

CASE REPORT

FES-rowing attenuates bone loss following spinal cord injury as assessed by HR-pQCT

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Neurologically motor complete spinal cord injury (SCI) presents a unique model of bone loss whereby specific regional sites are exposed to a complete loss of voluntary muscle-induced skeletal loading against gravity. This results in a high rate of bone loss, especially in the lower limbs where trabecular bone mass decreases by ~50–60% and cortical bone mass decreases by 25–34% before the rate of bone loss slows. These SCI-induced losses that are likely superimposed on continual age-related bone losses, increase the risk of low-impact fragility fracture. The fracture incidence 20 years post SCI is reported to be 4.6% per year. An intervention that effectively prevents, attenuates, or reverses bone loss is therefore highly desirable. We present a case study of an individual with chronic complete SCI, where bone loss has been attenuated following long-term functional electrical stimulation (FES)-rowing training. In this case study, we characterize the ultradistal tibia and ultradistal radius of the FES-rower with chronic complete SCI using high-resolution-peripheral quantitative computed tomography. These data are compared with a group of FES-untrained individuals with chronic complete SCI and to a normative non-SCI cohort. The evidence suggests, albeit from a single individual, that long-term FES-rowing training can attenuate bone loss secondary to chronic complete SCI. Indeed, key FES-rower's bone metrics for the ultradistal tibia more closely resemble normative age-matched values, which may have clinical significance since the majority of fragility fractures in chronic SCI occur in the lower extremities.

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Neurologically motor complete spinal cord injury (SCI) presents a unique model of bone loss whereby specific regional sites are exposed to a complete loss of voluntary muscle-induced skeletal loading against gravity. The initial loss of bone mineral content is estimated to approach 4% per month in trabecular sites, and 2% per month in cortical sites.¹ Trabecular bone mass is estimated to decrease by 48% of pre-injury values in the femur, reaching a new slower rate of loss by 3 years and 58% in the tibia by 5 years. Cortical bone mass decreases by 34% of pre-injury values in the femur by 5 years, and 25% in the tibia by 7 years.² These losses are likely superimposed on continual age-related bone loss.³ SCI-induced bone loss often leads to the development of osteoporosis, which increases the risk of low-impact fragility fracture.⁴ The fracture incidence 20 years post SCI is reported to be 4.6% per year.⁵ Thus, even active individuals with complete SCI at age 20 undergoing conventional standard of care have a very high likelihood of sustaining a fragility fracture in their lifetime, markedly affecting quality of life and increasing the risk of mortality.⁶ An intervention that effectively prevents, attenuates or reverses bone loss is therefore highly desirable.⁷

One such intervention under investigation is functional electrical stimulation (FES) rowing.⁸ In the present configuration, FES-rowing is an exercise that combines voluntary upper-limb concurrent with FES-assisted lower-limb exercise involving the quadriceps, hamstrings, tibialis anterior and triceps surae musculature. The training program, which involves three 30-min FES-rowing sessions and four 60-min FES-activated knee extension exercise sessions per week, has been previously described in detail.⁹ FES-rowing is well tolerated by males and females with complete and incomplete injury between C4 (with

a zone of partial preservation that permits a level of elbow flexion) and T12. In addition, there have been no reported medical complications related to FES-rowing training. Furthermore, any increased spasticity is managed by daily FES-training without the need for anti-spasmodic medications.

Gibbons *et al.*⁸ recently reported the potential to modulate bone loss in a male, age of 59 years with a T4 complete SCI following a road traffic accident, who is highly trained (> 12 years) in FES-rowing. The individual was 14 years post injury, with no history of musculoskeletal complications or medications known to affect bone (calcium, vitamin D or antiresorptive drugs). Although lower limb spasticity has increased since this individual's injury, daily FES-assisted exercise and passive standing successfully manages the condition without the need for anti-spasmodic medications. In Gibbons *et al.*⁸ bilateral trabecular bone density in the proximal tibia was reported using peripheral quantitative computed tomography (Stratec XCT3000, Pforzheim, Germany). The FES-rower's bone metrics were compared with data from a previous study involving a non-SCI cohort ($n=14$; mean age of 37 years) and a cohort of chronic SCI who were FES-untrained ($n=9$; mean age of 32 years; mean of 6.6 years post injury).¹⁰ Compared with the non-SCI cohort, the FES-rower had mean proximal tibia bone density in the non-dominant limb that was 0.57 s.d. lower, which is considered within the normal range based on the criterion of no more than 1.0 s.d. lower than the general population mean. The chronic SCI cohort had an average bone density that was 1.66 s.d. lower than the mean for non-SCI individuals. However, key limitations of that study include the lack of age-matched normative data, the relatively small number of

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Table 1. Comparison of subject demographics and ultradistal tibia and ultradistal radius metrics of FES-rower to normative non-SCI and chronic SCI cohorts

Variable	Normative non-SCI	Chronic SCI	FES-rower	FES-rower s.d. difference from	
	(50–59 yrs)	(FES-untrained)		non-SCI	Chronic SCI
<i>Demographics</i>					
Age (yrs)	55.3 ± 2.6	41.8 ± 7.8	59		
Sex	male n = 22	male/female n = 9	male		
Injury		T3-T12 AIS A & B	T4 AIS A		
Time post injury (yrs)		11.4 ± 9.4	14		
References	UCSF data	Wuermser <i>et al.</i>			
<i>Ultradistal tibia</i>					
Total.Ar (mm ²)	754.9 ± 103.1		612.9	-1.4	
Tb.Ar (mm ²)	602.5 ± 106.0		472.1	-1.2	
Total.vBMD (mg HA per cm ³)	326.3 ± 54.1	167.7 ± 64.7	305.1	-0.4	2.1
Tb.vBMD (mg HA per cm ³)	179.0 ± 36.7	67.5 ± 54.6	151.8	-0.7	1.5
Tb.N (1 per mm)	1.90 ± 0.39	1.09 ± 0.51	1.47	-1.1	0.7
Tb.Sp (mm)	0.468 ± 0.109	1.15 ± 0.82	0.593	1.1	0.7
Tb.Th (mm)	0.079 ± 0.013	0.04 ± 0.03	0.086	0.5	1.5
Ct.vBMD (mg HA per cm ³)	890.2 ± 64.8	809.8 ± 52.9	855.6	-0.5	0.9
Ct.Po (%)	4.2 ± 1.5		11.1	4.6	
Ct.Po.Dm (mm)	0.187 ± 0.021		0.264	3.7	
Ct.Th (mm)	1.51 ± 0.26	0.80 ± 0.28	1.52	0	2.4
Ct.Ar (mm ²)	148.0 ± 27.9	86.5 ± 30.5	144.4	-0.1	1.9
Ct.TMD (mg HA per cm ³)	984.1 ± 29.6		997.4	0.4	
Stiffness (kN mm ⁻¹)	209.3 ± 30.1		118.6	-3.0	
App Modulus (N mm ⁻²)	2099 ± 468		1554	-1.2	
Predicted failure load (N)	11177 ± 1257		6744	-3.5	
Cortical load fraction (distal) (%)	45.5 ± 8.5		56.0	1.2	
<i>Ultradistal radius</i>					
	n = 28				
Total.Ar (mm ²)	260.7 ± 41.7		223.7	-0.9	
Tb.Ar (mm ²)	336.7 ± 44.7		310.0	-0.6	
Total.vBMD (mg HA per cm ³)	329.7 ± 42.9		416.2	2.0	
Tb.vBMD (mg HA per cm ³)	178.6 ± 30.9		233.7	1.8	
Tb.N (1 mm ⁻¹)	2.01 ± 0.27		2.28	1.0	
Tb.Sp (mm)	0.434 ± 0.074		0.353	1.1	
Tb.Th (mm)	0.075 ± 0.010		0.085	1.0	
Ct.vBMD (mg HA per cm ³)	898.1 ± 56.5		961.7	1.1	
Ct.Po (%)	2.0 ± 1.3		2.4	-0.3	
Ct.Po.Dm (mm)	0.168 ± 0.015		0.164	0.3	
Ct.Th (mm)	1.02 ± 0.14		1.15	0.9	
Ct.Ar (mm ²)	73.6 ± 10.5		84.6	1.0	
Ct.TMD (mg HA per cm ³)	978.3 ± 30.0		994.0	0.5	
Stiffness (kN mm ⁻¹)	84.2 ± 16.0		76.4	-0.5	
App Modulus (N mm ⁻²)	1609 ± 402		1530	-0.2	
Predicted failure load (N)	4578 ± 614		4442	-0.2	
Cortical load fraction (distal) (%)	41.7 ± 6.3		40.8	-0.1	

Abbreviations: AIS, American Spinal Cord Injury Impairment Scale; A, motor and sensory complete injury; B, motor complete sensory incomplete injury; Ct.Ar, cortical area; Ct.Po, cortical porosity; Ct.Po.Dm, cortical porosity diameter; Ct.Th, cortical thickness; Ct.TMD, cortical tissue mineral density; Ct.vBMD, cortical vBMD; FES, functional electrical stimulation; HA, hydroxyapatite; SCI, spinal cord injury; T, thoracic; Total.Ar, total area; Tb.Ar, trabecular area; Tb.N, trabecular number; Tb.Sp, trabecular separation; Tb.Th, trabecular thickness; Total.vBMD, total volumetric bone mineral density; Tb.vBMD, trabecular vBMD; UCSF, University of California San Francisco. Note (1): The s.d. difference values in the right hand columns are calculated as the difference between the FES-rower mean and the normative non-SCI mean in units of the normative non-SCI s.d., and the difference between the FES-rower mean and the chronic SCI mean in units of the chronic SCI s.d. Note (2): Positive s.d. values indicate that the FES-rower values are greater than, and negative s.d. values less than, the reference cohort values. Note (3): Boxed values highlight age and time post injury comparisons, and the FES-rower s.d. difference values which are within 1 s.d. of non-SCI and ≥ 1.5 s.d. greater than chronic SCI.

subjects in the comparison cohorts and the lack of data on bone microstructure and strength.

The aim of the present study, was to measure bone density and microstructure, and to estimate strength, of the ultradistal radius and ultradistal tibia of a highly trained FES-rower with chronic complete SCI and to compare those measurements to a chronic SCI cohort, who were FES-untrained,¹¹ and normative age- and sex-matched non-SCI cohort (UCSF data including a subset of data previously reported¹²).

The same individual with chronic complete SCI reported by Gibbons *et al.* was the FES-rower in this study. During the 24-months before imaging, the FES-rower performed an average of three weekly 30-min rows at 30 strokes per minute, exposing the lower limbs to ~2700 loading cycles per week. The ultradistal radius and ultradistal tibia of the FES-rower was imaged using a high-resolution-peripheral quantitative computed tomography (HR-pQCT) scanner (XtremeCT I, Scanco Medical AG, Bruttisellen, Switzerland) and data were analyzed using established protocols.

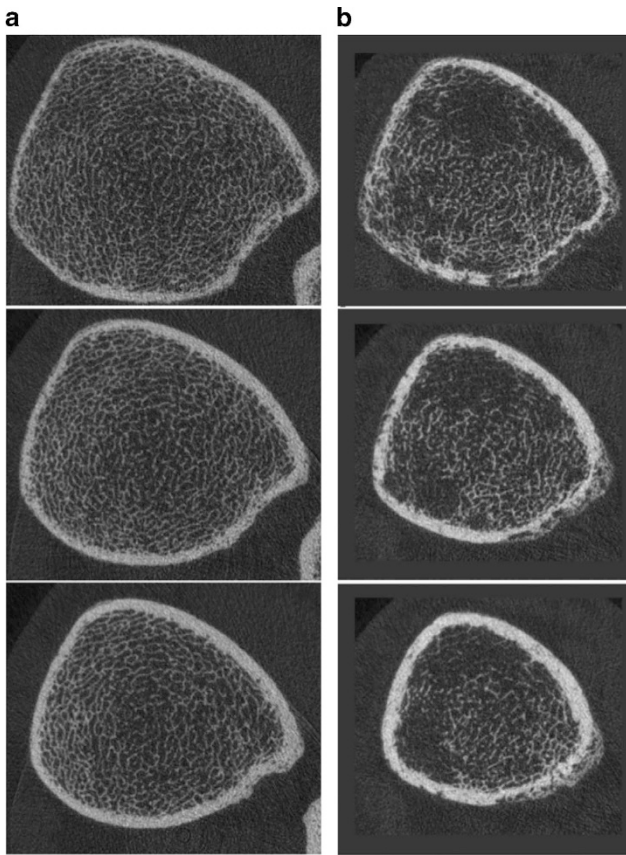


Figure 1. Representative HR-pQCT images from the most distal (top row), middle, and most proximal (bottom row) slice of the ultradistal tibia. **(a)** Normative 57-year-old non-SCI white male. **(b)** FES-rower. Images are equally scaled in both size and grayscale intensity.

The standard in-vivo imaging protocol (60 kVP, 900 mA, 100 ms integration time, 126 mm field of view) was used. The non-dominant side (before injury) was scanned. The extremity was immobilized using a carbon fiber cast anchored in the scanner to minimize motion. A single anteroposterior scout radiograph was obtained to allow correct positioning of the tomographic scan volume. A total scan volume of 9.02 mm length (110 slices) was then acquired starting 9.5 mm or 22.5 mm proximal to the joint line in the distal radius and tibia, respectively. Images were reconstructed to a 1536×1536 matrix, allowing for a final nominal, isotropic resolution of $82 \mu\text{m}$. To calculate densitometric bone parameters, image attenuation values were calibrated against the attenuation values derived from a standardized hydroxyapatite phantom. Analysis was performed using the standard clinical evaluation protocol in Image Processing Language (IPL v5.08b, Scanco Medical AG), as described in detail previously.¹³ Cortical parameters were assessed using an extended cortical bone analysis that provides direct calculation of cortical thickness as well as measures of porosity.^{14,15} Linear micro-finite element analysis was performed to measure apparent biomechanical properties (Scanco FE Software Version 1.12, Scanco Medical AG) using previously described techniques.^{16–18} Stiffness, apparent modulus, cortical compartment load fraction and failure load estimate were computed for each model.

Individual and group demographics are presented in Table 1. At the tibia, the FES-rower metrics were 0.7–2.4 s.d. better than the chronic SCI cohort means. However, while FES-rower densitometric and trabecular parameters were within ~ 1 s.d. of normative non-SCI age- and sex-matched control cohort means, cortical

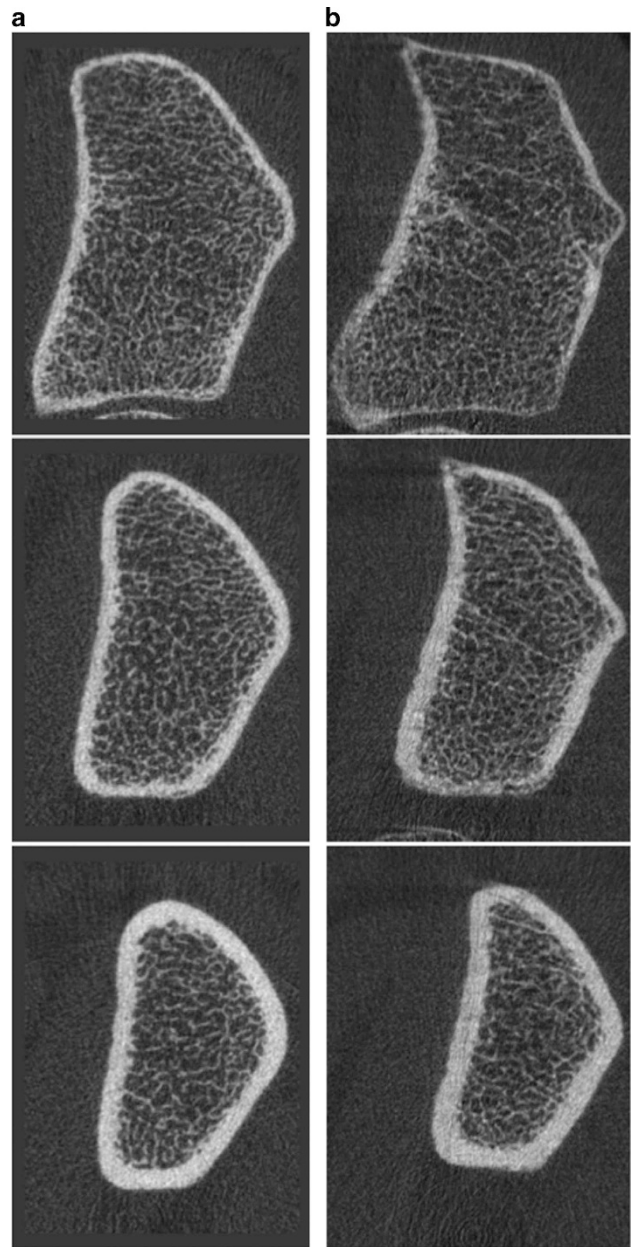


Figure 2. Representative HR-pQCT images from the most distal (top row), middle, and most proximal (bottom row) slice of the ultradistal radius. **(a)** Normative 57-year-old non-SCI white male. **(b)** FES-rower. Images are equally scaled in both size and grayscale intensity.

porosity and mean pore diameter were higher (+4.6 and +3.7 s.d., respectively) and biomechanical indices lower (-1.2 to -3.5 s.d.), which imply lower bone strength (Table 1). Figure 1 shows representative HR-pQCT ultradistal tibia images of a normative non-SCI age- and sex-matched individual (column A) compared to the FES-rower (column B). The images show increased heterogeneity of structure and density within both trabecular and cortical compartments in the FES-rower. It is important to note that the two individuals are not body-size matched. At the radius, FES-rower densitometric and trabecular metrics were 0.9–2.0 s.d. better than the normative non-SCI cohort means, while cortical and biomechanical indices were within 1 s.d. (Table 1). Figure 2 shows representative HR-pQCT ultradistal radius images of a normative non-SCI age- and sex-matched individual (column A) compared with the FES-rower (column B). The images show similar

trabecular and cortical bone uniformity between the non-SCI individual and the FES-rower. No radius data exist for the chronic SCI cohort.

We aimed to characterize the density, microstructure and strength of the ultradistal radius and tibia of a highly trained FES-rower with chronic complete SCI and compare these data to a chronic SCI cohort, who were FES-untrained, and normative non-SCI age- and sex-matched cohort. At the tibia, the majority of trabecular and cortical parameters for the tibia in the FES-rower were within ~ 1 s.d. of age- and sex-matched non-SCI controls, which suggests bone loss has been mediated in the FES-rower compared to the chronic SCI cohort. However, while habitual FES-rowing biomechanical loading generally appears to have attenuated bone loss in the tibia, bone strength is lower than age- and sex-matched non-SCI controls likely due to the markedly larger percent cortical porosity (+4.6 s.d.) and mean pore diameter (+3.7 s.d.). This suggests that the mechanobiological response that controls cortical porosity in the tibia is in some way different from the response for other bone parameters. At present, the nature of that difference is unknown. Moreover, the high cortical porosity and pore diameter values may explain the markedly lower estimated structural stiffness and failure load in the tibia of the FES-rower. However, the HR-pQCT assessment of a larger number of individuals who have undergone long-term FES-rowing training is required to more fully understand this issue. At the radius, the superior key bone metrics in the FES-rower compared with normative non-SCI controls is consistent with the greater daily bone stimulus for an active manual wheelchair user.

For this single participant age 59 years and over 14 years post injury, a number of the key FES-rower's bone metrics at the ultradistal tibia site more closely resemble normative non-SCI values. This may have clinical significance, since the vast majority of fragility fractures in chronic SCI occur in the lower extremities.⁵ Future studies that investigate the effect of long-term FES-rowing training on a large cohort of FES-naïve people with complete SCI are recommended.

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COMPETING INTERESTS

The authors declare no conflict of interest.

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