CaMKII inhibition rectifies arrhythmic phenotype in a patient-specific model of catecholaminergic polymorphic ventricular tachycardia

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Induced pluripotent stem cells (iPSC) offer a unique opportunity for developmental studies, disease modeling and regenerative medicine approaches in humans. The aim of our study was to create an in vitro ‘patient-specific cell-based system’ that could facilitate the screening of new therapeutic molecules for the treatment of catecholaminergic polymorphic ventricular tachycardia (CPVT), an inherited form of fatal arrhythmia. Here, we report the development of a cardiac model of CPVT through the generation of iPSC from a CPVT patient carrying a heterozygous mutation in the cardiac ryanodine receptor gene (RyR2) and their subsequent differentiation into cardiomyocytes (CMs). Whole-cell patch-clamp and intracellular electrical recordings of spontaneously beating cells revealed the presence of delayed afterdepolarizations (DADs) in CPVT-CMs, both in resting conditions and after β-adrenergic stimulation, resembling the cardiac phenotype of the patients. Furthermore, treatment with KN-93 (2-[N-(2-hydroxyethyl)]-N-(4-methoxybenzenesulfonyl)]amino-N-(4-chlorocinnamyl)-N-methylbenzylamine), an antiarrhythmic drug that inhibits Ca\(^{2+}\)/calmodulin-dependent serine–threonine protein kinase II (CaMKII), drastically reduced the presence of DADs in CPVT-CMs, rescuing the arrhythmic phenotype induced by catecholaminergic stress. In addition, intracellular calcium transient measurements on 3D beating clusters by fast resolution optical mapping showed that CPVT clusters developed multiple calcium transients, whereas in the wild-type clusters, only single initiations were detected. Such instability is aggravated in the presence of isoproterenol and is attenuated by KN-93. As seen in our RyR2 knock-in CPVT mice, the antiarrhythmic effect of KN-93 is confirmed in these human iPSC-derived cardiac cells, supporting the role of this in vitro system for drug screening and optimization of clinical treatment strategies.

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Induced pluripotent stem cell (iPSC) technology has been proposed as a valuable approach for studying the pathophysiology of human diseases in vitro. iPSCs are generated by the reprogramming of somatic cells through the expression of ectopic transcription factors, and have been shown to be able to differentiate into all cell types of the body, including functional cardiomyocytes (CMs).1–3

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**Abbreviations:** AP, action potential; APD, action potential duration; APD90, action potential duration at 30% of repolarization; APD50, action potential duration at 50% of repolarization; APD90, action potential duration at 90% of repolarization; CaMKII, Ca\(^{2+}\)/calmodulin-dependent serine–threonine protein kinase II; CASQ2, calsequestrin 2; CD-15 or SSEA1, stage-specific embryonic antigen 1; CM, cardiomyocyte; CPVT, catecholaminergic polymorphic ventricular tachycardia; DADs, delayed afterdepolarizations; DAPI, 4',6-diamidino-2-phenylindole; dCa\(^{2+}\)/dtna, rate of intracellular calcium increase; EBs, embryoid bodies; ECG, electrocardiogram; ES, embryonic stem cells; FACS, fluorescence-activated cell sorting; FBS, fetal bovine serum; FH, fetal heart; Fluo-4, 2′,7′-bis(carboxyethyl)aminomethyl fluorescein; GFA, GATA-binding factor 6; HEK-293T, human embryonic kidney 293 T cells; HGFRT, hypoxanthine–guanine phosphoribosyltransferase; iPSC, induced pluripotent stem cells; iso, iso-proterenol; KN-92, 2-N-(4-chlorocinnamyl)-N-methylbenzylamine, phosphate; KN-93, 2-N-(2-hydroxyethyl)Y-N-(4-methoxybenzenesulfonyl)]amino-N-(4-chlorocinnamyl)-N-methylbenzylamine, KRT14, epidermal keratin 14; MDP, maximum diastolic potential; MEF, mouse embryonic fibroblasts; MHC, myosin heavy chain, subunit α; MHCβ, myosin heavy chain, subunit β; MSE, mean standard error; NCAM1, neural cell adhesion molecule 1; OCT4, octamer-binding transcription factor 4; PCR, polymerase chain reaction; PECAM1, platelet endothelial cell adhesion molecule 1; PFA, paraformaldehyde; RyR2, ryanodine receptor, type 2; SOX17, SRY-box containing gene 17; SOX2, SRY (sex determining region Y)-box 2; SSEA4, stage-specific embryonic antigen 4; TA, triggered activity; TRA1–60, tumor rejection antigen 1–60; WT, wild type

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The usefulness of reprogrammed cells in modelling diseases in vitro has recently been tested for several human congenital cardiovascular diseases: iPSCs have been used for the investigation of monogenic disorders (i.e. long QT (prolongation of Q–T interval combined with torsades de pointes and manifests as several different forms) syndromes, Brugada syndrome/cardiac conduction disease, catecholaminergic polymorphic ventricular tachycardia and other forms of congenital arrhythmias) and disorders in which cardiac defects are part of a multigene phenotype (i.e. Leopard and Timothy syndromes). These reports confirmed that patientspecific iPSCs are able to give rise to differentiated CMs that possess the main functional and morphological aberration typical of the disease in vivo. Furthermore, the importance of iPSC in drug discovery applications is useful to identify novel therapeutic molecules and devise patientspecific therapies.

In our study, we focused on catecholaminergic polymorphic ventricular tachycardia (CPVT), a highly lethal inherited arrhythmogenic disease that may cause syncope and sudden death early in life owing to adrenergically mediated bidirectional ventricular tachycardia. Two genetic forms of the disease have been described: one transmitted as an autosomal dominant trait, caused by mutations in the gene encoding the ryanodine receptor type 2 (RyR2), and another very rare form due to recessive mutations in the cardiac-specific isoform of the casequestrin gene, CASQ2 (casequestrin 2). Both genes are involved in regulating Ca2+ handling within the CM and thus are key in determining excitation-contraction coupling.

Although the mortality rate associated with the disease is extremely high (30–35% by the age of 35 years), therapies are limited. Treatment with β-adrenergic blockers are usually effective in preventing recurrences of arrhythmias in the majority of patients, but approximately 30% of patients still experience at least one episode of life-threatening adrenergically induced arrhythmia (leading to syncope or cardiac arrest) despite therapy and will need implantable cardioverter defibrillators.

Therefore, development of model systems facilitating screening of new therapeutic molecules for the treatment of CPVT is highly advisable. Among the putative players in determining the CPVT phenotype, Ca2+/calmodulin-dependent serine–threonine protein kinase II (CaMKII) has been recently implicated in arrhythmic events elicited by β-adrenergic activation, and we recently demonstrated that its inhibition is able to prevent ventricular arrhythmogenesis in a mouse model of CPVT.

With these considerations in mind, our intent was to create a patient-specific cell-based system that could be used as an in vitro model to facilitate the screening of new therapeutic molecules for the treatment of CPVT. For this purpose, we generated an iPSC-based cardiac model from a patient carrying a heterozygous mutation in the gene encoding RyR2 and with phenotypic manifestations of the disease. In a first instance, we verified that the disease phenotype was recapitulated in the CMs derived from these iPSC. Subsequently, we inhibited the Ca2+-CaMKII pathway, which affects calcium handling, to test whether we could rescue the disease phenotype in human cardiac cells to confirm the clinical relevance of the observation made in myocytes derived from knock-in mice carriers of a heterozygous defect in RyR2 and presenting the clinical phenotype of CPVT.

Our results support the view that iPSC technology is likely to have clinical applicability to predict response to therapy in individual patients.

**Results**

**Clinical history.** In June 2006, the team of our outpatient clinic for inherited arrhythmia at the Maugeri Foundation was contacted for the assessment of a family with a history of juvenile sudden cardiac death. The proband (Figure 1A, subject II-2), a 42-year-old female reported that two of her children died suddenly before age 10 years (Figure 1A, subjects III-1 and III-2) both in a condition of adrenergic stress. Ill-1 died at the age of 8 years while riding on a carousel and III-2 died suddenly at the age of 9 years running in a school competition. The mother also reported that III-1 experienced a syncopal spell during physical activity a few months before dying. At that time, the boy was taken to the emergency room, but resting electrocardiogram (ECG) and echocardiogram were unremarkable and he was discharged. The other child of the proband, that is, III-2, died at the age of 9 years with no previous symptoms. At first clinical evaluation, the mother (II-2) reported two previous episodes of loss of consciousness during physical activity (at the age of 41 and 42 years) and reported that in a previous exercise stress test there was documentation of isolated premature ventricular contractions and a ventricular coupllet that resulted in the interruption of the test. We recorded her resting ECG (Figure 1B) and echocardiogram, which were unremarkable. However, maximal exercise stress test documented the onset of sustained bidirectional ventricular tachycardia (Figure 1B). CPVT diagnosis was established and β-blocker therapy was administered. A second exercise stress test after 5 days of therapy with nadolol (2 mg/kg) showed suppression of arrhythmias after maximally tolerated effort. The patient has remained asymptomatic, with no evidence of arrhythmias as of September 2012. Sequencing of the entire open reading frame of the RyR2 gene identified the c.6933 G>C nucleotide transversion in exon 46, leading to the p.Glu2311Asp missense mutation. Unfortunately, no post-mortem samples were available for the deceased children. Genetic testing also identified the same mutation in the asymptomatic two-year-old daughter (III-3), who was promptly treated with oral nadolol (2 mg/kg). Holter monitoring off therapy showed rare supraventricular and ventricular ectopic beats that disappeared after therapy.

**Generation of patient-specific CPVT-iPSC and their characterization.** CPVT-iPSCs were generated from primary fibroblasts isolated from a skin biopsy of the proband through lentiviral transduction with OCT4 (octamer-binding transcription factor 4), SOX2 (SRY (sex determining region Y)-box 2), NANOG (homeobox transcription factor) and LIN-28 (zinc-finger CCHC domain-containing protein 1).

Before induction, isolated primary skin cells exhibited the morphology (Figure 1Ca) and antigenic expression pattern of human fibroblasts (Supplementary Figure 1). Several
patient-specific iPSC clones were generated from them and clones have been selected by their morphological similarity to human ES cells and expanded (Figure 1C). Two iPSC lines were chosen, further characterized and used for differentiating into patient-specific CMs. As a control, iPSCs generated from a healthy subject were used (Supplementary Figure 2).23

As a first step, we verified that iPSCs generated were genetically matched to the donor and that those derived from the patient carried the heterozygous p.Glu2311Asp RyR2 gene mutation (RyR2-He+/−/C0), by direct sequencing (Figure 1D). No chromosomal abnormalities were detected by karyotype analysis (Figure 1E).

To establish that reprogramming had occurred correctly and that the selected iPSC clones were pluripotent, we tested whether these lines expressed pluripotency markers by verifying alkaline phosphatase activity (Figure 1Cc and Supplementary Figure 2C), the expression of 'stemness'-associated antigens (tra1-60 (TRA1–60) and stage-specific embryonic antigen 4 (SSEA4)) and transcription factors (OCT4, REX1 (RNA exonuclease 1 homolog), DNA (cytosine-5)-methyltransferase 3β (DNMT3B)) with different approaches, that is, immunofluorescence staining (Figure 1C and Supplementary Figure 2), real-time polymerase chain reaction (PCR) (Supplementary Figure 3A) and fluorescence-activated cell sorting (FACS) analysis (Supplementary Figures 3B and C).

Pluripotent cells are by definition capable of differentiating into all cell types of the body. Accordingly, iPSCs are able to spontaneously differentiate into cell types derived from each of the three germ layers when cultured in suspension to form EBs. To test the developmental properties of the selected iPSC lines, we induced differentiation with the EB aggregation method: immunohistochemical analysis (Figure 2A and Supplementary Figure 4) and semiquantitative real-time PCR (Figure 2B) revealed that the EBs contained cells expressing markers of the ectodermal (NCAM1 (neural cell adhesion molecule 1), KRT14 (epidermal keratin 14), βIII-tubulin, nestin), mesodermal (α-smooth muscle actin, desmin, PECAM1 (platelet/endothelial cell adhesion molecule 1) and cardiac genes) and endodermal (GATA6, SOX17 (SRY-box containing gene 17) and α-fetoprotein) lineages.

Moreover, control- and CPVT-iPSC injected into immunocompromised mice had the ability to form teratomas containing derivatives of all the three germ layers. This provided more stringent evidence of the pluripotency of these lines (Figure 2C).

Altogether, these data indicate that we have reprogrammed fibroblasts from a patient with CPVT into iPSC.

Figure 1 Generation of iPSC from a CPVT patient skin biopsy. (A) Pedigree of the RyR2-He+/−/C0 CPVT kindred modeled in this study. Proband (II-2) is indicated by an arrow. Filled symbols indicate clinically and genetically affected subjects. Half-black symbols indicate genetically affected individuals, and upper half-black symbols indicate sudden cardiac death cases. Square = male; circle = female. (B) Example of bidirectional ventricular tachycardia recorded off-therapy in the proband (paper speed 25 mm/s). (C) Representative images of dermal fibroblasts derived from the CPVT patient (a) and of an iPSC colony derived from the patient’s fibroblasts (b) showing alkaline phosphatase activity (c) and positivity for the pluripotency markers OCT4 (d), TRA1-60 (e) and SSEA4 (f). Scale bars = 100 μm. (D) Sequencing analysis confirming that the CPVT-iPSC line (He) carried the specific G-to-C mutation on one allele of the RyR2 gene, whereas control-iPSC (WT) did not show any genetic alteration. (E) iPSC lines maintained a normal karyotype after expansion
Cardiac differentiation. As a next step, we induced iPSC to differentiate toward the cardiac lineage. Control- and CPVT-iPSC lines developed spontaneously contracting areas (Supplementary Movie 1) expressing cardiac-specific channel and structural genes (Figures 3a and b). Importantly, western blot analysis revealed specific expression of RyR2 in iPSC-derived beating explants, either wild-type (WT) or CPVT, at comparable levels (Figures 3b and c). Immunostaining analysis confirmed the presence and the distribution of RyR2 in cells isolated from the contracting areas (Figure 3d and Supplementary Movie 2).6,24

Characterization of iPSC-derived CMs reveals differentiation of heterogeneous populations of cardiac cells. As a first step in the characterization process, we evaluated the electrical activity of control- and CPVT-iPSC-derived CMs.

The analysis of the main electrical features evidenced that the mean amplitude of the AP was around 20 mV higher in CPVT-CMs than in control-CMs, whereas the action potential duration at 30, 50 and 90% of repolarization (APD30, APD50 and APD90 respectively), the maximum diastolic potential (MDP) as well as the maximal rate of depolarization and...
maximal upstroke velocity were similar in both groups (Figure 3f).

Importantly, during the diastolic depolarization phase, CPVT-CMs had delayed afterdepolarizations (DADs), a prominent feature of mature CMs with a CPVT phenotype (Figure 3e).

Furthermore, as already reported in the literature, a more detailed electrical characterization of both control and CPVT cell lines showed that the differentiation process of these cells reflected the heterogeneity seen in the heart.3,11,24 In particular, analyzing the cells for a variety of parameters, including the maximal upstroke velocity (dV/dt max), APD50, APD90, and AP amplitude, it was possible to cluster two distinct populations of iPSC-derived CMs (iPSC-CMs) in both the cell lines: nodal cells (i.e. cells from the AV node), which were distinguishable because of their pronounced phase 4 depolarization preceding the onset of the AP, and working myocardial cells (i.e. atrial and ventricular chamber), which presented the typical plateau phase and, therefore, had the longest AP duration (Supplementary Figure 5).11,24

β-Adrenergic stimulation-induced DADs and triggered activity in CPVT-CMs. To assess the effect of β-adrenergic stimulation on CPVT-CMs, we recorded evoked (Figure 4a) and spontaneous (Figure 4b) APs before and after superfusion with the β-adrenergic agonist isoproterenol (Iso) (1 μM). CPVT-CMs presenting spontaneous AP developed DADs already at basal conditions in 43% of the cases (15 out of 35), and β-adrenergic stimulation significantly increased the frequency of this phenomenon, a hallmark of mature CMs from CPVT patients (75% of the cases, 12 out of 16; Figure 4b, indicated with black arrows). Furthermore, CPVT-CMs also developed DADs and triggered activity (TA) when paced at 0.5 Hz (12%, 2 out of 16, Figure 4a), confirming the
ability of our model system to recapitulate the disease phenotype in vitro.

**KN-93 exerts an antiarrhythmic effect on CVPT-CMs.** To prove the feasibility of our iPSC-based model for drug discovery and for testing the efficacy of novel therapeutic molecules, we cotreated CPVT-CMs with KN-93 (2-[N-(2-hydroxyethyl)]-N-(4-methoxybenzenesulfonyl)]-N-methylbenzylamine), a CaMKII inhibitor. We recently reported that KN-93 is the most powerful antiarrhythmic agent that prevents ventricular arrhythmias in our RyR2 R4496C+/−/C0 knock-in mouse model of CPVT. KN-93 was also capable of abolishing DADs and TA in isolated CMs in vitro.21

Consistent with the findings obtained in CMs derived from the CPVT knock-in mice, 1 μM KN-93 blunted Iso-induced DADs in iPSC-derived CPVT-CMs (n = 7, versus Iso, P < 0.05; Figure 4), whereas the inactive stereoisomer KN-92 (2-[N-(4-methoxybenzenesulfonfyl)]amino-N-(4-chlorocinnamyl)-N-methylbenzylamine, phosphate) (1 μM) has no effect on these arrhythmic events (n = 5, P < 0.05 versus KN-93).

Importantly, the AP morphology of control-iPSC-derived CMs did not undergo any noticeable changes when exposed to KN-93 (data not shown).

We repeated the same protocol using 3D beating clusters of control- and CPVT-iPSC-derived CMs, optically assessing intracellular calcium transients after loading with Fluo-4 (2-[3-[(2-[bis(carboxymethyl)amino]acetic acid). We found that in CPVT-CMs, calcium transients initiated from multiple locations within each beating cluster (n = 8; Figure 5b and Supplementary Movie 3) and that the impulses collided during the propagation, generating a non-homogeneous spreading of calcium transients. This condition was exacerbated by β-adrenergic stimulation during superfusion of the preparation with Iso. In this setting, the number of areas generating spontaneous calcium transients was increased. When we superfused the CPVT multicellular preparations, we observed that KN-93 stabilized calcium activation, reverting to the typical pattern observed in normal tissue, characterized by a single initiation site located on the periphery of the preparation where the...
source-to-sink load was favorable.\textsuperscript{25} As expected, control beating clusters had a single region of calcium impulse initiation under basal conditions and during Iso administration \((n = 6; \text{Figure 5a})\). Moreover, in 75% of the experiments (six out of eight), the upstroke of the Ca\textsuperscript{2+} transient in CPVT clusters in the presence of Iso had a double slope before reaching the peak (Figure 5b, middle panel). To note, KN-93 recovered this abnormal feature of the calcium upstroke. This may explain why the rate of intracellular calcium increase \((\text{dCa}^{2+}/\text{dt})\) after the addition of the CaMKII inhibitor slightly decreased (Figure 6c, versus Iso, not statistically significant), whereas the time to reach the peak was significantly reduced \((P < 0.05, \text{versus Iso}; \text{Figure 6b})\).

**Discussion**

A little more than a decade ago, mutations in the cardiac ryanodine receptor gene \((\text{RyR2})\) were first associated with CPVT, a life-threatening inherited arrhythmogenic disorder\textsuperscript{15}. Since then, much has been learnt about the pathogenesis of this disease: experimental findings from lipid bilayers as well as knock-in and knockout mouse models suggested that the mechanism underlying the onset of arrhythmia in CPVT patients strictly relies on defective Ca\textsuperscript{2+} mobilization within the CM during excitation-contraction coupling. Diastolic Ca\textsuperscript{2+} leak from the sarcoplasmic reticulum is believed to be the major player for the development of DADs, typical markers of electrical instability in CPVT-CMs. DADs are elicited by intracellular calcium load, which activates the membrane Na\textsuperscript{+}/Ca\textsuperscript{2+} exchanger in an electrogenic mode derived by the exchange of one Ca\textsuperscript{2+} for three Na\textsuperscript{+}, leading to diastolic membrane depolarizations that may reach the activation threshold for inward sodium current and generate triggered beats that may eventually lead to sustained arrhythmias\textsuperscript{26,27}.

The development of novel therapeutic approaches has been limited and the use of implantable defibrillators remains the therapy of choice for patients unresponsive to the therapeutic options. Furthermore, the only disease models of CPVT are the knock-in mice that have been used by us, and others, to test new drugs\textsuperscript{21}. However, the results obtained in myocytes from mice leaves investigators with the uncertainty of whether the antiarrhythmic effect seen is replicated in humans.

Clearly, the inability to study the disease and test new treatments in human diseased CMs represents a major limitation. Furthermore, accessibility to human cardiac tissue is limited to heart surgery or to post mortems.

The advent of human iPSC technology may solve these issues and revolutionize the investigation of pathological molecular events driving human diseases: these cells offer an
This study confirms the potential for human iPSC to model inherited arrhythmic syndromes by establishing a patient-specific model of CPVT in which to test therapeutic interventions that have been successful in our murine model of CPVT.

We performed a systematic study of the electrical properties of iPSC-derived CMs from WT and CPVT genotypes using the whole-cell patch clamp technique on spontaneously beating cells at 90 days after the formation of embryoid bodies (EBs). We did not find great heterogeneity in the electrical parameters between the two different groups of cells, but it was possible to highlight a pronounced within-group variability. Furthermore, although the distinction between atrial- and ventricular-like myocytes was not so defined, because of the existence of cells with an intermediate phenotype, we were able to cluster the cells into ‘working-like’ and ‘nodal-like’ CMs. However, the impressive heterogeneity in the electrical parameters is a striking feature of the published electrophysiology data on iPSC-derived myocytes. This could be because of the different stage of maturation of the cells and/or to the sophisticated architecture of the electrical structure of the heart reproduced in vitro.

Importantly, whole-cell patch clamp and microelectrode recordings revealed that CPVT-iPSC-derived CMs exhibited the key features of catecholaminergic-mediated arrhythmogenesis. Hence, clear DADs were observed in most of the CPVT-iPSC-CMs studied. This became even more prominent after activation of the β-adrenergic signaling pathway and gave rise to TA beats.

We also performed impulse-initiation optical mapping in order to determine the electrophysiological phenotypic defects in the context of the spontaneous beating cluster and thus to seek the main electrical disturbance of CPVT-induced arrhythmia. We have used this technique for the first time to build a spatiotemporal high-resolution map for studying the characteristics of intracellular calcium propagation in the context of the whole CPVT cluster. We detected that calcium transients were initiated within a delay of a few milliseconds from multiple regions within the cluster; this resulted in a double rise in the intracellular calcium transients, leaving the reuptake not affected. This is in contrast with recent observations by Novak and co-workers, where a prominent diastolic calcium rise was demonstrated as an indication of DADs. This discrepancy may be due to the diverse technical approaches employed to optically assess calcium transients (ratiometric versus non-ratiometric dyes). On this point, it is worth noting that both types of dyes possess high affinity for Ca-release and Ca, and this can artificially prolong the calcium transient and confound interpretation in CMs and tissues.

Our results offer a ‘side’ view of CPVT, in which the occurrence of calcium transient initiation at multiple foci constitutes the arrhythmogenic substrate by underlying altered gating kinetics of RyR2 (which may explain the double upstroke).

Furthermore, in agreement with data previously obtained in the RyR2(R4496C) CPVT mouse model, we demonstrate that CaMKII inhibition prevents β-adrenergically induced arrhythmogenesis also in patient-specific CMs. Thus, this technique opens up the possibility of testing the response to therapy of individual patients within the clinic. This transition from bench to bedside is most exciting. However, the technology needed to generate iPSC-derived CMs is still expensive and time consuming. Nonetheless, we anticipate the advent of novel technology that will reduce the ‘biopsy-to-human-CMs’ time.

A few tests of substances as putative therapeutic agents on iPSC-based CPVT models have already been reported. For example, flecainide has been recently proposed as an antiarrhythmic drug in mice and human. However, there are still uncertainties on the mechanism that drives its antiarrhythmic activity. Although some authors believe that flecainide acts by inhibiting RyR2’s open state, we supported an alternative hypothesis and demonstrated that the sodium channel blockers of the drug is preventing DADs to activate and generates triggered automaticity. This hypothesis was recently supported by Sikkel et al. Another potential
therapeutic agent for CPVT is dantrolene, a unique and very effective therapeutic option for malignant hypertrophias: this substance has been shown to act by stabilizing interdomain interaction of RyR2 and decreasing loss of Ca\(^{2+}\) from sarcoplasmic reticulum.\(^{6,34,35}\) In the present report, we propose inhibition of CaMKII as a potential therapeutic option for treating arrhythmias in CPVT. CaMKII is activated by multiple pathways and, within the CM, primarily acts by phosphorylating the main components of the calcium handling machinery and, as such, has a clear relevance in the pathophysiology of CPVT. Inhibition of this pathway has been shown to be potentially advantageous compared with \(\beta\)-blockers, the conventional therapy for CPVT patients; however, the use of CaMKII inhibitors in the clinical setting is still limited by the lack of molecules with target- and tissue-specificity.\(^{36}\) The development of a human CPVT model system and the demonstration of its ability to specifically respond to KN-93 (no activity of the inactive analog KN-92 \(b\) was detected) is instrumental to future investigations on identifying specific targets and devising effective strategies for the use of CaMKII inhibition in the clinical setting.

In conclusion, our work contributes to the implementation of the available CPVT mutant models, which is mandatory for establishing a direct relationship between specific mutations and the observed functional effects, as well as determining potential side effects and is fundamental for validating such findings in the perspective of customized patient treatment.

Materials and Methods

Cell culture. Dermal fibroblasts were obtained by enzymatic digestion from 3 to 4 mm skin biopsies of a patient diagnosed with CPVT after written informed consent. Isolated fibroblasts were cultured in DMEM–low glucose/F12 (1:3) supplemented with 10% fetal bovine serum (FBS), glutamine, 0.1 mM non-essential amino acids and antibiotics.

Mouse embryonic fibroblasts (MEFs) were isolated from E12.5–13.5 embryos, following a standard protocol.\(^{37}\) Inactivated MEFs were prepared from cells at passage 3 by treatment with mitomycin C (10 \(\mu\)g/ml) for 3 h at 37 C. After derivation, iPSCs were initially grown on a MEF feeder layer in human embryonic stem cell (ES) medium, that is, knockout DMEM supplemented with 20% knockout serum replacement, 2 mM glutamax, 0.1 mM non-essential amino acids, 1% B27 supplement without vitamin A, 1% N2 supplement, 0.1 mM \(\beta\)-mercaptoethanol, 50 \(\mu\)g/ml penicillin, 50 \(\mu\)g/ml streptomycin (all from Invitrogen), Life Technologies, Carlsbad, CA, USA), and 20 ng/ml human basic fibroblast growth factor FGF (Milltenyi Biotech, Bergisch Gladbach, Germany). At passage 2–3, iPSC lines were adapted to grow on Matrigel (human ES-qualified Matrix from BD Biosciences, Franklin Lakes, NJ, USA) in mTESR1 medium (STEM Cell Technologies, Vancouver, BC, Canada) as described.\(^{23}\)

Human iPSC generation and characterization. Reprogramming was induced by lentiviral infection, as described.\(^{38,39}\) In brief, lentiviral particles were produced in human embryonic kidney 293T cells (HEK-293T cells) by independent transfections of the four ‘pluripotency’ genes Oct4, Sox2, Nanog and Lin28 (Addgene plasmids 16579, 16577, 16578 and 16 580 from Thomson Laboratory, University of Wisconsin, Madison, WI, USA) using the calcium phosphate method.\(^{40}\) Viral supernatants were collected at 30 h and used fresh for the infection.

Low-passage fibroblasts were seeded at 8 \times 10\(^3\) cells per 100 mm dish on the day before the infection. The cells were then infected two times using an equal amount of lentiviral particles for each gene in the presence of 4 \(\mu\)g/ml polybrene. Six days later, infected fibroblasts were seeded onto MEF feeders at a low density (5 \times 10\(^3\) cells per 100 mm dish). The next day, the medium was replaced with regular human ES cell culture medium supplemented with basic FGF.\(^{38}\) Valproic acid (0.5 mM) was applied for 10 days\(^{31}\) to improve the efficiency of the reprogramming process. iPSC colonies became evident around days 21–25 after infection and were mechanically isolated based on their ES-like morphology. Isolated clones were expanded and their pluripotency characterized through the evaluation of ‘stemness’ marker expression and the analysis of their developmental competence in vitro (EBs assay) and in vivo (teratoma formation assay).\(^{3}\)

Two clones for each subject were used for the experiments.

Immunohistological analysis and alkaline phosphatase activity. Cells were fixed in 4% paraformaldehyde (PFA) for 20 min and permeabilized with 0.2% Triton for 10 min. Blocking of unspecific sites was achieved by incubation with 10% donkey or goat serum (Sigma-Aldrich, St. Louis, MO, USA) for 1 h at room temperature. Cells were stained with several primary antibodies, specific for either ‘stemness’ or differentiation markers: human fibroblast surface protein (Clone B1/16, mouse monoclonal, 1:100; Sigma-Aldrich), human Oct4 (mouse monoclonal, 1:500; Millipore, Billerica, MA, USA), human TRA–1–60 (mouse monoclonal, 1:100; Stem Cell Technologies), human SSEA-4 (mouse monoclonal, 1:100; Stem Cell Technologies), human \(\alpha\)-tubulin (mouse monoclonal, 1:100; Promega, Madison, WI, USA), human nestin (mouse monoclonal, 1:100; Millipore), human smooth muscle actin (mouse monoclonal, 1:20; Dako, Glostrup, Denmark), human \(\alpha\)-1-fetoprotein (rabbit polyclonal, 1:100; Dako), human \(\alpha\)-sarcoceric actin (rabbit polyclonal, 1:400; Abcam, Boston, MA, USA), \(\alpha\)-actinin (mouse monoclonal, 1:500; Sigma-Aldrich) and ryanodine receptor 2 (rabbit polyclonal, 1:100; Alomone labs, Jerusalem, Israel). Alexa-Fluor-488- and -555-conjugated secondary antibodies were used for specific detection, whereas nuclei were stained with 4',6-diamidino-2-phenylindole (DAPI). Coverslips were mounted using Vectashield mounting medium (Vector Laboratories, Burlingame, CA, USA). Confocal microscopy was performed with a Leica TCS-SP2 digital scanning confocal microscope equipped with a HCX PL APO \(\times 40\)numerical aperture = 1.25 oil immersion objective. The pinhole diameter was kept at Airy 1. Images were exported to Adobe Photoshop (Adobe Systems, Mountain View, CA, USA) and created with Adobe illustrator (Adobe Systems).

Alkaline phosphatase activity of iPSC lines was determined using the Alkaline Phosphatase Detection Kit (Millipore), after cell fixation in 4% PFA, according to the manufacturer’s instructions. Lines were considered positive when alkaline phosphatase activity was detected in more than 95% of iPSC lines (two clones each condition were analyzed).

RNA extraction and RT-PCR. Total RNA was isolated using Trizol (Invitrogen), treated with amplification grade DNase I (Invitrogen) and reverse transcribed to cDNA (SuperScript III First-Strand Synthesis System; Invitrogen). Real-time PCR was carried out on an ABI7900HT (Applied Biosystems, Life Technologies, Carlsbad, CA, USA) using either the Taqman Gene Expression Assay or the SybrGreen PCR Master Mix (Applied Biosystems), and data were analyzed with REST (Relative Expression Software Tool) software (http://www.gene-quantification.de/rest.html).\(^{42}\) Expression of cardiac-specific genes in cDNA from beating explants was assessed with regular RT-PCR using specific primers. A complete list of the primers used in these experiments is provided in Supplementary Table 1.

Flow cytometry analysis. Dermal fibroblasts and iPSCs were harvested and dissociated into single cells using Trypsin and Tryple Express (Invitrogen). A list of the antibodies used for surface markers was assessed on fresh cell samples. Anti-CD13 (mouse monoclonal, 1:100; eBioscience), anti-CD45-APC, anti-CD15-PE, anti-SSEA4-FITC and anti-TRA1-60-PE were from BD Pharmingen (San Diego, CA, USA). Analyzes were carried out on a FACS Canto flow cytometer (Beckton Dickinson, Franklin Lakes, NJ, USA). Data were analyzed with DIVA software (Beckton Dickinson).

Western blot analysis. Whole-cell lysates were obtained from control (WT) and CPVT iPSC-derived beating explants and analyses preformed using 25 \(\mu\)g of proteins following standard procedures. Proteins from human fetal heart (FH) were used as positive control. Monoclonal anti-RyR2 (1:1000; Thermo Fisher, Waltham, MA, USA) and polyclonal anti-\(\beta\)-Actin (1:2000; Santa Cruz Biotechnologies, Dallas, TX, USA) antibodies were used for detection. Quantification of RyR2 expression levels was determined using Fiji software (Open Source image processing package available at the website: http://fiji.sc/Fiji).\(^{36}\)

Genomic sequencing and karyotyping. Genomic DNA was isolated from control- and CPVT-derived iPSC lines (two clones each) by DNeasy Blood
and tissue kit (Qiagen, Venlo, Limburg, Netherlands). Purified DNA was amplified with Bid Dye Terminator v.1.1 Sequencing RR-100 (Applied Biosystem) with specific primers and analyzed with a 3130x Genetic Analyzer (Applied Biosystem and Hitachi, Chiyoda, Tokyo, Japan). Chromosomal G-banding analysis was performed by the University of Milan-Bicocca Cytogenetics Laboratory (Milan, Italy), using standard procedures.

**Spontaneous differentiation and cardiac induction.** Control and CPVT iPSCs were differentiated by aggregation into EBs: iPSC colonies were detached using 1 mg/ml dispase (Roche, Basel, Switzerland) and plated onto ultra-low-attachment plates (Corning, Incorporated, Corning, NY, USA) in EB differentiation medium, that is, DMEM/F12 medium supplemented with 20% FBS, 0.1 mM non-essential amino acids, glutamine and antibiotics. After 7 days, EBs were plated onto gelatin-coated dishes for further differentiation.

For cardiac lineage induction, ascorbic acid (50 μg/ml) was added to the medium. Spontaneously contracting areas, which appeared 12–30 days after EB plating, were manually microdissected and plated onto fibronectin-coated plates for further differentiation for an additional 45–90 days. Explants were maintained in EB differentiation medium supplemented with FBS at only 2%. For single-cell analyses (electrophysiological and immunofluorescence analyses), cells were dissociated as described previously and plated onto fibronectin-coated plastic or glass 2-well chamber slides (Nunc, Naige Nunc International, Penfield, NY, USA).

**Teratoma assay.** iPSC lines were harvested by dispase treatment, resuspended in X-VIVO medium (Lonza, Basel, Switzerland), and injected subcutaneously into immunodeficient mice (NOD-SCID or Rag2−/− mice homozygous for the scid mutation (severe-combined immunodeficiency) are severely deficient in functional B and T lymphocytes)). Teratomas formed 9–15 weeks after injection were collected and processed according to standard procedures for paraffin embedding and hematoxylin-eosin and immunohistochemical staining.

**Recording of APs.** Cells were seeded on poly-lysi ne-like-covered slides (Lab-Tek II, Nunc) and kept in differentiation medium for about 2 months. APs from spontaneously contracting iPSC-CMs were recorded using the patch-clamp technique in the whole-cell configuration with a Multiclamp 700B amplifier (Axon Instruments, Sunnyvale, CA, USA). The experiments were performed at 37 °C under continuous perfusion of extracellular solution containing (in mM): 140 NaCl, 4 KCl, 2 CaCl2, 1 MgCl2, 10 Hepes and 5 glucose (pH adjusted to 7.40 with NaOH). Patch-clamp pipettes, formed from borosilicate glass with a P-97 horizontal puller (Sutter Instruments, Novato, CA, USA), and had a resistance of 2–3 MΩ when filled with an intracellular solution containing (in mM): 20 KCl, 120K-aspartate, 1 MgCl2, 4 Na2-ATP, 0.1 GTP, 10 glucose and 10 Hepes (pH adjusted to 7.20 with KOH). Some experiments were carried out with intracellular electrophysiology recordings. In this case, spontaneously beating EBs were impaled using sharp glass microelectrodes with resistances ≥10 MΩ. Electrode capacitance was nulled and the recordings were made using the previously described Multiclamp 700B amplifier in gap-free mode. Solutions containing 1 μM Isos, 1 μM KN-93 or KN-92 were prepared fresh before the experiments and applied using a gravitational flow system for 2–3 min before data collection. All signals were acquired at 10 kHz, digitized (Digidata 1332A, Axon Instruments) and analysed with PCLAMP 9.2 software (Axon Instruments).

**Definition of delayed APs and TA.** We defined DADs as low-amplitude depolarizations following completion of repolarization, and have an amplitude ≥5% of the preceding AP. TA was defined as an AP developing from a DAD rather than from an external stimulus.

**Fast optical mapping of intracellular calcium transient.** Intracellular calcium transient characteristics were measured as described previously. Brieﬂy exotropic clusters from CPVT and WT were excised and recultured onto 22 mm glass coverslip. After 48–96 h, the coverslips were immersed in a 1 ml solution containing culture medium plus 2.5 μmol of Fluor-4 AM (Invitrogen, Life Technologies) and incubated for 20 min at 37 °C. Afterwards, the coverslips were mounted onto a custom-made microscope chamber and perfused with Tyrode solution at 37°C containing (in mM): 140 NaCl, 4 KCl, 2 CaCl2, 1 MgCl2, 10 Hepes and 5 glucose (pH adjusted to 7.40 with NaOH). Optical recording of intracellular calcium transient were assessed using a CMOS fast resolution camera (Ultima L, Someda, Costa Mesa, CA, USA) mounted on an inverted microscope (Nikon T/U from Nikon Instruments, Chiyoda, Tokyo, Japan) and acquired for 8 s at 0.5 KHz at ×10 magnification. To reduce the light exposure, the synchronization of the light shutter and the acquisition was achieved using Digidata 1440A (Molecular Devices, Sunnyvale, CA, USA; Criel IT) by programming a dedicate protocol of acquisition. Recordings were analysed using B-analys 1.1208 (Somedia).

**Statistical analysis.** Data are represented as mean ± MISE (or mean ± S.D. where indicated). The significance of differences between two groups was evaluated with unpaired Student’s t-test. P < 0.05 was considered statistically significant. Single asterisk indicates P < 0.05, whereas double asterisks indicate P < 0.01.

**Conflict of Interest**

The authors declare no conflict of interest.

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**Ethical Statement**

The study has been conducted in a safe and ethical manner after the approval of the Institutional IRB. All the subjects involved in the study gave their informed consent to the use of their biological material to this purpose.

**Author Contributions**

EDP, MD, CN, GC and SGP conceived the research and planned the experiments; EDP, FL, MM, JEAC, HH and PP performed the experiments; MD contributes to the discussion of the data; EDP, FL, MM, JEAC, MD, CN, GC and SGP discussed and analyzed the data; and EDP, FL, MM, MD, ON, GC and SGP wrote the article.

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