The effects of poliomyelitis on motor unit behavior during repetitive muscle actions: a case report

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Abstract

Background: Acute paralytic poliomyelitis is caused by the poliovirus and usually results in muscle atrophy and weakness occurring in the lower limbs. Indwelling electromyography has been used frequently to investigate the denervation and innervation characteristics of the affected muscle. Recently developed technology allows the decomposition of the raw surface electromyography signals into the firing instances of single motor units. There is limited information regarding this electromyographic decomposition in clinical populations. In addition, regardless of electromyographic methods, no study has examined muscle activation parameters during repetitive muscle actions in polio patients. Therefore, the purpose of this study was to examine the motor unit firing rates and electromyographic amplitude and center frequency of the vastus lateralis during 20 repetitive isometric muscle actions at 50% maximal voluntary contraction in healthy subjects and one patient that acquired acute paralytic poliomyelitis.

Case presentation: One participant that acquired acute type III spinal poliomyelitis (Caucasian male, age = 29 yrs) at 3 months of age and three healthy participants (Caucasian females, age = 19.7 ± 2.1 yrs) participated in this study. The polio participant reported neuromuscular deficiencies as a result of disease in the hips, knees, buttocks, thighs, and lower legs. None of the healthy participants reported any current or ongoing neuromuscular diseases or musculoskeletal injuries.

Conclusion: An acute bout of poliomyelitis altered motor unit behavior, such as, healthy participants displayed greater firing rates than the polio patient. The reduction in motor unit firing rates was likely a fatigue protecting mechanism since denervation via poliomyelitis results in a reduction of motorneurons. In addition, the concurrent changes in motor unit firing rates, electromyography amplitude and frequency for the polio participant would suggest that the entire motorneuron pool was utilized in each contraction unlike for the healthy participants. Finally, healthy participants exhibited changes in all electromyographic parameters during the repetitive muscle actions despite successfully completing all contractions with only a slight reduction in force. Thus, caution is warranted when quantifying muscular fatigue via motor unit firing rates and other electromyographic parameters since the parameters changed despite successful completing of all contractions with only a moderate reduction in strength in healthy subjects.

Keywords: Electromyography, Firing rates, Isometric, Motorneuron, Vastus lateralis

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Background

Acute paralytic polio is a disease caused by the poliovirus that typically occurs in young children. The poliovirus spreads along nerve pathways and primarily destroys motorneurons of the anterior horns [1]. The poliovirus usually results in asymmetrical paralysis with muscle atrophy and weakness occurring in the lower limbs more so than the upper limbs [2]. Numerous studies have used indwelling electromyography (EMG) to investigate the denervation and reinnervation characteristics of the affected muscle (i.e., motor unit number and amplitude, motor unit firings, conduction velocity, etc.) [3-8]. For example, Larsson et al. [7] reported that healthy participants had greater motor unit (MU) firing rates of the tibialis anterior muscle than the poliomyelitis participants during slowly increasing isometric voluntary muscle actions. In addition, McComas et al. [9] reported a reduction in motor unit number after an acute bout of poliomyelitis at twice the rate than healthy subjects aged > 60 years, whereas, Herda and Cooper [10] reported a reduction in MUs recruited above 20% of maximal voluntary contraction (MVC) for a poliomyelitis patient in comparison to healthy participants.

Recently, De Luca et al. [11] developed technology able to decompose the raw surface EMG signals recorded during isometric muscle actions into its constituent MU action potential trains with the Precision Decomposition (PD) III algorithm. The resulting output allows for the examination of the firing instances for each detected MU. The PD III algorithm has typically detected 30-40 MUs and occasionally up to 60 MUs during a single isometric trapezoid muscle action at various force levels (5 – 100% MVC). Previously, De Luca et al. [11] and Nawab et al. [12] validated the PD III algorithm and reconstruct-and-test (used for testing accuracy of MU action potential trains) algorithms. This decomposition technology has been used to examine the MU control scheme, which states that lower threshold MUs have higher firing rates than higher threshold MUs during isometric muscle actions [13-15]. However, there is limited information available regarding the use of EMG decomposition in clinical populations [10,16]. It would be expected that an acute bout of paralytic polio would influence the overall MU control scheme and be detectable by the EMG decomposition technique developed by De Luca et al. [11], which was supported by Herda and Cooper [10].

Previous studies have utilized repetitive isometric muscle actions to further elucidate mechanisms of fatigue on MU behavior [17,18]. For example, Carpentier et al. [18] indicated that central drive intensified, as measured by EMG amplitude, while MU discharge rates decreased progressively during the repetitive muscle actions of the first dorsal interosseous at 50% MVC. In addition, the authors reported that the level of change in MU behavior was related to recruitment threshold (low- versus high-threshold), which was also supported by Farina et al. [17]. To date, it is unknown the influence of poliomyelitis on MU firing rates and other EMG parameters (amplitude and center frequency) during repetitive muscle actions. Since polio patients have far less motorneurons as a result of the denervation and reinnervation process, it is conceivable that their MU behavior during the repetitive tasks would be altered in comparison to healthy subjects. Therefore, the purpose of this study was to examine the MU control scheme and EMG amplitude and frequency of the vastus lateralis (VL) during 20 repetitive isometric muscle actions at 50% MVC for healthy individuals and one individual that acquired acute paralytic poliomyelitis.

Case presentation

In this case report, we present one participant that acquired acute type III spinal poliomyelitis (mean ± standard deviation [SD], Caucasian male, age = 29 yrs, weight = 63.4 kg, height = 150.7 cm) at 3 months of age and three healthy participants (Caucasian females, age = 19.7 ± 2.1 yrs, weight = 67.5 ± 9.4 kg, height = 170.7 ± 2.3 cm). Previous research has reported no sex-related differences in EMG amplitude, EMG frequency, or MU behavior during fatiguing contractions [19-25].

The participant with acute type III spinal poliomyelitis (PO) reported neuromuscular deficiencies as a result of disease in the hips, knees, buttocks, thighs, and lower legs. None of the healthy (HE) participants reported any current or ongoing neuromuscular diseases or musculoskeletal injuries. Each participant signed a written informed consent document. This study was approved by the Human Subjects Committee – Lawrence at the University of Kansas.

Strength testing

Each participant was seated with restraining straps over the pelvis, trunk, and contralateral thigh, and the lateral condyle of the femur was aligned with the input axis of the Biodex System 3 isokinetic dynamometer (Biodex Medical Systems, Inc., Shirley, NY, USA). All isometric leg extensor strength assessments were performed on the right leg at a flexion of 90°. Isometric strength for the leg extensors muscles was measured using the force signal from a load cell (LC402, Omegadyne, Inc., Sunbury, OH, USA) that was fitted to the isokinetic dynamometer. During the experimental trial, participants completed a warm-up consisting of three to five voluntary isometric muscle actions from 30%-80% MVC. Participants then performed three isometric MVCs with strong verbal encouragement for motivation followed by twenty repetitive submaximal isometric trapezoid muscle actions at 50% MVC. The trapezoid trajectory contained five segments: a 5 sec quiescent period for baseline noise calculation, an up-ramp increased at a rate of 10% MVC/sec, a
constant force of 50% MVC for 12 sec, a down-ramp decreased at 10% MVC/sec, and a 3 sec quiescent period. Therefore, the duration of each repetitive trial lasted 22 sec and subjects were given a rest period of 8 – 9 sec between trials. Each participant was instructed to maintain their force output as close as possible to the target force presented digitally in real time on a computer monitor. Following the repetitive muscle actions, participants performed an additional MVC.

Electromyography signal detection and processing
Surface EMG signals were recorded from the VL using a 5 pin surface array sensor (Delsys, Boston, MA). Prior to sensor placement, the surface of the skin was prepared by shaving, removing superficial dead skin with adhesive tape, and sterilized with an alcohol swab. In addition, EMG signals must successfully pass the Delsys Signal Quality Check (SQC) prior to the recording of EMG for decomposition. The SQC requires the signal to noise ratio to be > 2, baseline noise < 4.8 μV root mean square, and line interference < 1.0 (normalized magnitude power spectrum at 50/60 Hz) (Delsys, Inc., User’s Guide). The surface EMG sensor was placed over the belly of the VL and fixed with adhesive tape while the reference electrode was placed over the patella. Refer to De Luca et al. [26] for detailed information regarding the signal processing of the EMG signals. Decomposition techniques (PD III algorithm) were applied to the surface EMG signals to extract action potentials and the firing events of single MUs [11,12,14,27]. Furthermore, the accuracy level of each MU was assessed by a reconstruct-and-test procedure [11]. Only MUs with > 90% accuracy were included for analysis. For each MU, the recruitment thresholds (expressed relative to MVC) and the mean firing rate (MFR) at the targeted contraction level (pulses per second [pps]) were analyzed. The MFR was calculated as the average value of the mean firing rate trajectory during steady force [26]. Linear regressions were performed on the MFR versus recruitment threshold relationships for each muscle action. Slopes and y-intercepts were calculated for each linear regression model. The y-intercept values specify a maximal firing rate (pps) sustainable by the lowest threshold MUs.

In addition, EMG amplitude and mean power frequency (MPF) was investigated. The signals were collected at a frequency of 20 kHz and bandpass filtered (fourth-order Butterworth) at 10-500 Hz. Signal processing was performed with custom programs written with LabVIEW software (Version 11.0, National Instruments, Austin, TX). The amplitude for the EMG signal was calculated with root mean square (RMS). For the isometric muscle actions, EMG RMS and MPF were calculated by averaging the values across the four EMG channels during the entire 12 sec targeted contraction force. These data were normalized to the EMG RMS and MPF values corresponding to the first 50% repetitive muscle action.

Results
There was a total of 383 MUs analyzed for the HE participants with 80 MUs analyzed for the PO participant (Table 1). For the pre- and post-MVCs, peak force values decreased 10.3% and 28.0% for the HE (548.67 ± 121.01 N, 492.00 ± 147.15 N) and PO (37.80 N, 27.20 N), respectively. The normalized (i.e., 1st repetition = 100%) EMG RMS was 117.68 ± 9.36% and 126.97 ± 23.13% while the MPF was 89.90 ± 1.90% and 88.19 ± 4.06% for the HE during the 10th and 20th repetition, respectively (Figure 1). For the PO, the normalized EMG RMS was 93.37% and 101.71% while the MPF was 101.78% and 100.00% during the 10th and 20th repetitive contraction (Figure 1). For the y-intercepts, the mean ± SD for the HE were 33.42 ± 2.41 pps, 29.27 ± 2.12 pps, and 29.12 ± 1.83 pps for the 1st, 10th, and 20th repetition and the y-intercepts for the PO were 25.64 pps, 23.48 pps, and 25.26 pps for the 1st, 10th, and 20th repetition (Figure 2). For the slopes, the mean ± SD for the HE were -0.47 ± 0.05, -0.34 ± 0.02, and -0.41 ± 0.11 for the 1st, 10th, and 20th repetition and the slopes for the PO were -0.42, -0.26, and -0.41 (Figure 2).

Discussion
Previously, Larsson et al. [7] reported differences in MU firing rates between healthy and prior polio patients during “slowly increasing isometric voluntary contractions” (p.190). The y-intercept from the MFR versus recruitment

Table 1 Motor unit (MU) number, slope, y-intercept and R values for the healthy (HE) and acute paralytic poliomyelitis (PO) participants and repetitive (Rep) contractions (1st, 10th, and 20th) for the mean firing rate (MFR) versus recruitment threshold

| Subject | MU Number | Rep# | Slope | Y-intercept | R  |
|---------|-----------|------|-------|-------------|----|
| HE      | 28        | 1st  | -0.452| 34.697      | 0.925|
|         | 36        | 10th | -0.356| 31.632      | 0.947|
|         | 41        | 1st  | -0.526| 30.641      | 0.858|
|         | 55        | 10th | -0.323| 27.535      | 0.874|
|         | 51        | 20th | -0.523| 30.283      | 0.914|
|         | 43        | 1st  | -0.439| 34.926      | 0.862|
|         | 50        | 10th | -0.345| 28.641      | 0.909|
|         | 53        | 20th | -0.313| 27.011      | 0.866|
|         | 24        | 1st  | -0.418| 25.642      | 0.798|
|         | 23        | 10th | -0.261| 23.478      | 0.755|
|         | 33        | 20th | -0.412| 25.257      | 0.836|

All relationships were significant (P < 0.05).
threshold relationships theoretically represents a maximal firing rate sustainable by the lowest threshold MUs [14]. In the present study, the MFRs for the HE were higher (y-intercept range: 33.42 ± 2.41 – 29.12 ± 1.83 pps) than the PO (y-intercept range: 25.64 - 23.48 pps) for each repetition (Figure 2). Negative slopes were reported from the linear regressions for each participant and contraction and, thus, earlier recruited MUs tended to have greater firing rates than the later recruited MUs. Referring to Table 1 and Figure 2, there were no observed differences between the slopes between PO and HE participants. Therefore, the inverse relationship between the firing rate and recruitment threshold typically reported in healthy muscle was present for the PO muscle [14]. The overall MU control scheme (i.e., negative slopes) was similar between the PO and HE participants, however, the MFRs were less (i.e., y-intercepts) for the PO than the HE participants. High-threshold MUs are typically described as fast fatiguing for the VL. De Luca and Hostage [14] suggested that if the high-threshold MUs discharged at relatively higher rates, high-threshold MUs would fatigue more quickly and force will not be maintained. Since it has been reported that there are far less motorneurons in the PO than HE participants as a result of denervation [5,8], the lower MFRs throughout the force spectrum for the PO is possibly a fatigue protecting mechanism.

EMG RMS is a global measure of MU recruitment and firing rates while EMG MPF reflects the average muscle fiber conduction velocity [28]. Classically, muscle fatigue has been monitored with these EMG parameters and indicators of fatigue are believed to be evident by increases in RMS and decreases in MPF [29]. An increase in EMG RMS is the result of synchronization of active MUs and the recruitment of new MUs while the decrease in MPF has been primarily attributed to a reduction in muscle fiber conduction velocity [30]. For the 1st to 10th repetition, the HE had an increase (17.7%) in EMG RMS while EMG MPF declined (10.0%). Concurrently, the y-intercepts from the MFR versus recruitment threshold relationships decreased (12.4%) for the HE participants and, thus, would suggest that the HE participants recruited additional MUs and/or had greater MU synchronization during the 10th repetition in comparison to the 1st repetition. In contrast, the y-intercepts from the MFR versus recruitment threshold relationships and EMG RMS decreased (8.4% and 6.6%) simultaneously while EMG MPF (+1.8%) remained stable from the 1st to 10th repetition for the PO participant. For the 10th to 20th repetition, there was relatively no change in the y-intercepts (0.5%) or EMG MPF (-1.7%), however, EMG RMS increased 9.3% for the HE. For the PO, the y-intercept from the MFR versus recruitment threshold and EMG RMS returned to baseline levels (i.e., increased) during the 20th repetition. The EMG parameters indicated that MU behavior differed between the PO and HE during the repetitive muscle actions. Subsequently, MU firing rates and EMG RMS responded in a similar manner for the PO, in contrast, MU firing rates and EMG MPF decreased while EMG RMS increased during the repetitive muscle actions for the HE. Future studies may want to consider normalizing these parameters to an MVC rather than a submaximal muscle action as in the present study to further elucidate neural mechanisms of fatigue.

Previously, Larsson et al. [7] reported that polio patients used muscle fibers in an all or none manner during slowly increasing isometric voluntary muscle actions. In the present study, the changes in EMG RMS mirrored the
changes in the y-intercepts from the MFR versus recruitment threshold relationships with relatively no change in MPF for the PO, which tentatively suggested that all MUs were recruited during each muscle action at 50% MVC. Since EMG RMS is the concurrent representation of MU recruitment and firing rates, the increase in EMG RMS with a corresponding decrease in MFR would suggest that HE relied on recruitment of additional MUs and/or synchronization of MU during the repetitive tasks. However, the mirrored changes between EMG RMS and MFR for the PO suggested that no additional MUs were recruited nor was there an increase in
MU synchronization during the repetitive tasks. Interestingly, HE participants were able to complete all 20 contractions with a moderate (10.3%) reduction in post-MVC strength. Although the PO had fewer changes in all parameters measured (MU MFR, EMG RMS and MPF), the decrease in post MVC for the PO was much greater than the HE (28.0% vs. 10.3%). Thus, EMG parameters changed during the repetitive tasks despite a lack of fatigue in the HE as indicated by the successful completion of the tasks with only a moderate reduction in strength.

Conclusions

In summary, the denervation and reinnervation process as a result of an acute bout of poliomyelitis altered MU behavior in comparison to the HE participants. Specifically, the PO participants had lower MFRs and the y-intercepts from the MFR versus recruitment threshold relationships mirrored changes in EMG RMS from the 1st to 20th repetition, unlike the HE participants. Thus, suggesting that all MUs were recruited during each contraction for the PO participant. In addition, caution is warranted when examining EMG RMS and MPF and MU MFR behavior as indicators of muscular fatigue during repetitive muscle actions since the parameters changed in the HE despite successfully completing all muscle actions with only a moderate decrease in strength. Future research should examine the influence of high-intensity exercise training on the fatigability of polio patients as it is evident that the alterations in MU control scheme provide a mechanism to respond to such training despite the denervation and reinnervation process.

Consent

Written informed consent was obtained from all participants for publication of this Case Report. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

Abbreviations

EMG: Electromyography; HE: Healthy participant; MFR: Mean firing rate; MPF: Mean power frequency; MU: Motor Unit; MVC: Maximal voluntary contraction; PO: Polio participant; PD: Precision decomposition; PPS: Pulses per second; RMS: Root mean square; SQC: Signal quality check; SD: Standard deviation; VL: Vastus lateralis.

Competing interests

The authors declare that they have no competing interests.

Authors’ contributions

MT carried out the statistical analysis and interpretation of data, and drafted the manuscript. TH participated in the design of the study, data collection, assisted in the analyses and interpretation of data, and helped revise the manuscript. MT assisted in the design of the study, data collection, and interpretation of data. All authors read and approved the final manuscript. No funding was provided for the authors or for the manuscript.

Acknowledgements

We are grateful for the contributions of the undergraduate research assistants in the Neuromechanics Laboratory for their assistance in performing the experiments.

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Received: 11 April 2014 Accepted: 2 September 2014 Published: 6 September 2014

References

1. Swan C. The anatomical distribution and character of the lesions of Poliomyelitis. Australian J Exp Biol Med Sci 1939, 17:345–364.
2. Love S, Wiley CA. Poliomyelitis and polioencephalitis. In Greenfield’s Neuropathology, Volume 1. Edited by Love DNLAxW. S. London: Hodder Arnold; 2008:1285–1291.
3. Borg K, Borg J, Dhoot G, Edstrom L, Grimby L, Thonnell LE. Motoneuron firing and isomyosin type of muscle fibres in prior polio. J Neurol Neurosurg Psychiatry 1989, 52:1141–1148.
4. Ravits J, Hallett M, Baker M, Nilsson J, Dalakas M. Clinical and electromyographic studies of postpoliomyelitis muscular atrophy. Muscle Nerve 1990, 13:567–674.
5. Roeleveld K, Sandberg A, Stalberg EJ, Stegeman D: Motor unit size estimation of enlarged motor units with surface electromyography. Muscle Nerve 1998, 21:878–885.
6. Borg K, Borg J. Conduction velocity and refractory period of single motor nerve fibres in antecedent poliomyelitis. J Neurol Neurosurg Psychiatry 1987, 50:443–446.
7. Larsson L, Li X, Toltbäck A, Grimby L: Contractile properties in single muscle fibres from chronically overused motor units in relation to motoneuron firing properties in prior polio patients. J Neurol Sci 1995, 132:182–192.
8. Stalberg E, Grimby G: Dynamic electromyography and muscle biopsy changes in a 4-year follow-up: study of patients with a history of polio. Muscle Nerve 1995, 18:699–707.
9. McComas AJ, Quah C, Griggs RC: Early and late losses of motor units after poliomyelitis. Brain 1997, 120:1415–1421.
10. Herda TJ, Cooper MA: Examination of motor unit control properties of the vastus lateralis in an individual that had acute paralytic poliomyelitis. J Clin Neurophysiol 2014. In Press.
11. De Luca CJ, Adam A, Wotiz R, Gilmore LD, Nawab SH: Decomposition of surface EMG signals. J Neurophysiol 2006, 96:1646–1657.
12. Nawab SH, Chang SS, De Luca CJ: High-yield decomposition of surface EMG signals. Clin Neurophysiol 2010, 121:1602–1615.
13. De Luca CJ, Le Fever RS, McCue MP, Xenakis AP: Behaviour of human motor units in different muscles during linearly varying contractions. J Physiol 1982, 329:113–128.
14. De Luca CJ, Hostage EC: Relationship between firing rate and recruitment threshold of motoneurones in voluntary isometric contractions. J Neurophysiol 2010, 104:1034–1046.
15. Kamen G, Sison SV, Du CC, Patten C: Motor unit discharge behavior in older adults during maximal-effort contractions. J Appl Physiol 1995, 79:1908–1913.
16. Suresh N, Xiaoyan L, Ping Z, Rymer WJ: Examination of motor unit control properties in stroke survivors using surface EMG decomposition: a preliminary report. In Engineering in Medicine and Biology Society, EMBC, 2011 Annual International Conference of the IEEE, Aug. 30 2011-Sept. 3 2011. I n Engineering in Medicine and Biology Society, EMBC. IEEE, 2011:8243–8246.
17. Farina D, Holobar A, Gazzoni M, Zazula D, Merletti R, Enoka RW: Adjustments differ among low-threshold motor units during intermit- tent, isometric contractions. J Neurophysiol 2009, 101:350–359.
18. Carpenter A, Duchateau J, Hainaut K: Motor unit behaviour and contractile changes during fatigue in the human first dorsal intersosseus. J Physiol 2001, 534:903–912.
19. Cramer JT, Housh TJ, Evetovich TK, Johnson GO, Ebersole KT, Perry SR, Bull AJ: The relationships among peak torque, mean power output, mechanomyography, and electromyography in men and women during maximal, eccentric isokinetic muscle actions. Eur J Appl Physiol 2002, 86:226–232.
20. Cramer JT, Housh TJ, Weir JP, Johnson GO, Beningham JM, Perry SR, Bull AJ: Gender, muscle, and velocity comparisons of mechanomyographic and electromyographic responses during isokinetic muscle actions. Scand J Med Sci Sports 2004, 14:116–127.
21. Pincivero DM, Green RC, Mark JD, Campy RM: Gender and muscle differences in EMG amplitude and median frequency, and variability during maximal voluntary contractions of the quadriceps femoris. J Electromyogr Kinesiol 2000, 10:189–196.

22. Carneiro JG, Goncalves EM, Camata TV, Altimari JM, Machado MV, Batista AR, Guerra Junior G, Moraes AC, Altimari LR: Influence of gender on the EMG signal of the quadriceps femoris muscles and performance in high-intensity short-term exercise. Electromyogr Clin Neurophysiol 2010, 50:326–332.

23. Defreitas JM, Beck TW, Ye X, Stock MS: Synchronization of low- and high-threshold motor units. Muscle Nerve 2014, 49:575–583.

24. Stock MS, Beck TW, Defreitas JM: Effects of fatigue on motor unit firing rate versus recruitment threshold relationships. Muscle Nerve 2012, 45:100–109.

25. Stock MS, Beck TW, DeFreitas JM, Ye X: Sex comparisons for relative peak torque and electromyographic mean frequency during fatigue. Res Q Exerc Sport 2013, 84:343–352.

26. De Luca CJ, Contessa P: Hierarchical control of motor units in voluntary contractions. J Neurophysiol 2012, 107:178–195.

27. Chang SS, De Luca CJ, Nawab S: Aliasing rejection in precision decomposition of EMG signals. Conf Proc IEEE Eng Med Biol Soc 2008, 2008:4972–4975.

28. Lindstrom LH, Magnusson RI: Interpretation of myoelectric power spectra: a model and its applications. Proc IEEE 1977, 65:653–662.

29. Štrin I, Jarm T, Stojnik V: Evaluation of the mean power frequency of the EMG signal power spectrum at endurance levels during fatiguing isometric muscle actions. Kinesiologia Slov 2008, 14:28–38.

30. Lowery M, Nolan P, O’Malley M: Electromyogram median frequency, spectral compression and muscle fibre conduction velocity during sustained sub-maximal contraction of the brachioradialis muscle. J Electromyogr Kinesiol 2002, 12:111–118.

doi:10.1186/1756-0500-7-611

Cite this article as: Trevino et al.: The effects of poliomyelitis on motor unit behavior during repetitive muscle actions: a case report. BMC Research Notes 2014 7:611.