



Timothy syndrome associated with a novel likely pathogenic CACNA1C variant: A neonatal case report

Ramón Miguel Vargas-Vera, PhD^{1,2*}

¹Universidad Católica Santiago De Guayaquil, Guayaquil, Ecuador

²Universidad De Guayaquil, Guayaquil, Ecuador

Abstract

Timothy Syndrome (TS) is a rare multisystem channelopathy caused by pathogenic variants in CACNA1C, classically associated with severe cardiac involvement and neurodevelopmental abnormalities. Variants affecting codon 402 represent a critical functional hotspot of the CaV1.2 L-type calcium channel. We report a preterm male neonate presenting with microcephaly, small anterior fontanelle, mild fetal ventriculomegaly, cardiomyopathy, bradycardia, and neonatal respiratory failure. Whole-exome sequencing identified a heterozygous likely pathogenic variant in CACNA1C (NM_000719.7:c.1205G>A; p.Gly402Asp), not previously reported in the literature but affecting a codon with established pathogenic relevance in Timothy syndrome. This case expands the genotypic and phenotypic spectrum of Timothy syndrome and highlights the importance of early clinical-genetic correlation in neonates with severe cardiac manifestations.

Keywords: Timothy syndrome, CACNA1C, Neonatal cardiomyopathy, Channelopathy, Case report

Introduction

Timothy syndrome (OMIM #601005) is an extremely rare genetic disease, inherited in an autosomal dominant manner, caused by pathogenic variants in the CACNA1C gene, which encodes the $\alpha 1C$ subunit of the voltage-dependent L-type calcium channel (CaV1.2) [1-3]. This channel plays a key role in cardiac depolarization, myocardial contractility, and neurological development.

The variants classically associated with the syndrome affect highly conserved domains of the channel, particularly codon 402, causing altered voltage-dependent inactivation and a sustained increase in intracellular calcium influx [2,4,5]. Clinically, the syndrome manifests with severe arrhythmias, cardiomyopathy, neurological abnormalities, craniofacial anomalies, and, in some cases, syndactyly.

Case presentation

Male newborn, product of a pregnancy complicated by fetal hemodynamic alterations evidenced by Doppler echocardiography, who was delivered by emergency cesarean section due to acute fetal distress. He presented with severe respiratory

distress at birth, requiring immediate admission to the neonatal intensive care unit.

On initial physical examination, microcephaly was observed (head circumference below the percentile for gestational age), a small anterior fontanelle, without evident major facial dysmorphisms. The cardiovascular system showed signs of hemodynamic compromise and bradycardia. Imaging studies confirmed cardiomyopathy and mild fetal ventriculomegaly. He developed episodes of neonatal hypoglycemia and suspected neonatal sepsis.

Genetic findings

Whole exome sequencing was performed using genomic DNA obtained from peripheral blood, a methodology widely validated for the diagnosis of rare genetic diseases in the neonatal period [6,7]. The analysis identified a heterozygous variant likely pathogenic in CACNA1C: NM_000719.7:c.1205G>A, resulting in the amino acid change p.Gly402Asp.

This variant has an extremely low frequency in population databases (gnomAD <0.001%) and in silico predictions highly suggest a damaging effect (REVEL 0.95; 3Cnet 0.97). Although it has not been previously reported in patients, other variants at the

same codon (p.Gly402Ser and p.Gly402Val) have been described as pathogenic in association with Timothy syndrome, reinforcing the functional relevance of this protein region [4,8].



Figure 1. Neonatal facial appearance showing subtle craniofacial features without major dysmorphisms

Figure 1. Neonatal clinical appearance. Frontal facial view showing subtle craniofacial features without major dysmorphisms. Note the reduced cranial size consistent with microcephaly and a small anterior fontanelle. Images were obtained during the neonatal period in the neonatal intensive care unit, prior to the genetic diagnosis.



Figure 2. Neonate in the neonatal intensive care unit under cardiorespiratory monitoring

Figure 2. General clinical status in the neonatal intensive care unit. The patient is shown in a neonatal incubator under cardiorespiratory monitoring, illustrating the need for intensive neonatal care due to respiratory distress and cardiovascular instability.



Figure 3. Lower limb and plantar view showing absence of syndactyly, consistent with atypical Timothy syndrome

Figure 3. Lower limb examination. Image shows the plantar aspect of the foot and lower limb with no evidence of classical syndactyly. The absence of limb fusion supports an incomplete or atypical presentation of Timothy syndrome.

Discussion

The identification of a probably pathogenic variant at codon 402 of CACNA1C in a patient with severe neonatal cardiac involvement and early neurological abnormalities is highly suggestive of Timothy syndrome, even in the absence of classic syndactyly [3,9,10]. The absence of syndactyly in this patient supports the existence of incomplete or atypical forms of the syndrome, as previously described in the literature. This case contributes to expanding the mutational spectrum of CACNA1C and reinforces the functional relevance of codon 402 as a critical region for CaV1.2 channel physiology. It also highlights the importance of early genetic diagnosis for clinical

management, prognostic stratification, and family genetic counseling.

Conclusion

A neonatal case compatible with Timothy syndrome associated with a new likely pathogenic variant in CACNA1C is described. The clinical-genetic correlation supports the causal involvement of the p.Gly402Asp variant and emphasizes the need for close cardiac and neurological monitoring in these patients.

Ethics statement

Written informed consent was obtained from the patient's parents for genetic testing and for publication of this case report and any potentially identifiable data/images.

Author contributions

RMVV conceptualized the study, interpreted the genetic findings, and drafted the initial manuscript. MVP-I contributed to clinical data collection, literature review, and critical revision of the manuscript. KSV-S contributed to the cardiology assessment, clinical interpretation, and manuscript revision. IMT-R and MJUB contributed to patient data acquisition, clinical follow-up, and manuscript review. All authors approved the final version of the manuscript and agree to be accountable for all aspects of the work.

Data availability statement

The data supporting the conclusions of this article are included within the article. Further inquiries can be directed to the corresponding author.

Informed consent was obtained from the parents for the performance of genetic studies and the publication of clinical information in an anonymized manner.

Acknowledgments

To the patient's family and the neonatology and clinical genetics team for their collaboration in the comprehensive management of the case.

Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Declarations of interest: None. Funding: This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

References

1. Splawski I, Timothy KW, Sharpe LM, et al. CaV1.2 calcium channel dysfunction causes a multisystem disorder including arrhythmia and autism. *Cell.* 2004;119(1):19–31. <https://doi.org/10.1016/j.cell.2004.09.011>
2. Splawski I, Timothy KW, Decher N, et al. Severe arrhythmia disorder caused by cardiac L-type calcium channel mutations. *Proc Natl Acad Sci USA.* 2005;102(23):8089–8096. <https://doi.org/10.1073/pnas.0502506102>
3. Splawski, Igor et al. "Severe arrhythmia disorder caused by cardiac L-type calcium channel mutations." *Proceedings of the National Academy of Sciences of the United States of America* vol. 102,23 (2005): 8089-96; discussion 8086-8. doi:10.1073/pnas.0502506102
4. Boczek, Nicole J et al. "Novel Timothy syndrome mutation leading to increase in CACNA1C window current." *Heart rhythm* vol. 12,1 (2015): 211-9. doi: 10.1016/j.hrthm.2014.09.051
5. Wemhöner, Konstantin et al. "Gain-of-function mutations in the calcium channel CACNA1C (Cav1.2) cause non-syndromic long-QT but not Timothy syndrome." *Journal of molecular and cellular cardiology* vol. 80 (2015): 186-95. doi: 10.1016/j.jmcc.2015.01.002
6. Clark, Michelle M et al. "Meta-analysis of the diagnostic and clinical utility of genome and exome sequencing and chromosomal microarray in children with suspected genetic diseases." *NPJ genomic medicine* vol. 3 16. 9 Jul. 2018, doi:10.1038/s41525-018-0053-8
7. Kingsmore, Stephen F et al. "A Randomized,

Controlled Trial of the Analytic and Diagnostic Performance of Singleton and Trio, Rapid Genome and Exome Sequencing in Ill Infants." *American journal of human genetics* vol. 105,4 (2019): 719-733. doi: 10.1016/j.ajhg.2019.08.009

8. Delinière, Antoine et al. "Phenotypic Characterization of Timothy Syndrome Caused by the *CACNA1C* p. Gly402Ser Variant." *Circulation. Genomic and precision medicine* vol. 16,3 (2023): 280-282. doi:10.1161/CIRCGEN.122.004010

9. Underwood, Jack F G et al. "Timothy Syndrome and *CACNA1C*-Related Disorder: First International Language and Management Guidelines Consensus Statement." *Research square* rs.3.rs-8058536. 17 Nov. 2025, doi:10.21203/rs.3.rs-8058536/v1. Preprint.

10. Dufendach, K, Timothy, K, Ackerman, M. et al. Clinical Outcomes and Modes of Death in Timothy Syndrome: A Multicenter International Study of a Rare Disorder. *J Am Coll Cardiol EP*. 2018 Apr, 4 (4) 459-466. <https://doi.org/10.1016/j.jacep.2017.08.007>

11. Brugada, Josep et al. "Present Status of Brugada Syndrome: JACC State-of-the-Art Review." *Journal of the American College of Cardiology* vol. 72,9 (2018): 1046-1059. doi: 10.1016/j.jacc.2018.06.037

12. Wilde, Arthur A M, and Ahmad S Amin. "Clinical Spectrum of SCN5A Mutations: Long QT Syndrome, Brugada Syndrome, and Cardiomyopathy." *JACC. Clinical electrophysiology* vol. 4,5 (2018): 569-579. doi: 10.1016/j.jacep.2018.03.006