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ELECTROMYOGRAPHIC CHARACTERIZATION IN AN ANIMAL MODEL OF  
DYSTONIA

A thesis submitted in partial fulfillment of the requirements for the degree of Masters of  
Science at Virginia Commonwealth University.

by

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# Abstract

## ELECTROMYOGRAPHIC CHARACTERIZATION IN AN ANIMAL MODEL OF DYSTONIA

By Kunal D Chaniary, M.S.

A Thesis submitted in partial fulfillment of the requirements for the degree of Masters of Science at Virginia Commonwealth University.

Virginia Commonwealth University, 2008

Director: Paul A. Wetzel, Ph.D.  
Associate Professor, Department of Biomedical Engineering

Kernicterus causes damage to the auditory system and the basal ganglia in humans. Although the Gunn rat model of kernicterus has been extensively used to characterize the auditory features, this model has not been utilized to systematically investigate the movement disorder. In the present study, spontaneously jaundiced (jj) 16 day old Gunn rat pups were treated with sulfadimethoxine to exacerbate bilirubin neurotoxicity and compared to saline treated jjs and non-jaundiced (Nj) littermates. Electromyographic (EMG) activity was recorded from antagonistic hip muscles in dystonic and in normal

appearing rats. Raw EMG signals were decomposed using the Discrete Wavelet Transform based multi-resolution analysis, and signal coefficients corresponding to the dominant EMG frequency band were chosen. Gunn rats exposed to sulfadimethoxine developed a stable clinical state characterized by prolonged abnormal axial and appendicular postures. Coherence plots revealed 4-7 Hz co-activation in antagonistic muscles that was significantly more prominent in jj sulfa treated dystonic compared to normal rats. The EMG findings support the presence of dystonia in sulfadimethoxine exposed jj Gunn rats.

## Introduction

Dystonia is characterized by sustained, abnormal stereotyped posturing.<sup>1,2,3</sup> The movement disorder is characterized physiologically by co-contractions of antagonistic muscles and overflow contraction of nearby muscles.<sup>2,4</sup> Dystonia can be classified based on the age of onset, anatomical distribution, and etiology.<sup>3</sup> Classification by anatomical distribution includes focal dystonia in which only a single body part is affected, segmental dystonia where two or more body parts are involved and generalized dystonia characterized by involvement of muscles throughout the body.<sup>5</sup> Another way to classify dystonia is based on etiology, including primary dystonia *describing* inherited or sporadic cases and secondary dystonia, which, for example, can result from strokes or the effects of drugs.<sup>5</sup>

Kernicterus or bilirubin encephalopathy, brain damage due to excessive neonatal jaundice (hyperbilirubinemia) involves damage to the brainstem auditory nuclei and the basal ganglia.<sup>6-8</sup> Kernicterus is caused by high amounts of bilirubin in the infant's blood. Bilirubin a breakdown product of heme catabolism is lipid soluble, water insoluble and neurotoxic. Under normal circumstances, unconjugated bilirubin in the blood is bound to albumin and any extra bilirubin is conjugated in the liver by uridine diphosphate glucuronosyl transferase (UDPGT) and excreted in bile. However, in infants with kernicterus, due to immaturity of glucuronosyl transferase the bilirubin binding capacity of

serum albumin is exceeded and the levels of unconjugated-indirect bilirubin in the blood increases. This extra bilirubin crosses the blood-brain barrier and penetrates the brain causing permanent brain damage. Depending on the level of exposure, the effects range from unnoticeable to severe brain damage. The basal ganglia and the auditory system are most susceptible to damage from the bilirubin toxicity. Damage to the auditory system can lead to auditory neuropathy and under prolonged exposures can cause permanent hearing loss. Damage to the basal ganglia results in dystonia, athetosis and spasticity. The syndrome in term infants can be identified by high pitched cry, poor sucking or nursing, weakness, abnormalities of muscle tone including increased muscle tone (hypertonia) and decreased muscle tone (hypotonia), lethargy, arching of the back and spine, fever and abnormal auditory brainstem potentials.<sup>9</sup>

Permanent neurological damage in kernicterus can be prevented if detection and diagnosis is done at an early stage. Currently the clinical diagnosis in kernicterus can be confirmed using magnetic resonance imaging (MRI) and Brainstem Auditory Evoked Potentials (BAEPs). BAEP's measured in hyperbilirubinemic infants show an increase in interwave interval indicating increased conduction time and decrease in amplitude indicates auditory dys-synchrony.<sup>6-10</sup> One of the most common method used for hyperbilirubinemia treatment is phototherapy.<sup>9,11</sup> In this method light causes photoisomerization of bilirubin into non-toxic water soluble products thereby preventing toxic accumulation of bilirubin in the body to prevent permanent brain damage. Another method used to treat severe hyperbilirubinemia is exchange transfusion.<sup>9,11</sup> In this therapy

the baby's blood is replaced with donor's blood that does not contain bilirubin. The goal of this procedure is to remove enough bilirubin to prevent severe complication of brain damage due to kernicterus.

The jaundiced (jj) Gunn rat model is a well established model of bilirubin encephalopathy and kernicterus.<sup>12,13</sup> This model provides an opportunity to study kernicterus and bilirubin toxicity in the laboratory and helps to understand the mechanisms, and pathophysiology that underlie this disorder. The homozygous recessive (jj) Gunn rat lacks the enzyme uridine diphosphate glucuronosyl transferase, which is responsible for the conversion of unconjugated bilirubin to conjugated bilirubin.<sup>12,13</sup> In jj Gunn rats, the serum levels of total and unconjugated bilirubin peak at about postnatal day 16 and remain moderately elevated throughout the animal's life. Their usually mild bilirubin encephalopathy can be accentuated by injecting the jj rats with sulfadimethoxine (sulfa) on day 16 to displace bilirubin from blood albumin sites into tissues, including the brain.<sup>6,12-14</sup> Within hours after sulfa injection, the rats develop auditory system dysfunction and abnormal prolonged axial and appendicular posturing, which closely resembles the clinical syndrome seen in human kernicterus. Heterozygous non-jaundiced Nj rats have about 50% of the normal enzyme activity and are phenotypically normal.<sup>14</sup> The jj Gunn rat has many of the same clinical symptoms and histopathologic lesions that are exhibited by newborn suffering from kernicterus.

Previous studies on the Gunn rat model have focused particularly on the damage in the auditory brainstem pathways.<sup>12-19</sup> In contrast, the clinical aspects and the pathophysiological correlate of the movement disorder have never been systematically investigated in this animal model. The purpose of this study is to use electromyographic (EMG) recordings and coherence analysis to support our clinical impression that the movement disorder in jaundiced Gunn rats represents dystonia. These methods could also importantly be utilized to support the validity of other phenotypic animal models of dystonia.

## MATERIALS AND METHODS

All animals and procedures used for the study were approved by the Institutional Animal Care and Use Committee (IACUC) at Virginia Commonwealth University. A total of 32 animals were obtained from our in house breeding colony for this study. Animals from six different litters were used to account for inter-litter variability. The pups were weighed on day 16 and blood samples were drawn via a cheek puncture to determine hematocrit (Hct) and total plasma bilirubin (TB) levels.

### **Groups and Clinical Assessments:**

A total of 18 jj animals received an intraperitoneal injection of sulfa (100 mg/kg) on day 16. Five of these animals developed mild-to-moderate motor disability and comprised the experimental group. Of the remaining animals, six were severely affected and did not survive the acute bilirubin encephalopathy, and seven were not affected and were not studied further. Control groups consisted of jjs given saline (n = 5) and Njs given sulfa (n = 5) or no treatment (n = 4). Littermate jjs and Njs were randomly assigned to either experimental sulfa or control groups. All animals were carefully monitored each day for loss of body weight, which was compensated with an oral feeding of kitten milk formula or a subcutaneous injection of 5% dextrose in 0.45% NaCl or a combination of both. Animals were clinically examined daily, and a clinical score (CS) between 0 to 5 was assigned based on the severity of the movement disorder (0- normal, 1- slight limb

dystonia & gait abnormality, 2-mild limb dystonia & gait abnormality, and impaired righting reflex, 3- moderate limb dystonia and gait abnormality, with prolonged righting reflex, 4- severe failure of ambulation, general lack of spontaneous movement with occasional bursts of hyperactivity, and no righting reflex, and 5- moribund, including seizures and agonal respiration),<sup>19</sup> and 0.5 was added to the score of animals appearing to be midway between a category and the next higher one. The five dystonic jj sulfa animals used in this study received scores between 1.0 and 2.5. Six dystonic jj sulfa animals given scores  $\geq 3.5$ , two each at 3.5, 4.0 and 4.5, became severely dystonic and did not survive to 34 days of age; seven additional jj sulfa animals, all receiving scores of 0.5, were clinically unaffected and not used for the study. Thus, for this study, the scale functionally ranges from 0 to 4.

### **Recording of EMG Activity:**

EMG activity was recorded on day 34, more than two weeks after sulfa injection. We targeted day 34 as an optimal age for recording EMG activity to allow sufficient time for the animal to recover from the acute bilirubin encephalopathy and for their neurological symptoms to stabilize, and for the animal to grow larger, allowing easier targeting of muscles for percutaneous insertion of fine wire electrodes.

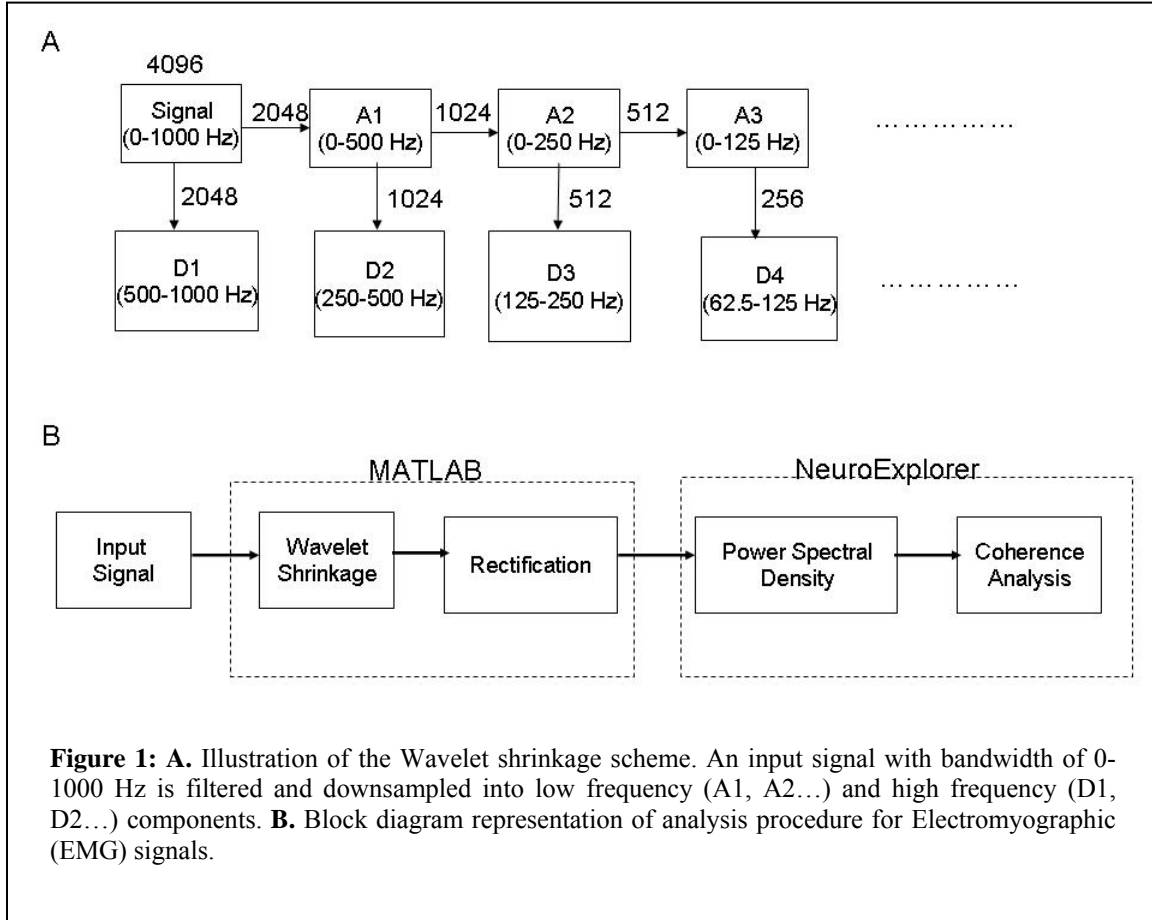
On the day of recording, the lower backs of the rats were shaved to allow targeting of the muscles for electrode insertion. Rats were lightly anesthetized with an intramuscular injection of ketamine (30 mg/kg) and acepromazine (3 mg/kg). Throughout the recording, the body temperature was monitored with a rectal temperature probe and



held at  $37^0 \pm 0.1^0\text{C}$  with a heating pad. Teflon coated 50  $\mu\text{m}$  stainless steel fine wire electrodes (A-M systems, Carlsborg, WA) were inserted percutaneously into antagonistic hip muscles, the gluteus superficialis (hip flexion) and the gluteus medius (hip extension) via a 30 gauge needle. Muscles were stimulated electrically (Grass Technologies, West Warwick, RI) and the hip motion was observed in order to verify correct placement of the electrodes. Recordings were assessed as the animals awakened from anesthesia. In the partially sedated state, a small pinch was applied to the toes and EMG activity was recorded in response to this stimulus. Multiple such stimuli were applied and EMG activity was recorded for a minimum of at least 120 sec. The signals were amplified (x 1000) and filtered (10 Hz - 1 KHz) through a differential AC amplifier (A-M Systems, Carlsborg, WA) and digitized through a NI-DAQ card (National Instruments Co., Austin, TX) at a sampling frequency = 4 KHz. EMG activity was monitored continuously on a desktop computer using Sort Client (Plexon Inc., Dallas, TX).

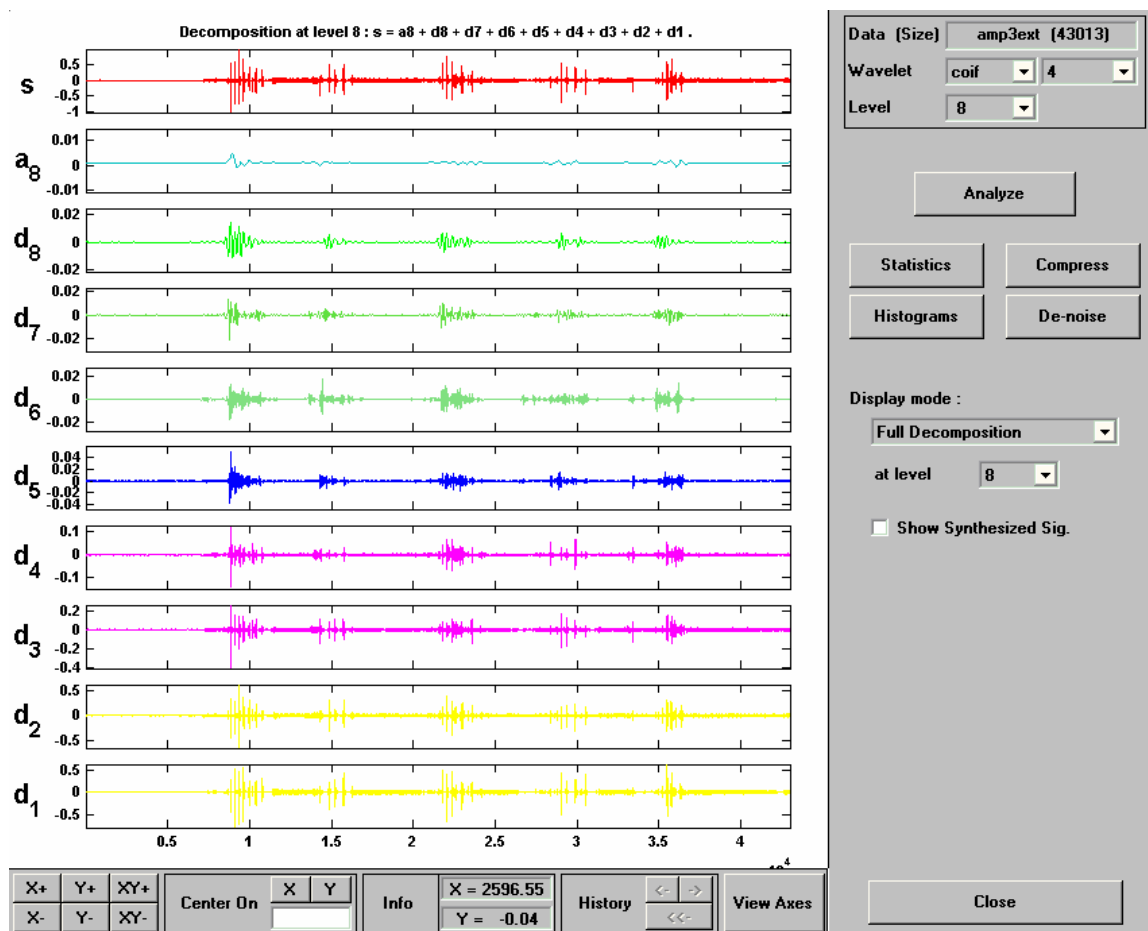
### **Decomposition of the EMG Signals into Relevant Frequency Bands:**

The raw EMG signal in dystonia consists of repetitive bursts superimposed on sustained hypertonic activity.<sup>20,21</sup> For accurate estimation of coherence between antagonist muscle pairs, it is necessary to separate the high frequency bursts from sustained hypertonic muscle activity.<sup>20,21</sup> An effective way to achieve this separation is to decompose the EMG signal using the Discrete Wavelet Transform (DWT) based multi-resolution approach.<sup>20-24</sup> The DWT approach involves passing the EMG signals progressively through a series of filters with different cutoff frequencies; thereby, decomposing the original signals into different frequency bands.<sup>23,24</sup>



At each level of decomposition, the raw EMG signal is decomposed into approximate (A1, A2...) and detail (D1, D2...) coefficients (Fig. 1A). The approximate coefficients are the high scale, low frequency components of the signal obtained by passing the original signal through a highpass scaling filter and downsampling by two. The detail coefficients are the low scale, high frequency coefficients of the signal obtained by passing the original signal through a lowpass scaling filter and downsampling by two. The high pass and low pass filters at each stage are related to each other and are called as quadrature mirror filters. The advantage of using a DWT analysis is that the time localization of frequencies is not lost during the decomposition of the

signal.<sup>24</sup> Because bursts in EMG signals are dominant in the 50-150 Hz frequency range,<sup>20,21</sup> we subjected the raw signals to level eight decomposition, which permitted us to investigate the pattern of EMG-EMG coherence in 125-250 Hz (D3), 62.5-125 Hz (D4), and 31.25-62.5 Hz (D5) frequency bands. The EMG signals were decomposed and then rectified using Matlab (MathWorks Inc., Natick, MA). Power spectral and coherence analyses were performed using Neuroexplorer (Nex Technologies, Littleton, MA) (Fig 1B). A representative decomposition of an EMG signal using Discrete Wavelet Transform is shown in Fig. 2.



**Figure 2:** Representative decomposition of an EMG signal by Discrete Wavelet Transform (DWT) using MATLAB. d1-d8 are the detail coefficients and a8 is approximate coefficient.

### **Power Spectral Density and Coherence Analysis:**

Power spectral density is the measure of the average power in a signal per unit of frequency as a function of frequency.<sup>25</sup> Coherence provides a measure of how similar two signals are and the linear association between them.<sup>2,26</sup> Based upon the degree of association, coherence can be assigned values from 0 to 1 in which 0 corresponds to minimal or no coherence and 1 indicates maximal coherence. The coherence estimation of two signals a, b can be represented as

$$\lambda_{ab}(f) = \frac{C_{ab}(f)}{\sqrt{C_{aa}(f) * C_{bb}(f)}}$$

$C_{ab}(f)$ = cross spectral density of the two signals, and

$C_{aa}(f)$ ,  $C_{bb}(f)$ = auto-spectra of each signal.

The data were smoothed by using a moving average filter (width = 1 sec) and coherence measures were obtained across multiple windows of one second duration along the entire length of each signal.

### **Statistics:**

Statistical analyses were carried out by the one way analysis of variance (ANOVA) for multiple comparisons followed by Tukey-Kramer honestly significant difference (HSD) test. A probability value of < 0.05 was considered to be statistically significant for making comparisons between groups.

## RESULTS

Table 1 provides a summary of physiological parameters and clinical scores (CS) for the animals undergoing EMG analysis. Gender was evenly distributed between the jj and Nj groups. Sulfa treated jj animals weighed significantly less than the animals in the other three groups at 34 days of age ( $p < 0.001$ ). Fig. 3A depicts a control animal and Fig. 3B represents the movement disorder seen in sulfa treated jj rats, characterized by sustained abnormal postures in the trunk and limbs. In sulfa treated jj animals, gait and balance were also affected in proportion to the degree of abnormal posturing.

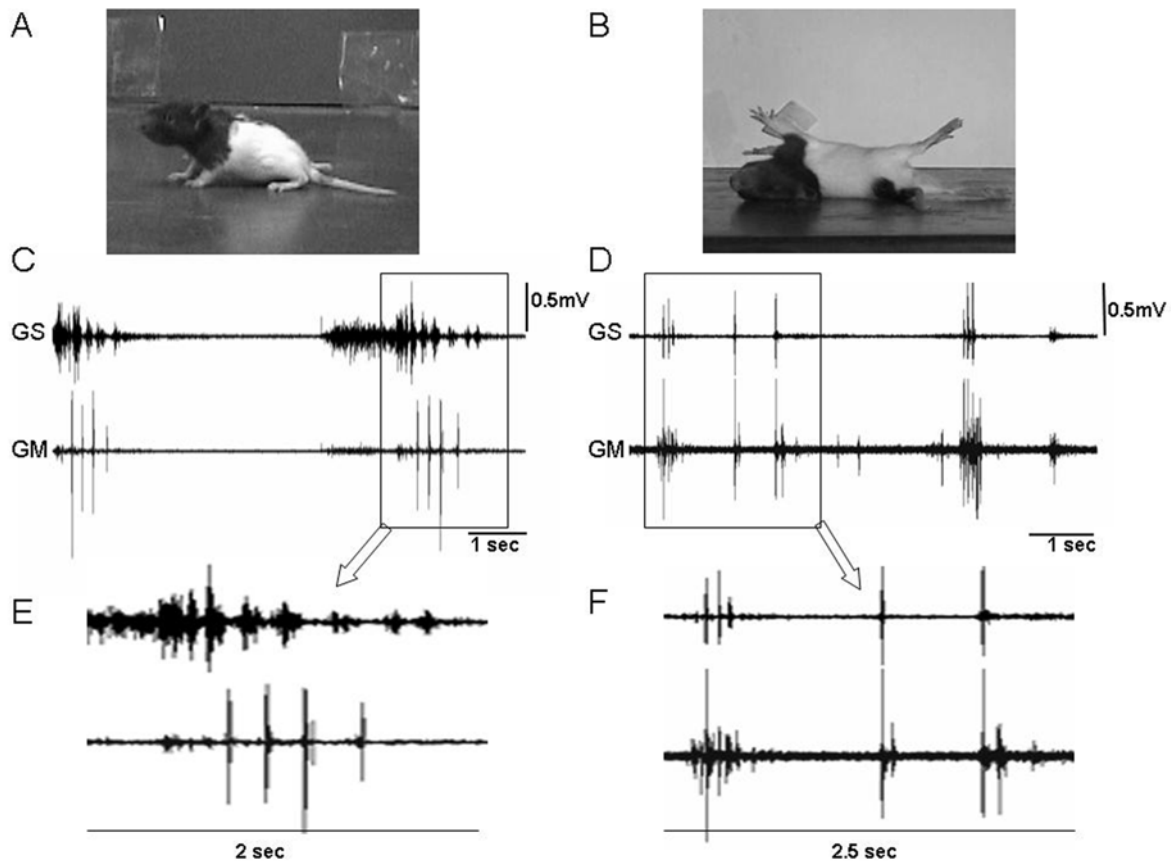
**Table 1:** Summary of physiological parameters and clinical scores of animals used for Electromyographic (EMG) characterization.

Group	Day 16			Day 34	
	Weight(gms)	TB(mg/dL)	CS	Weight(gms)	CS
jj sulfa (n=5)	26.1 ± 2.7	9.8 ± 0.8	0.5 ± 0	61 ± 5.5	1.5 ± 0.5
jj saline (n=5)	27.8 ± 2.2	10.33 ± 1.4	0.5 ± 0	79.8 ± 7.6	0.5 ± 0
Nj sulfa (n=5)	27.6 ± 4.4	0 ± 0	0 ± 0	87.5 ± 5.0	0 ± 0
Nj untreated (n=4)	-	-	-	107.5 ± 10.7	0 ± 0

(TB- Total Bilirubin, CS- Clinical Scores)

Under light anesthesia, a tail pinch produced clear volitional movements typically lasting 10-15 seconds. In sulfa-treated animals, these movements closely resembled the abnormal spontaneous motor activity observed in unanesthetized dystonic animals and clearly differed from the movements induced in lightly anesthetized control animals. Subsequent to a tail pinch, EMG activity in all Nj and saline treated jj animals showed a pattern dominated by alternating activation of agonist and antagonist hip muscles. Representative EMG activity in a sulfa treated Nj animal is shown in Fig 1C. In contrast,

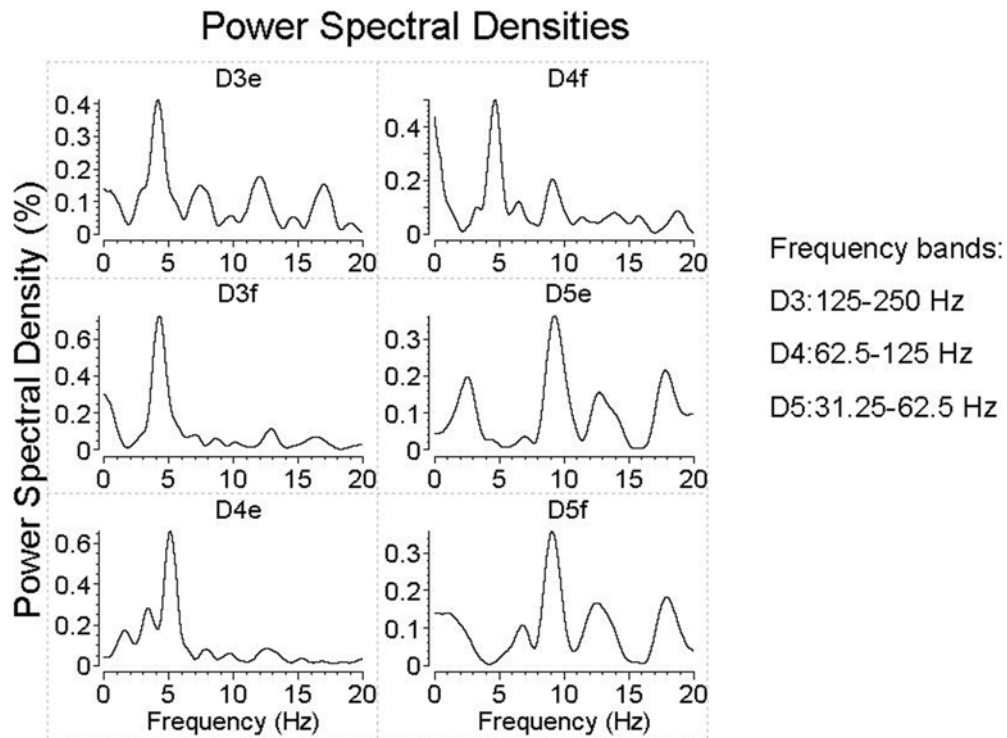
co-contractions were regularly seen in agonist and antagonistic hip muscles in sulfa treated jj animals (Fig 1D).



**Figure 3:** **A.** Normal nonjaundiced (Nj) rat 24 hours after sulfa treatment **B.** Dystonic posture in a jaundiced (jj) rat 24 hours after sulfa treatment. Note the abnormal dystonic posture, especially in the hindlimbs **C.** Raw Electromyographic (EMG) activity recorded in the Nj rat shows alternating pattern of activation in antagonistic hip muscles (gluteus superficialis [GS], gluteus medius [GM]) **D.** Raw EMG activity in jj rat shows co-contractions in antagonistic hip muscles. **E, F.** Magnified view of EMG activity in C, D, respectively.

In sulfa treated jj animals, EMG activity showed a prominent peak in the power spectral density at 4.5 Hz in D3 signals and at 5 Hz in D4 signals (Fig 4). In D5 signals, a peak in power spectral density response is seen at greater than 8 Hz. These frequency ranges however are not relevant to studies of dystonia<sup>2</sup>, and hence they have not been

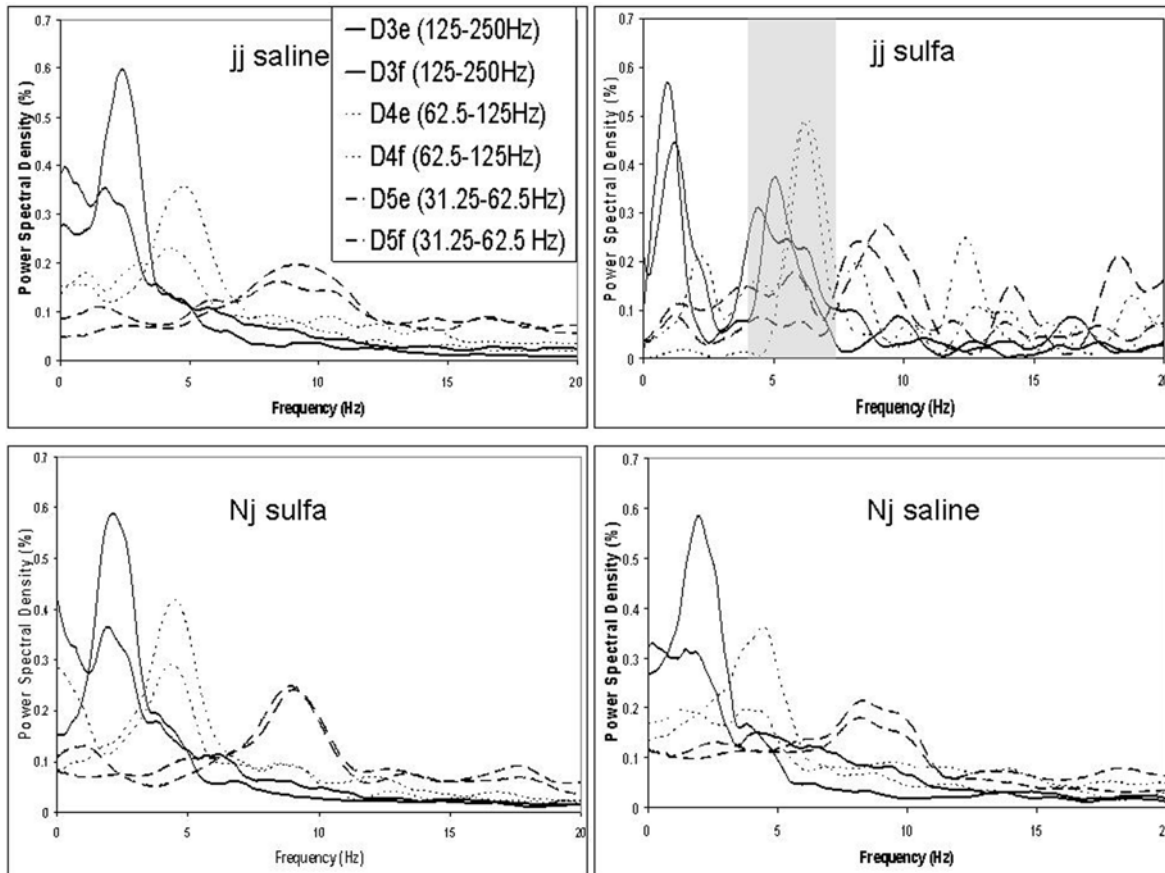
considered for further analysis. Of note, the peak response shifted towards higher frequencies with the increasing levels of decomposition (i.e. from D3 to D5).



**Figure 4:** Representative power spectral density distribution from a sulfa treated jaundiced (jj) animal. Peak in the spectral density response can be seen at 4.5 Hz in D3 (125-250 Hz) signals and at 5 Hz in D4 (62.5-125 Hz) signals for extensor (e) and flexor (f) muscles.

To compare groups, we calculated the average power spectra obtained in the four different groups of animals. The 4-7 Hz peak in power spectra was significantly prominent in D3 and D4 signals among the sulfa treated jj animals as compared to the other three groups (Fig 5). To compare the PSD's in the 4-7 Hz range in the four different groups of animals the unequal variance F test was used. The F value for signals D3e, D3f, D4e, and

D4f were 161.7, 143.2, 40.25, 9.5 respectively among the four groups of animals (all  $p$  values  $< 0.001$ ). Post hoc tests revealed that in all the 4 signals (i.e D3e, D3f, D4e, D4f) the jj sulfa group was significantly different that the other 3 groups (Tukey's HSD).

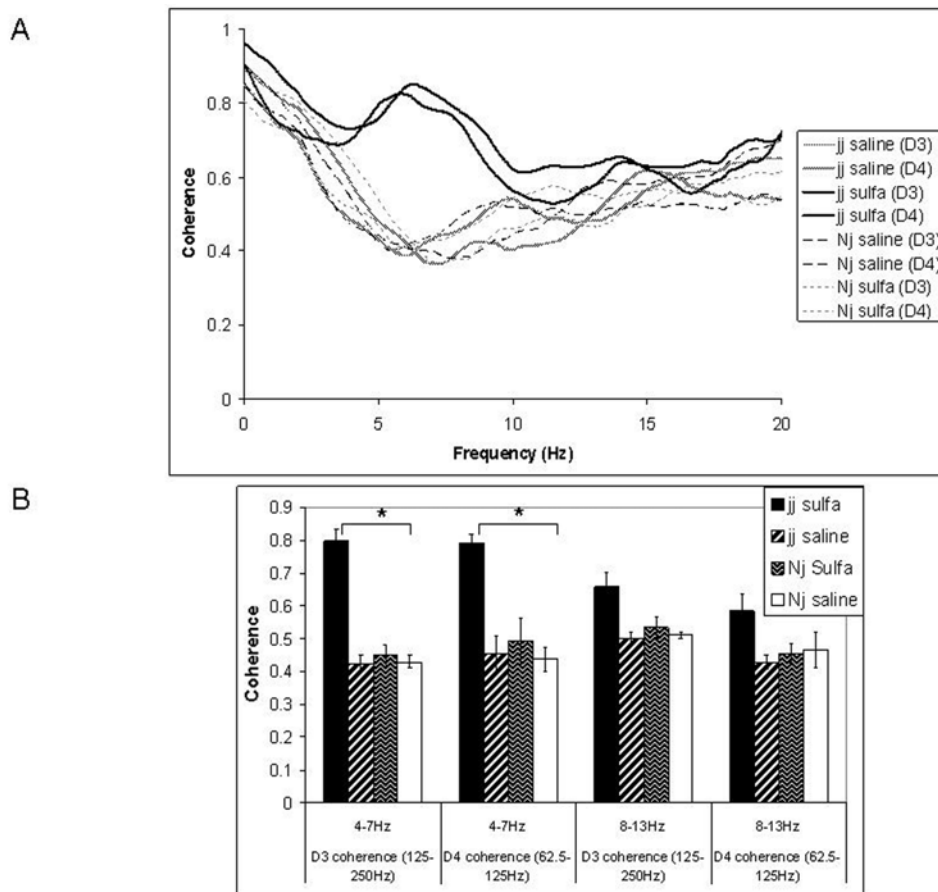


**Figure 5:** Percentage power spectral density in 4 experimental groups. There is an increase in the power spectra in 4-7 Hz range (highlighted area) in jaundiced (jj) sulfa treated animals in D3 (125-250 Hz) and D4 (62.5-125 Hz) signals for extensor (e) and flexor (f) muscles.

To look for antagonist muscle co-activation, we examined the coherence spectra in the hip muscle pairs among the four different groups of animals. Only sulfa treated jj animals showed an abnormal, prominent peak in the coherence spectra in the 4-7 Hz range (Fig 6A). The coherence for the D3 and D4 signals at 4-7 Hz were compared across the



four groups using the unequal variance F-test and were found to be different (D3,  $F[3, 324.6] = 3049.3$ ; D4,  $F[3, 325.77] = 2802.2$ ; both p-values  $< 0.0001$ ) (Fig 6B). Using Tukey's HSD, sulfa treated jj animals (D3 mean=0.796, D4=0.789) were significantly different from the other three groups. There were no significant differences in coherence between the other three groups (D3 combined mean=0.433, D4=0.461), and there was no significant correlation between the power spectral density of the D3 and D4 signals and the dystonia rating of the rat.



**Figure 6:** **A.** Coherence spectra of the D3 (125-250 Hz) and D4 (62.5-125 Hz) signals in the 4 groups of animals. A clear prominent peak in the coherence spectra for antagonistic muscles can be seen in sulfa treated jj animals in the 4-7 Hz range **B.** Summary of coherence in the 4 groups (\* $P < 0.05$ ). Error bars indicate the standard error of the mean.

## DISCUSSION

In this study, EMG recordings from antagonistic hip muscle pairs revealed prominent co-contractions in impaired, but not control rats, supporting our clinical impression that jaundiced (jj) Gunn rats with kernicterus develop dystonia. Using the DWT methodology to enhance the separation of burst from background tonic activity, an abnormal 4-7 Hz drive was observed in antagonistic hip muscles in dystonic jj rats. With this methodology, a distinct 4-7 Hz peak in the PSD was shown more prominently in 125-250 Hz and 62.5-125 Hz signals, which encompass the usual EMG burst frequency band ranges. Prior studies in humans with dystonia have demonstrated a comparable 4-7 Hz drive in antagonistic muscle pairs thereby further supporting the validity of the jaundiced Gunn rat as a model for studying human dystonia. Moreover, because similar 4-7 Hz co-contraction drives have been established in primary human dystonias, studies in the Gunn rat model may have direct implications for primary as well as secondary human dystonias.

The present findings, along with findings from human studies of dystonia, suggest that a central 4-7 Hz drive to dystonic muscle pairs underlies the pathophysiology for many common forms of dystonia. Tijssen<sup>27</sup> showed an abnormal synchronizing drive of 5-7 Hz in antagonistic cervical muscle pairs during involuntary contraction in patients with cervical dystonia. Grosse<sup>2</sup> examined the differences in EMG discharge patterns in different etiological groups of patients with dystonia, and observed an abnormal synchronizing drive of 4-7 Hz in patients with primary dystonia. Wang<sup>21</sup> studied the functional coupling

between EMG's and local field potentials (FP) of the internal globus pallidum (GPi) in a patient with dystonia. A strong coherence was reported between the activity in the GPi and the muscles of the upper arm at frequencies of 4-7 Hz in the dystonic patient.

To our knowledge, this is the first time that co-contractions have been observed in antagonistic muscles in secondary dystonia. Grosse and his colleagues<sup>2</sup> found an abnormal pattern of synchronizing descending drive of 4-7 Hz only in the distal lower limb of symptomatic patients with DYT1 gene mutation (i.e. primary dystonia) and not in patients with secondary dystonias. One possibility to explain the discrepancy may be that many secondary dystonias do not have the 4-7 Hz drive in antagonistic muscle pairs. Another possibility could be that our Gunn rat model, despite being a model of secondary dystonia, may have pathophysiological aspects that are more typical of primary dystonias.

Other animal models have previously been introduced to investigate the pathophysiological aspects of dystonia. However though, to our knowledge, the presence of dystonia has never been well characterized via EMG studies in these models. The dystonic *dt<sup>sz</sup>* (sz-epileptic seizures) hamster model shows a close resemblance to generalized paroxysmal dystonia in humans and has been used extensively to investigate neurophysiological correlates of basal ganglia in dystonia.<sup>5,28</sup> Another animal model of generalized dystonia is the *dt* rat. Expression of dystonia in these rats has been linked to the biochemical abnormalities within their cerebellum.<sup>29</sup> Byl and colleagues<sup>30</sup> have introduced a chronic model of focal hand dystonia in owl monkeys produced by repetitive

hand opening and closing. The focal feature of this model offers an advantage of producing less distress to the animals than many chronic models, though its utility is also somewhat limited by this feature. In monkeys, dystonia has been produced by administration of various toxins including 4-phenyl-1,2,3,6-tetrahydropyridine (MPTP) and 3-nitropropionic acid (3-NP). These agents however only inconsistently induce dystonia and cannot be targeted effectively towards milder symptoms.<sup>31</sup>

The present study has a number of shortcomings. Firstly, the limited range of severity of the dystonia prevented us from assessing for possible correlations between EMG power spectral density of the D3 and D4 signals and the severity of dystonia. However, the lack of correlation suggests that wavelet analysis may be relatively more sensitive to dystonia than more traditional approaches. Secondly, the significance of our findings depends largely on the premise that co-contractions support a clinical impression of dystonia. Yet, co-contractions however can occur naturally in humans, for instance, during isometric exercise<sup>32</sup> and moreover, dystonia may be exhibited with little or no coincident antagonist muscle involvement.<sup>33</sup> Despite these caveats, we contend that the present demonstration of a 4-7 Hz co-contraction drive, comparable to that established for human dystonias and limited to animals exhibiting clinical dystonia, is strong evidence in support of the presence of clinically relevant dystonia. Finally, although the animals appeared alert and displayed their naturally dystonic or normal clinical states during the EMG data collection periods, our recording conditions imposed the limitations, especially the possible effects of anesthesia on the responses. However a clear difference in findings

between our experimental and control groups argue that the effect is not due to anesthesia or other experimental conditions, which would have affected all groups equally. Future studies, are necessary to exclude the possibility that alert rats, moving naturally would, for example, exhibit a different pattern of EMG activity or would exhibit a stronger coherence pattern than lightly anesthetized, stimulated animals.

In summary, the present findings suggest that EMG may provide an objective means to verify the presence of dystonia in our animal model. This approach may provide a convenient and reliable means to objectively quantify dystonia for further experiments in the Gunn rat, as well as in other animal models of dystonia. The jaundiced Gunn rat model of kernicterus should serve as a reliable model to investigate pathophysiological mechanisms for dystonia, and to test novel therapies.

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## VITA

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Major: Biomedical Engineering

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C. Professional Positions

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Installations and servicing of imaging modalities like cathlabs, CT scan, MRI and PET scanners.

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