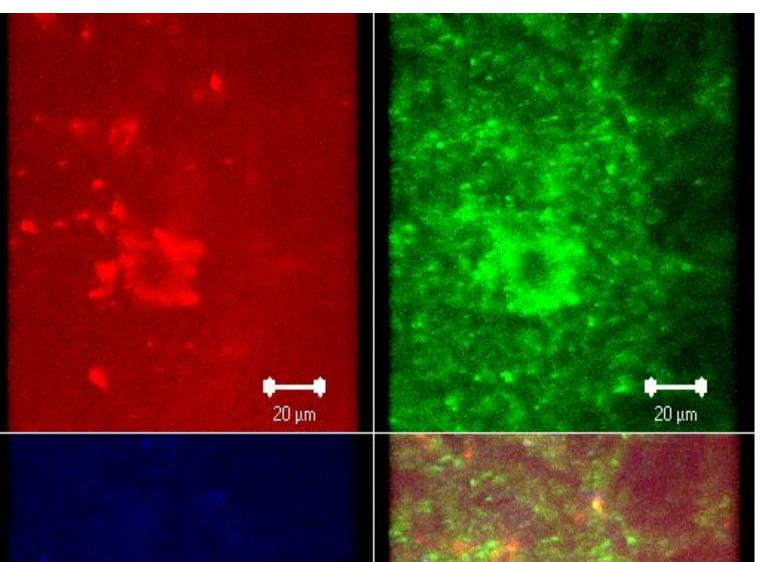


Results

20 un

Figure 4. Lumbar segments of the spinal cord from 2year-old control animals were fixed and bound with rabbit anti-GDNF (green), mouse anti-ChAT (red), and goat anti-CGRP (blue) and were viewed on a confocal microscope.

> **Exercised 2 year old Spinal Cord**



Conclusions

These results show that short term voluntary increases GDNF protein levels in the spinal cord of young and old animals. Immunohistochemical results suggest that motor neuron size is affected by exercise. Our western blot results suggest that expression of GDNF is higher following voluntary exercise in the spinal cord. Short term voluntary exercise may increase GDNF protein content in the spinal cord of aged individuals and may act as a measure to prevent motor neuron loss commonly associated with senescence.



Methods

Test Subjects:

23 male Sprague Dawley rats (Charles River Co.) aged 6 months (n = 12) and 2 years old (n = 11).

Sedentary (11 animals): Age-matched control animals had no running wheel access.

Exercised (12 animals): Animals were housed in individual cages with continuous access to voluntary running wheels for 2 weeks.

Determination of GDNF protein content:

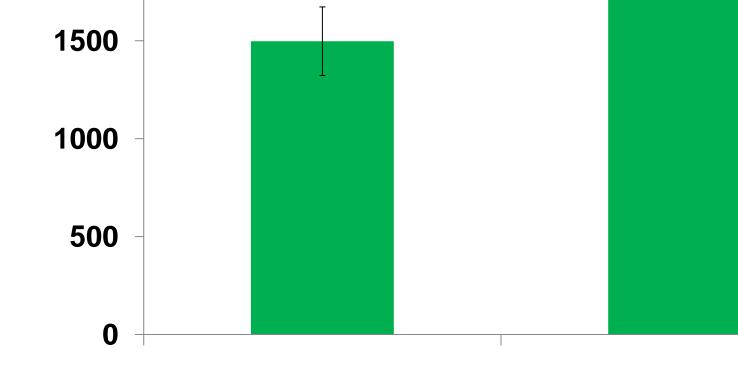
Spinal cords were removed and the lumbar region was processed. The sections smashed into a powder that was homogenized in a buffered salt solution containing protease inhibitors followed by centrifugation at 23,700 x g for 30 minutes. The supernatant was collected and GDNF protein content was measured using an enzyme-linked immunosorbant assay (ELISA). GDNF values were quantified using a known standard curve and expressed as pg of total.

Statistical Analysis:

Data are represented as the mean Standard Error of the Mean (SEM). Data were analyzed using one-way ANOVA to test for differences among the independent groups. Significance is established as $p \le 0.05$.

Immunohistochemistry:

Spinal cord sections were removed and fixed in 4% paraformaldehyde overnight at 4 C. Tissues were then washed in phosphate buffered saline (PBS) and placed in a 30% sucrose solution overnight at 4 C for cryoprotection. Tissues were stored at -80 C until ready for processing. The spinal cord was sectioned on the cryostat. Fixed tissues were bound with antibodies against GDNF, choline acetyltransferase (ChAT), and calcitonin gene related peptide (CGRP). Tissues were washed and then bound with secondary antibodies conjugated to fluorescent molecules (AlexaFluor 488, AlexaFluor 568 and AlexaFluor 647, respectively). Samples were placed on slides and viewed using a confocal microscope.



control

exercise

Figure 2. GDNF protein content in spinal cord measured via ELISA. Spinal cord was taken from both control animals aged 6 months and exercised animals with 2 weeks of voluntary exercise. Data represent the mean SEM ($p \le 0.05$).

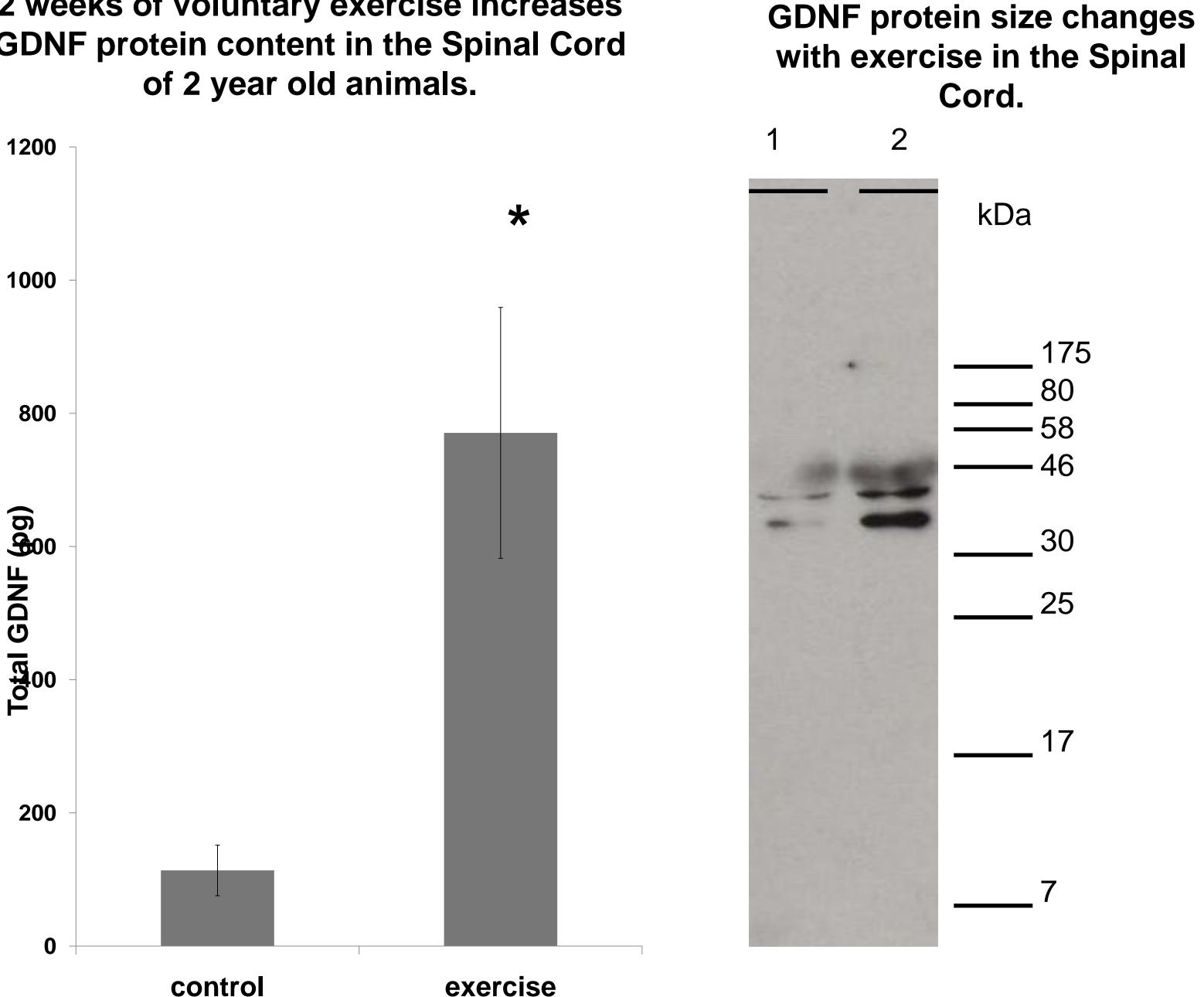
2 weeks of voluntary exercise increases **GDNF** protein content in the Spinal Cord of 2 year old animals.

DNF

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20 μm 20 μm

Figure 5. Lumbar segments of the spinal cord from 2year-old animals that were exercised for 2 weeks. Sections were fixed and bound with rabbit anti-GDNF (green), mouse anti-ChAT (red), and goat anti-CGRP (blue) and were viewed on a confocal microscope.



Acknowledgements

Special thanks to the WMU Biology Imaging Center. This work

Western Blot:

Homogenized tissue supernatant was used to determine protein concentration using a BCA protein assay with bovine serum albumin as the protein standard. Samples were prepared for loading into polyacrylamide gels by adding Laemmli 2X Loading buffer and boiled for 5 minutes. A protein ladder and a positive control of a GDNF standard were added to the gel. The gel was run at two different voltages from a power source, 100 Volts though the stacking gel and 200 Volts through the loading gel. Once all the samples migrated to the bottom, the gel was transferred to a polyvinylidene difluoride (PVDF) membrane where it was transferred by a power supply of 12 volts for 1 hour. Once the transfer finished, bands from the protein ladder were marked for visualization, and the PVDF membrane was blocked nonspecifically for 1 hour at 4 C under agitation. The membrane was incubated with a primary antibody against GDNF overnight at 4 C under agitation. The membrane was washed under agitation, then incubated with a HRP-conjugated secondary antibody for 1 hour at 37 C under agitation. The ECL detection kit was used to visualize the proteins and x-ray films were developed.

Figure 3. GDNF protein content in spinal cord measured via ELISA. Spinal cord was taken from both control animals aged 2 years and exercised animals with 2 weeks of voluntary exercise. Data represent the mean SEM ($p \le 0.05$).

Figure 6. Western blot using antibodies specific for GDNF. Spinal cord was taken from control animals aged 6 months, and 6 month old animals with 2 weeks of exercised. Lane 1 is the 6 month old control sample. Lane 2 is the 6 month old exercised animal.

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