Understanding Cardiac Calcium Channelopathies

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Timothy syndrome is a rare genetic disorder characterized by QT prolongation (designated LQTS), arrhythmias and sudden death, structural heart disease, cognitive defects with autism, syndactyly (webbed fingers and toes), hypoglycemia, and immune deficiencies.1,2 A single mutation (G406R) in exon 8a of the cardiac L-type calcium channel (CACNA1C, Ca,1.2, α1c) was shown to cause Timothy syndrome in multiple unrelated subjects, whereas mutations (G406R, G402S) in the alternatively spliced exon 8 (which is expressed at ~3-fold–higher levels than exon 8a) cause a similar syndrome lacking syndactyly.3,4 These 3 mutations decrease voltage-dependent inactivation of Ca,1.2, which is predicted to slow the inactivation of ICa,L during each action potential, prolong action potential and QT interval duration, increase the amplitude and duration of Ca2+ transients, and predispose to afterdepolarizations and arrhythmias.3–5

The findings by Thiel et al are potentially important for several reasons. First, the mechanistic link between changes in ICa,L and arrhythmia susceptibility in Timothy syndrome appears more complex than previously thought. Second, this is the first evidence that CaMKII directly participates in arrhythmia susceptibility in a human inherited channelopathy. Third, although Timothy syndrome is rare, abnormalities in Ca2+ handling may cause arrhythmias and sudden death in common cardiac conditions including heart failure.12–14 CaMKII-mediated changes in ICa,L facilitation and ryanodine receptor leak can be studied in these other pathological states,
and the potential efficacy of CaMKII inhibition can be assessed. Finally, the findings raise the intriguing possibility that CaMKII-related proteins may contribute to the extracardiac phenotypes including autism.

A number of additional studies are required to confirm the conclusions. As pointed out by the authors, the findings need to be replicated in myocytes from a larger animal with cardiac action potentials more similar to those of humans. Although canine wedge studies have been performed using the Ca\(^{2+}\) channel opening drug BayK8644 to model increases in \(I_{\text{C\text{a},L}}\),\(^{16}\) additional studies using intact hearts and whole animal models expressing mutant channels are necessary. In addition, long-term expression of the mutation may lead to different findings than short-term viral exposure, as seen with the effects on K\(^+\) channel expression caused by transgenic overexpression of the AC3-I peptide.\(^{11}\) Studies using G406R Ca\(_{1.2}\) knockin mice could address many of these questions.

During the last 15 years, ion channelopathies have been shown to cause a number of inherited arrhythmia syndromes, including long-QT syndrome (K\(^+\), Na\(^+\), and Ca\(^{2+}\) channels), short-QT syndrome (K\(^+\) channels), Brugada syndrome (Na\(^+\) channels), short-QT and Brugada syndrome (Ca\(^{2+}\) channels), and catecholaminergic polymorphic ventricular tachycardia (SR Ca\(^{2+}\) release channels).\(^{17}\) Given the importance of \(I_{\text{C\text{a},L}}\) to cardiac electrical and mechanical function, it is surprising how few pathogenic mutations have been identified. Two potential causes could underlie this finding: Mutations could be rare because they are highly lethal, or they could be only rarely identified because they are well tolerated. The evidence suggests that both may be true. Marked increases in \(I_{\text{C\text{a},L}}\) occur as a result of physiological \(\beta\)-adrenergic stimulation; similarly, L-type Ca\(^{2+}\) channel blockers are well tolerated but do not prevent arrhythmias or sudden cardiac death despite decreasing intracellular Ca\(^{2+}\) load. The mutations that lead to Timothy syndrome are all located in the 8th exon and all affect voltage-dependent inactivation. Similarly, only a few mutations characterized by marked loss of Ca\(^{2+}\) channel function have been identified in individuals with an overlap syndrome consisting of short-QT intervals and Brugada-like ECG abnormalities.\(^{18}\) Thus, it is possible that whereas modest changes in the amplitude of \(I_{\text{C\text{a},L}}\) are well tolerated, changes in its time course disrupt important signaling mechanisms including CaMKII and are poorly tolerated. If true, heterozygous mutations that alter channel number or current amplitude would have a limited phenotype, whereas rare mutations that alter channel properties in specific ways would lead to highly lethal arrhythmia syndromes.

In summary, the study by Thiel et al has expanded our understanding of the pathophysiological mechanisms underlying Timothy syndrome. In addition, it may ultimately provide new insights into the role of CaMKII in arrhythmia susceptibility, into Ca\(^{2+}\) handling in other organs including the brain, and into the regulatory pathways that are controlled by cardiac Ca\(^{2+}\) channels. In addition, at least some hope exists that these findings could ultimately lead to novel therapeutic options to treat or prevent life-threatening arrhythmias.

Disclosures

None.

References


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