Propionic acidemia and long QT syndrome: A potentially serious combination

Acidemia propiónica y síndrome QT largo: una asociación potencialmente grave

Dear Editor:

Propionic acidemia (OMIM 606054) is an autosomal recessive organic acid metabolism disorder that involves a defective form of propionyl-CoA carboxylase (PCC). The disease usually manifests with acute and potentially fatal episodes of metabolic decompensation and delays in cognitive development. Its association with cardiomyopathy, usually dilated, is well known, and usually manifests during the episodes of metabolic decompensation. More recently, propionic acidemia has been associated with long QT syndrome\(^1\) (the normal range in children older than 1 year is defined by a heart-rate corrected QT interval [QTc] < 440 ms),\(^2\) which can predispose the patient to the development of ventricular arrhythmias and sudden death. Its exact aetiology is not known and its treatment is similar to that of long QT without associated propionic acidemia.

We present the case of one girl with propionic acidemia in whom the electrocardiogram (ECG) revealed prolongation of the QTc.

The patient was a girl aged 10 years diagnosed in the neonatal period as having propionic acidemia and followed up by the congenital errors of metabolism unit. The patient was undergoing treatment with l-carnitine and a protein-restricted diet. She was referred to the paediatric cardiology unit for assessment. The patient had no cardiovascular symptoms. There was no family history of sudden death or heart disease. The findings of the physical examination and echocardiogram were normal. The ECG revealed a sinus rhythm with normal voltages, but with a QTc interval of 500 ms (calculated by Bazett’s formula) and repolarization abnormalities with negative T waves in the inferior (II, III and aVF) and left precordial leads (V4–V6) (Fig. 1). Both parents were evaluated, and the ECG did not reveal any abnormal findings in either. The patient wore a Holter monitor for 24 h, which recorded a maximum QTc of 500 ms, with no evidence of arrhythmias. Laboratory tests were performed, including measurement of electrolyte, glucose, ammonia, plasma amino acids and acylcarnitine levels, the results of which were normal. Treatment of long QT syndrome was initiated with oral propanolol. Subsequent evaluations showed that the patient remained free of cardiovascular symptoms and tolerated therapy with propanolol, while prolongation of the QTc persisted in the ECG (QTc, 440–480 ms).

Propionic acidemia is caused by the deficient activity of PCC, a biotin-dependent mitochondrial enzyme needed for the conversion of propionyl-CoA to d-methylmalonyl-CoA. PCC is a heteropolymer composed of two types of subunits,
by symmetric ascending motor weakness and areflexia. GBS encompasses a group of acute immune-mediated acquired polyradiculoneuropathies that are usually preceded by gastrointestinal or respiratory infections. The most common forms are classified on the basis of neurophysiological criteria as acute inflammatory demyelinating polyradiculoneuropathy (AIDP) and acute motor axonal neuropathy (AMAN). While both forms may be clinically indistinguishable, their prognosis can be very different. We present a case in which serial neurophysiological studies were key in the identification of a case of GBS with an atypical presentation, its classification as AMAN, and the assessment of the patient’s outcome.

A male patient 7 years of age with no medical or surgical history of interest sought care in the emergency department for pain in the lumbar region and the right triceps surae accompanied by limited mobility in climbing and going down stairs lasting 11 days. We observed that the patient had difficulty walking on his toes, and the rest of the neurological examination was normal, including the deep tendon reflexes (DTRs). The findings of the lower limb X-rays were normal.

References


A. Fuertes Moure a,b,c, M. Centeno Jiménez a, R. Álvarez García-Rovés a,b, N. Gil Villanueva a, C. Medrano López a

a Instituto de Investigación Sanitaria, Hospital Universitario Gregorio Marañón, Madrid, Spain
b Sección de Cardiología Pediátrica, Hospital Universitario Gregorio Marañón, Madrid, Spain
c Corresponding author.
E-mail address: angelesfm@hotmail.es (A. Fuertes Moure).

Contribution of serial neurophysiological studies in atypical Guillain-Barré syndrome

Contribución de los estudios neurofisiológicos seriados en el síndrome de Guillain-Barré atípico

Dear Editor:

Guillain-Barré syndrome (GBS) is the most common cause of acute flaccid paralysis in children. It is characterised

by various medicines that may lengthen the QT interval and thus promote the development of ventricular arrhythmias.