Cardiac Tachyarrhythmias in Hereditary Long QT Syndromes Presenting as a Seizure Disorder

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ABSTRACT: Patients with hereditary long QT disorders — Romano-Ward Syndrome and Jervell Lange-Nielsen Syndrome — sometimes present with seizures due to cardiac tachyarrhythmias. Two such patients are presented, emphasizing diagnostic clues — syncope, seizures with onset in early life, precipitation of attacks by emotional or physical stress, positive family history of sudden death and/or seizures, normal EEG and prolonged QT interval on ECG. Treatment is usually with adequate doses of sympathetic betablockers.

In 1957, Jervell and Lange-Nielsen described autosomal recessive syndrome in children as characterised by prolonged QT interval, ventricular tachyarrhythmias, and clinical manifestations of syncope, congenital deafness and sometimes sudden death. A similar congenital disorder with autosomal dominant inheritance but without deafness was recognized independently by Romano in 1963 and Ward in 1964. These hereditary prolonged QT syndromes have since been described in the literature as “Jervell-Lange-Nielsen Syndrome” (JLNS) and “Romano-Ward Syndrome” (RWS). Patients sometimes present with seizures due to cardiac tachyarrhythmias and recognition of these potentially fatal but treatable disorders by neurologists is important. We present index patients from two families — one with RWS, the other with JLNS — to illustrate clinical features and diagnostic clues with a brief discussion on management.

CASE REPORTS

Case 1: This 33-year-old woman was referred to the Neurology Department for assessment following 3 seizures between the ages of 5 and 33. These were characterized by stiffening of the body and limbs, jerking of the upper limbs, cyanosis and urinary incontinence. Each attack lasted approximately two minutes. The third episode occurred while the telephone was ringing and other attacks during sleep. Three of her six siblings died suddenly during sleep between the ages of 12 and 22 years; two of them had prior history of chronic recurrent seizures. The patient’s 12-year-old daughter suffered “blackouts” and documented ventricular tachycardia from infancy; her attacks were characterized by stiffening, urinary incontinence and pallor.

Neurological examination, blood electrolytes, EEG and CT scan of the brain were normal. ECG monitoring during EEG suggested long QT interval and subsequent 12-lead ECG confirmed a prolonged QT interval. Her 80-year-old mother was found to have a prolonged QT interval. We were also able to document prolonged QT and attacks of ventricular arrhythmias associated with syncope in the patient’s 12-year-old daughter. (Figure). The patient remains seizure-free on Propranolol. Her daughter was given the same medication, but subsequently required atrial pacing because of profound bradycardia.

Case 2: This 17-year-old congenitally deaf girl presented with recurrent blackouts since 18 months of age. These consisted of generalized stiffening, jerking, cyanosis, and at times urinary incontinence. Most attacks were precipitated by minor emotional stress. One such attack in the hospital was documented and associated with ventricular fibrillation.

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Two of her cousins, one paternal and the other maternal, were deaf; one of them had seizures and died suddenly at 21 years during an apparent seizure. The patient’s parents shared a common great grandfather.

Neurologic examination showed bilateral nerve deafness. Blood electrolytes and ECG were normal. ECG monitoring during EEG showed a prolonged rate-corrected QT of approximately 500 milliseconds (normal: 400 milliseconds). This was subsequently confirmed by a regular 12-lead ECG. She has responded well to Propranolol.

**DISCUSSION**

Recognition of RWS and JLNS among seizure patients is important to prevent potential mortality both in patients and their relatives. Diagnostic clues include: i) onset of syncope and/or cardiogenic seizures in childhood and teens,7,8 ii) family history of syncope, seizures and sudden death, iii) normal ECG with prolonged QT interval on ECG. Patients with JLNS, in addition, are congenitally deaf. The ECG should be routinely monitored during EEG in these patients with attention to the QT interval. QT can be recognized from the EEG paper in approximately ½ of tracings and measurements greater than 400 milliseconds (two divisions at usual paper speed of 30 milliseconds per second) are suspicious. However, accurate measurements and confirmation of long QT interval should be based on regular 12 lead ECG. QT interval may be prolonged in a variety of non-hereditary disorders including myocardial ischemia, cardiomyopathy, mitral valve prolapse, electrolyte and metabolic abnormalities such as hypokalemia, hypocalcemia, hypomagnesaemia; antiarrhythmic drugs like quinidine, disopyramide, procainamide; tricyclic antidepressants and phenothiazines.9,10 Age and sex also affect QT interval.

A positive family history of syncope and seizures is not unique to RWS and JLNS, but may also be present in patients with congenital and adult onset heart block.11

Seizures and syncope in RWS and JLNS are due to sudden ventricular tachyarrhythmias with resulting impairment of cerebral perfusion. These arrhythmias are considered responsible for sudden death. The mechanism of prolonged QT interval in these syndromes is not established but has been proposed to result from asymmetric stimulation of the heart by right and left sympathetic chains.12-14 Prolongation of the QT interval, reduction of the threshold for ventricular fibrillation and enhancement of ventricular excitability occur in experimental animals and humans following stimulation of left stellate ganglion or right stellate ganglonectomy.12-16 Precipitation of tachyarrhythmias and syncope in RWS and JLNS by emotional outbursts, stress and exercise is well recognized7,10 and might support the role of the sympathetic nervous system in these disorders.

Betablocker therapy is used initially in most cases to prevent tachyarrhythmias and has been shown to reduce mortality.7 Phenytion6,8 and left sympathetic ganglonectomy16 are useful in propranolol-resistant cases. In intractable cases, atrial overdrive pacing devices capable of automatic cardioversion and defibrillation should be considered.

Frequency of RWS and JLNS among seizure patients is not known, but is likely small. However, Schott et al.17 noted that cardiac arrhythmias from various causes contributed to seizures in 20% of their patients with “idiopathic” epilepsy and emphasized the importance of 24-hour ECG monitoring to detect arrhythmias.

**REFERENCES**