Brugada Phenocopy Emerging as a New Concept. Response

Fenocopia de Brugada: surgimiento de un nuevo concepto. Respuesta

To the Editor,

The authors appreciate the interest in the case report we published in Revista Española de Cardiología concerning the observation of the Brugada electrocardiographic (ECG) pattern in a patient with hyperkalemia.1 We likewise welcome the introduction of the concept of phenocopy, an expression with which our finding is compatible.2,3 We also consider it opportune to stress that both the latest consensus on ECG diagnosis of Brugada syndrome and the introduction of the term phenocopy are more recent than the online publication of our case report in 2011.2–4 The definitions of the ECG patterns that are typical of Brugada syndrome and those that mimic this syndrome in the presence of serum electrolyte disturbances were introduced subsequent to our publication.

It is important to highlight reasons for attributing the changes observed in the ECG to hyperkalemia rather than to the acidosis and hypokalemia also observed in our patient. Reports of Brugada phenocopy associated with hypokalemia and acidosis have described the development of pseudo J waves in the QRS complex and ST segment depression in leads other than right precordial leads. These are precisely the features that differentiate this ECG pattern from type 1 Brugada pattern.5,6 Other possible causes (hyperglycemia, drugs, fever, and myocardial ischemia) were ruled out in the case discussed in our report.

We appreciate any contribution that aids in the understanding of the mechanisms involved in the induction of ECG patterns mimicking Brugada syndrome (phenocopies) and other patterns, such as early repolarization, which can also be associated with the risk of sudden cardiac death.7

Finally, we agree with Dr. Anselm on the importance of performing a challenge test with flecainide to rule out Brugada syndrome. Until the prognosis of patients presenting with Brugada phenocopy has been established, it is advisable to perform pharmacological challenge tests and, if appropriate, to induce ventricular arrhythmias by means of an electrophysiological study.

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RASophathies: From Noonan to LEOPARD Syndrome

RASophatías: del síndrome de Noonan al síndrome LEOPARD

To the Editor,

We have read with interest the article recently published by Carcavilla et al.1 titled “LEOPARD Syndrome: A Variant of Noonan Syndrome Strongly Associated With Hypertrophic Cardiomyopathy.” However, we would like to add some comments that we find interesting.

LEOPARD syndrome (LS) (OMIM 151100) and Noonan syndrome (NS) (OMIM 163950) are two disorders that are part of a newly classified family of autosomal dominant syndromes termed “RSopathies”, which are caused by germline mutations in components of the RAS-MAPK (mitogen-activated protein kinases) signal transduction pathway2 that is involved in the regulation of normal cell proliferation, survival, and differentiation.

Although the diagnosis of LS is made on clinical grounds by observation of key features, none of which is pathognomonic, to provide a valid overview on symptoms and features of LS we should not overestimate the reports published in the premolecular era, because these cohorts may consist of heterogeneous diseases. In fact, in early childhood the phenotype of LS can be typical of NS; however, with age other characteristic features of LS, including lentigines, hypertrophic cardiomyopathy (HCM), and hearing loss, may appear. For this reason, anomalies observed in molecularly confirmed cases should be regarded as more reliable.

Genetically, both syndromes share mutations in the PTPN11 (protein-tyrosine phosphatase, non-receptor type 11) gene on chromosome 12q24, as they are heterozygous missense mutations in PTPN11 observed in up to 90% and 50% of LS and NS cases, respectively. Similarly, mutations in the RAF1 (v-Raf-1 murine leukemia viral oncogene homolog 1) gene on chromosome 3p25.2 and mutation in the BRAF (v-Raf murine sarcoma viral oncogene homolog B1) gene on chromosome 7q34 are also seen. However, the point mutations identified in PTPN11 that are associated with NS are distinct from those associated with LS and therefore with different biochemical properties: gain-of-function mutations in PTPN11 are more frequent in NS patients3,4 while a loss-of-function or dominant-negative mutations in PTPN11 are more prevalent in patients with LS.5,6 These gain-of-function and loss-of-function mutations may explain the differences in phenotypes between these two syndromes.

Although it is not included in the LEOPARD acronym, HCM is the most frequent anomaly observed, representing a potentially life-threatening problem in these patients. HCM, which is generally...