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## Functional Effects of Schizophrenia-Linked Genetic Variants on Intrinsic Single-Neuron Excitability: A Modeling Study

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### Abstract

**Background**—Recent genome-wide association studies have identified a large number of genetic risk factors for schizophrenia (SCZ) featuring ion channels and calcium transporters. For some of these risk factors, independent prior investigations have examined the effects of genetic alterations on the cellular electrical excitability and calcium homeostasis. In the present proof-of-concept study, we harnessed these experimental results for modeling of computational properties on layer V cortical pyramidal cells and identified possible common alterations in behavior across SCZ-related genes.

**Methods**—We applied a biophysically detailed multicompartmental model to study the excitability of a layer V pyramidal cell. We reviewed the literature on functional genomics for variants of genes associated with SCZ and used changes in neuron model parameters to represent the effects of these variants.

**Results**—We present and apply a framework for examining the effects of subtle single nucleotide polymorphisms in ion channel and calcium transporter-encoding genes on neuron excitability. Our analysis indicates that most of the considered SCZ-related genetic variants affect the spiking behavior and intracellular calcium dynamics resulting from summation of inputs across the dendritic tree.

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**Conclusions**—Our results suggest that alteration in the ability of a single neuron to integrate the inputs and scale its excitability may constitute a fundamental mechanistic contributor to mental disease, alongside the previously proposed deficits in synaptic communication and network behavior.

### Keywords

Biophysical modeling; Functional genomics; Genome-wide association studies; Layer V pyramidal cell; Neuron excitability; Schizophrenia

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Schizophrenia (SCZ) is a severe mental disorder with heritability estimates ranging from .6 to .8 (1). A recent genome-wide association study (GWAS) identified more than a hundred genes exceeding genome-wide significance, confirming the polygenic nature of this psychiatric disorder (2). This remarkable success in gene discovery brings up the next big challenge for psychiatric genetics—translation of the genetic associations into biological insights (3). Attaining this goal is supported by the development of biophysically detailed neuron models, boosted by the recent launch of mega-scale neuroscience projects (4). These models make it possible to investigate SCZ disease mechanisms by computational means, ultimately aiming toward achieving better clinical treatments and disorder outcomes (5,6).

The 108 recently confirmed SCZ-linked loci span a wide set of protein-coding genes (2), including numerous ion channel-encoding genes. The disorder is associated with genes affecting transmembrane currents of all major ionic species, sodium ( $\text{Na}^+$ ), potassium ( $\text{K}^+$ ), and calcium ( $\text{Ca}^{2+}$ ). In addition, some of the SCZ-linked genes are involved in regulation of the  $\text{Ca}^{2+}$  concentration in the intracellular medium (2), which is another great contributor to excitability. It is thus reasonable to hypothesize that the SCZ-linked genes should have an impact on the excitability at the single-neuron level.

We focused our study on cortical layer V pyramidal cells (L5PCs) as a principal computational element of the cerebral circuit. An L5PC extends throughout the cortical depth with the soma located in layer V and the apical dendrite branching into the apical tuft in layer I, and its long axon may project to nonlocal cortical and subcortical areas. The tuft serves as an integration hub for long-distance synaptic inputs and is often considered a biological substrate for cortical associations providing high-level context for low-level (e.g., sensory) inputs to the perisomatic compartment (7). Therefore, the ability of L5PCs to communicate the apical inputs to the soma has been proposed as one of the mechanisms that could be impaired in the mental disease (7). In agreement with this hypothesis, recent psychiatric GWASs consistently reported association of genes coding for the subunits of voltage-gated  $\text{Ca}^{2+}$  channels as risk factors in SCZ and bipolar disorder (8–10).

In the present proof-of-principle study, we applied a model (11) of L5PCs to explore how genetic variants in SCZ-linked genes affect the single-cell excitability. We carried out our study by linking a documented effect of a genetic variant in an ion channel or  $\text{Ca}^{2+}$  transporter-encoding gene to a change in the corresponding neuron model parameter. It should, however, be noted that information does not generally exist for the effect of single nucleotide polymorphism (SNP) variants identified through GWASs on the biophysical parameters required for the computational models. We instead used information obtained

from in vitro studies of more extreme genetic variations, including loss-of-function mutations. A central assumption of this approach is that the effects of SNP variants can be represented as scaled-down versions of those of the more extreme variants and that the emergence of the full psychiatric disease phenotype results from the combined effect of a large number of subtle SNP effects (12,13). A deficit in synaptic communication is likely to contribute to SCZ (2,14–16) but is outside the scope of the present work.

## Methods and Materials

### The L5PC Model

The multicompartmental neuron model used in this work was based on a reconstructed morphology of a layer V thick-tufted pyramidal neuron [cell #1 in (11)]. The model includes the following ionic currents: fast inactivating Na<sup>+</sup> current, persistent Na<sup>+</sup> current, nonspecific cation current, muscarinic K<sup>+</sup> current, slow inactivating K<sup>+</sup> current (I<sub>Kp</sub>), fast inactivating K<sup>+</sup> current, fast noninactivating K<sup>+</sup> current, high-voltage activated Ca<sup>2+</sup> current, low-voltage activated Ca<sup>2+</sup> current, small-conductance Ca<sup>2+</sup>-activated K<sup>+</sup> current (I<sub>SK</sub>), and finally, the passive leak current. See Supplement 1 for the model equations, and for the simulation codes, see ModelDB entry 169457 (<http://senselab.med.yale.edu/ModelDB/showModel.cshtml?model=169457>).

### Genes Included in the Study

Since we did not aim to provide a comprehensive evaluation of a representative fraction of the genetic risk factors of SCZ but rather to provide a proof of principle of the computational modeling approach, we selected the genomic loci using the following approach. We based our study on a recent GWAS (2), which reported significantly associated SNPs that were scattered across hundreds of genes with a variety of cellular functions. We concentrated on those genes that encoded either ionic channels or proteins contributing to transportation of intracellular Ca<sup>2+</sup> ions.

We used the SNP-wise *p* value data of Ripke *et al.* (2), and for each gene of interest we determined the minimum *p* value among those SNPs that were located in the considered gene. We performed this operation for all genes encoding either subunits of voltage-gated Ca<sup>2+</sup>, K<sup>+</sup>, or Na<sup>+</sup> channel; subunits of a small-conductance calcium-activated potassium (SK), leak, or hyperpolarization-activated cyclic nucleotide-gated channel; or Ca<sup>2+</sup>-transporting ATPase. The genes *CACNA1C*, *CACNB2*, *CACNA1I*, *ATP2A2*, and *HCN1* possessed a small minimum *p* value each ( $p < 3 \times 10^{-8}$ )—these genes were also highlighted in the locus-oriented association analysis as performed in Ripke *et al.* (2). To extend our study to explore a larger set of genes, we used a more relaxed threshold ( $p < 3 \times 10^{-5}$ ) for the minimum *p* value and obtained the following genes in addition to the previously mentioned ones: *CACNA1D*, *CACNA1S*, *SCN1A*, *SCN7A*, *SCN9A*, *KCNN3*, *KCNS3*, *KCNB1*, *KCNG2*, *KCNH7*, and *ATP2B2*. Of these, we omitted the genes that are not relevant for the firing behavior of an L5PC. It should be noted that we used the SNPs reported in Ripke *et al.* (2) only to name the above SCZ-related genes, and due to lack of functional genomics data, we could not include the actual SCZ-related SNPs in our simulation study.

Instead, we searched in PubMed for functional genomic studies reporting the effects of any genetic variant of the above genes. For details, see Supplement 1.

## Results

### A New Framework for Bridging the Gap Between GWASs of SCZ and Computational Neuroscience

In this work, we reviewed the literature on effects of variants in SCZ-related genes on ion channel behavior and intracellular  $\text{Ca}^{2+}$  dynamics and interpreted the reported effects in the context of our neuron model parameters. An overview of the relevant studies is given in Table 1, while the effects of each variant on the L5PC model parameters are given in Table 2. These data gave us a direct interface for linking a change in the genomic data, such as an SNP or an alternative splicing, into a change in neuron dynamics. The reported data, however, often corresponded to variants with large phenotypic consequences that, in general, are absent in SCZ patients. To simulate subtle cellular effects caused by the common SNPs related to SCZ (17), we downscaled the variants of Table 2 by bringing the changed parameters closer to the control neuron values when the reported change caused too large an effect in the neuron firing behavior. Our approach is illustrated in Figure 1.

The variants in Table 1 were 23 in total, although some of them represented a range of effects of different variants (Table S1 in Supplement 1). The entries corresponded to variants of genes encoding for  $\text{Ca}^{2+}$  channel subunits (*CACNA1C*, *CACNB2*, *CACNA1D*, *CACNA1I*), intracellular  $\text{Ca}^{2+}$  pumps (*ATP2A2*, *ATP2B2*),  $\text{Na}^+$  channel subunits (*SCN1A*, *SCN9A*),  $\text{K}^+$  channel subunits (*KCNS3*, *KCNB1*, *KCNN3*), and a nonspecific ion channel subunit (*HCNI*). In the following, we present simulation results for the L5PC model equipped with some of these downscaled variants (we refer to these model neurons as variant neurons or simply as variants). As we did not know the quality of the effects of the actual SCZ-related polymorphisms, we performed the simulations for a range of differently scaled variants, including negative scalings (i.e., opposite effects with regard to the effects reported in Table 2). We concentrated our study on a representative sample of six variants, highlighted in Table 2. These variants represented six genes with different roles in L5PC electrogenesis and a wide range of observed effects (Table S2 in Supplement 1).

#### Variants Show Altered Intracellular ( $\text{Ca}^{2+}$ ) Responses to Short Stimuli

To characterize the implications of SCZ-related genes on the neuron excitability, we started by analyzing the effects of the downscaled versions of genetic variants in Table 2 on the neuron response to a short somatic suprathreshold square-pulse stimulus. Figure 2A and B shows the time course of the membrane potential and the intracellular  $\text{Ca}^{2+}$  concentration ( $[\text{Ca}^{2+}]$ ) for one variant, whereas Figure 2C shows the  $[\text{Ca}^{2+}]$  response in the phase plane for several different variants. The most typical effect of a variant was a deviation in the peak  $[\text{Ca}^{2+}]$ , but differences could also be observed in the rising phase of the  $[\text{Ca}^{2+}]$ , as shown in the phase plane representation in Figure 2C.

The results in Figure 2 show that the  $[\text{Ca}^{2+}]$  response served as a more sensitive indicator of genetic effects than the membrane potential. In *CACNA1C* and *CACNB2* variants, which

affect the high-voltage-activated  $\text{Ca}^{2+}$  current, and in the *CACNA1I* variant, which affects the low-voltage-activated  $\text{Ca}^{2+}$  current, the observed effects on the  $[\text{Ca}^{2+}]$  response were due to changed  $\text{Ca}^{2+}$  channel kinetics. Similarly, in the *ATP2B2* variant, which affects the plasma membrane  $\text{Ca}^{2+}$  ATPase (PMCA) activity, the observed effect was caused by altered intracellular  $\text{Ca}^{2+}$  dynamics. By contrast, the small yet observable differences between control and *SCN1A* variant  $[\text{Ca}^{2+}]$  phase planes were due to alterations in subthreshold membrane potential fluctuations, which caused variation in the activation of  $\text{Ca}^{2+}$  channels.

### Steady-State Firing Is Influenced by the Variants

Next, we investigated the steady-state behavior of the neuron when a direct current was applied to the soma. As shown in Figure 3, the f-I curves (firing frequency as a function of direct current amplitude) of many SCZ-associated variants were notably different from those obtained with the control neuron.

The deviations in the f-I curves in many of the variants can be explained by changes in the  $\text{Ca}^{2+}$ -activated SK current. These changes were caused either by direct alteration of the activation kinetics of SK channels (as in the *KCNN3* variant) or indirectly through the changes in the intracellular  $[\text{Ca}^{2+}]$  response associated with other variants (*CACNA1C*, *CACNB2*, and *ATP2B2*). As an example, the  $\text{Ca}^{2+}$  influx during an action potential was larger in the *CACNA1C* variant neuron compared with the control neuron (compare curves in blue and magenta in Figure 2C), and this led to an increase in the SK current activation, relative to that in the control neuron. The increased  $I_{\text{SK}}$ , in turn, delayed the induction of the next spike and hence resulted in a loss of gain (flattening) in the f-I curve (Figure 3). This finding is in line with previous modeling studies that also have highlighted the role of  $\text{Ca}^{2+}$ -activated  $\text{K}^+$  currents in modulating f-I curves (18,19).

We also studied the steady-state firing behavior by analyzing the limit cycle, i.e., the phase plane representation of the variable of interest after a large number of initial cycles. As the  $[\text{Ca}^{2+}]$  response served as the most sensitive biomarker, we used the  $[\text{Ca}^{2+}]$  limit cycle for this analysis. Figure 4 shows, for some of the gene variants, the  $[\text{Ca}^{2+}]$  limit cycle as a response to a long, square-pulse stimulus.

The different variants had different ways of impacting the shape of the  $[\text{Ca}^{2+}]$  limit cycle. An especially common trait was a shift or compression in the horizontal direction, representing a small increase or decrease of intracellular  $[\text{Ca}^{2+}]$  during steady firing. Figure S1 in Supplement 1 shows the range of  $[\text{Ca}^{2+}]$  values observed during the constant current injection for all variants used in this study. Interestingly, almost all variants gave deviations from the  $[\text{Ca}^{2+}]$  range observed in the control neuron.

The only genes that did not show notable variance for the range of  $[\text{Ca}^{2+}]$  during steady-state firing were *CACNA1I*, *KCNS3*, and *KCNB1*. Despite this, *CACNA1I* did have a clear effect on the  $[\text{Ca}^{2+}]$  phase plane in the single-pulse response (Figure 2C). Conversely, the *KCNS3* and *KCNB1* variants were not found to influence any of the considered aspects of neuron behavior (Figures S1 and S2 in Supplement 1). In these variants, the activation and inactivation of  $I_{\text{Kp}}$  have been modified. Our findings reflect the overall minor role that the current  $I_{\text{Kp}}$  plays in shaping the response properties of the L5PC model neuron: it is only

present in the soma and with relatively small maximal conductance (11). The minor contribution of this current to the model neuron behavior may be in contradiction with experimental evidence, which states that this current constitutes a major proportion of the total outward  $K^+$  currents (20).

### Variants Have an Effect on Integration of Somatic and Apical Inputs

A question that arises from the observation of the above trends is whether, and to what extent, the small deviations from control neuron behavior affect the information processing capabilities of the neuron. In Hay *et al.* (11), it was shown that the L5PC model can be used to describe the  $Ca^{2+}$  spike generation as a response to a combination of stimuli at the apical dendrite and at the soma. Moreover, they showed a good qualitative match with experimental data and model predictions for the neuron responses during an up or down state. In this work, we followed their definition for the down state as the resting state of the neuron and the up state as a state where a subthreshold current is applied to the proximal apical dendrite (11). We also adopted their protocol of combining an apical stimulus and somatic stimulus and studying the effect of order and interstimulus interval (ISI) between the two [see Figure 8 in Hay *et al.* (11)]. This gave us a temporal profile showing a range of ISI for which a large  $Ca^{2+}$  spike is produced. We then estimated the effect of our genetic variants on this temporal profile.

Figure 5 shows the temporal profiles for the six representative variants. Effects of the different variants on the neuron response were visible both during the down and during the up states. Particularly large effects were observed in *CACNA1I* and *SCN1A* variants, which code for the low-voltage-activated  $Ca^{2+}$  channels and transient  $Na^+$  channels that contribute to the sharp rise of voltage during an action potential. These channel types are expressed both in the soma and in the apical dendrite. Although all downscaled variants shared the basic form of the response curves, the shape was clearly altered in some variants. As shown in Hay *et al.* (11), the up state temporal profile of the control neuron showed elevated apical responses when the apical stimulus was applied shortly after the somatic stimulus but not when the apical stimulus was applied first. This order-specific response was changed in some of the variants (e.g., *SCN1A* variant) so that they also produced large  $Ca^{2+}$  spikes when the apical stimulus was applied shortly before the somatic stimulus and could hence alter the order sensitivity of the coincidence detection.

### Variants Show Differences in Inhibition of a Second Apical Stimulus

The alteration of information processing by the variants can also be seen in the sensitivity to successive synaptic inputs at the same locations. To explore this, we gave two successive stimuli to the apical dendrite and varied the interval (ISI) between them. The first stimulus activated long-time scale inhibitory currents, especially  $I_{SK}$  and muscarinic  $K^+$  current, and within some following time window, these currents could inhibit the generation of a subsequent spike. This could be interpreted as a form of single-cell prepulse inhibition: if the initial input caused the neuron to spike, the next input could fail to do so, even at times when the second stimulus was of larger amplitude.

Figure 6 shows how the threshold synaptic conductance needed for inducing a second spike, relative to the one needed for the first spike, depends on the ISI. Interestingly, the threshold conductance curve was affected by many of the variants, especially by the variants of  $\text{Ca}^{2+}$  channel genes. Although the shown effect was partly attributed to the differences in the spiking thresholds between the variants, differences between the variants could still be observed when absolute spiking threshold values for inducing a second spike were considered (data not shown). These findings raise the possibility that the disturbed prepulse inhibition observed in many SCZ patients might be an effect of altered ionic channel properties at the single-neuron level and not only of the modified elements in synaptic circuitry as previously thought (21,22) [cf. (23)].

## Discussion

We presented a framework for studying the effects of genetic variants in genes related to SCZ on layer V pyramidal cell excitability. Our results indicate that most of the studied variants predicted an observable change in the neuron behavior (Figures 2–6). Taken together, these data provide support for using neuronal modeling to study the functional implications of SCZ-related genes on neuronal excitability. Although the analyses presented in this article are specific to L5PCs and SCZ-related genes, our framework may be directly applicable to other cell types and other polygenic diseases, such as bipolar disorder and autism, given an identification of risk genes related to neuronal excitability. Furthermore, our approach can be directly extended to biophysically detailed models of neuronal networks, e.g., Hay and Segev (24).

The results shown in the present article are based on a multicompartmental neuron model (11) employing a certain set of ionic channels. Alternative models with comparable complexity have recently been published. The authors of Bahl *et al.* (25) present and apply a method for fitting the L5PC model to a simplified neuron morphology. In another recent work by Almog and Korngreen (26), the L5PC model was fit to each stained morphology and the corresponding recordings separately. To rule out our conclusions being an artefact of a certain specific morphology, we carried out our analyses on another neuron model fit [cell #1 equipped with channel conductance values from the first column of Table S1 in (11)] and an alternative neuron morphology [cell #2 of (11)]. The trend of the variant effects remained the same in these simulations (Figures S4 and S5 in Supplement 1). More comprehensive analysis could be carried out by using the above-mentioned alternative models or by employing a wider set of neuron morphologies (27), but this is left for future work.

Discovering the disease mechanisms of disorders with polygenic architecture, such as SCZ, requires considering the interplay between many different biological processes and entities. The computational framework presented here allows one to identify physiological mechanisms and biomarkers common across multiple risk-related genes. Although there are many more SCZ-related genes (2) than those listed in Table 1, the identified variants capture multiple types of changes in neuron excitability (Figures 2–6). This allows studying the effects of combinations of different genetic variants in a straightforward manner, as suggested in Figure S3 in Supplement 1, although one should be careful in the interpretation of the results, as certain variants could have nonadditive effects. The gene hits not



considered in the present work contribute to, e.g., expression level and function of ionotropic and G-protein coupled receptors, as well as phosphorylation and dephosphorylation of diverse proteins. It should be noted that several of the identified SCZ-related genes, such as *PTN*, *PAK6*, and *SNAP91*, may also affect neuron morphology (28–31). Our framework could be extended to include some of these effects, but studying the contribution of many signaling genes is out of the scope of the used neuron model.

Linking cellular mechanisms to SCZ can be extremely difficult owing to the high cognitive level on which the disease symptoms usually emerge. There is, however, one important exception, namely, altered prepulse inhibition, which is found in approximately 40% to 50% of SCZ patients (32). In the present study, we showed that the considered genetic variants may have had different tendencies to suppress a second synaptic stimulus, owing only to the differences in the (non-synaptic) ion channel gating or  $\text{Ca}^{2+}$  dynamics (Figure 6). This could be an important contributor to understanding the phenomenon of prepulse inhibition, although deficiencies in synaptic connections are most likely to play a role as well (21). Moreover, altered integration of local and distal inputs in the manner of Figure 5 could possibly underlie incomplete or excessive activation of pyramidal cells in certain brain areas linked to auditory hallucinations [(7); see also discussion of the role of spontaneous activity in primary auditory cortex in triggering verbal hallucinations in (33)]. Experimental testing of the shown model predictions for integration of inputs on a single-cell level is possible as well. Even without genetic manipulations, the downscaled variants of the ion channels could be imitated in vitro by partial pharmacologic blocks that are configured to cause changes in currents comparable with those in our variants. However, in such a case, the fine details on, e.g., current activation time constants would be dismissed.

The genetic variation of ion channels will affect not only the membrane potentials of the neurons but also the transmembrane currents generating the local field potential recorded inside the cortex and the electrocorticography signals recorded at the cortical surface (34,35). Further, the same transmembrane currents also determine the neuronal current dipole moments measured in electroencephalography and magnetoencephalography (36,37). Thus, a natural extension of the present work is to investigate the effects of the genetic variation of the single-neuron current dipole moments (38) and the corresponding single-neuron contribution to the electroencephalography signals. This is left for future work.

A severe limitation of our approach is the lack of data on how different specific SCZ-related genetic polymorphisms affect the ion channel function and  $\text{Ca}^{2+}$  dynamics. In this article, we used literature data on extreme variants (e.g., loss-of-function of function mutations) of the identified SCZ-related risk genes, from a range of cell types, including nonneural tissues. Ideally, one would want to use physiological measures for the relevant genetic variants in the actual cell types (i.e., L5PCs) in cortical circuits, but such data do not currently exist. On the other hand, the diversity of data in the literature that we used and the possible discrepancies in the experimental procedures therein could also reflect in the results shown in this study, although our downscaling procedure constrained such errors from above. Novel methods employing automated cell patching (39) to test the effects of subtle SCZ-related SNPs could provide the solution to both above limitations of our study.

The validity of the downscaling, which for the aforementioned reasons is a central procedure in our framework, is built upon two assumptions: 1) the SCZ-related gene variants do affect neuronal excitability on a single-neuron level, and 2) their effects on ion channel or calcium pump kinetics are smaller than those of certain already documented mutations with extreme phenotypic consequences, such as deafness, hemiplegic migraine, or other pain disorders. The first assumption can be argued for by the multitude of ion channel-encoding genes that have been shown to be related to SCZ risk and the second assumption by the lack of severe body malfunctioning (such as the abovementioned ones) in SCZ patients, but rigorous proofs for both assumptions are yet to be shown. A challenge for future studies is also to decipher how the ion channel densities are affected by the variants in various cell types, an issue that might be important in modeling the effects of SCZ-associated polymorphisms.

Our framework is an early attempt toward understanding the disease mechanisms of polygenic psychiatric disorders by computational means. It binds together genetic and single-neuron levels of abstraction in the big picture of modeling psychiatric illnesses, as recently sketched in Wang and Krystal (5) and Corlett and Fletcher (40). While earlier computational studies considered only effects of single variants on neuron firing behavior (41,42), our framework allows the screening of multiple types of variants and their implications on the neuron excitability. It may serve as a proof of principle for a novel biophysical psychiatry approach, enabling the integration of information from previously disparate fields of study including GWASs, functional genomics, and biophysical computational modeling using models of moderate complexity. We consider the chosen level of complexity a suitable trade-off for this purpose: while more approximate models, such as integrate-and-fire models, could not distinguish between the effects of different genetic variants in an acceptable precision, more detailed models including, e.g., dynamics of protein translation and phosphorylation are likely to be very computation-intensive. The challenge for future work is to extend the study to both larger and finer spatial and temporal scales.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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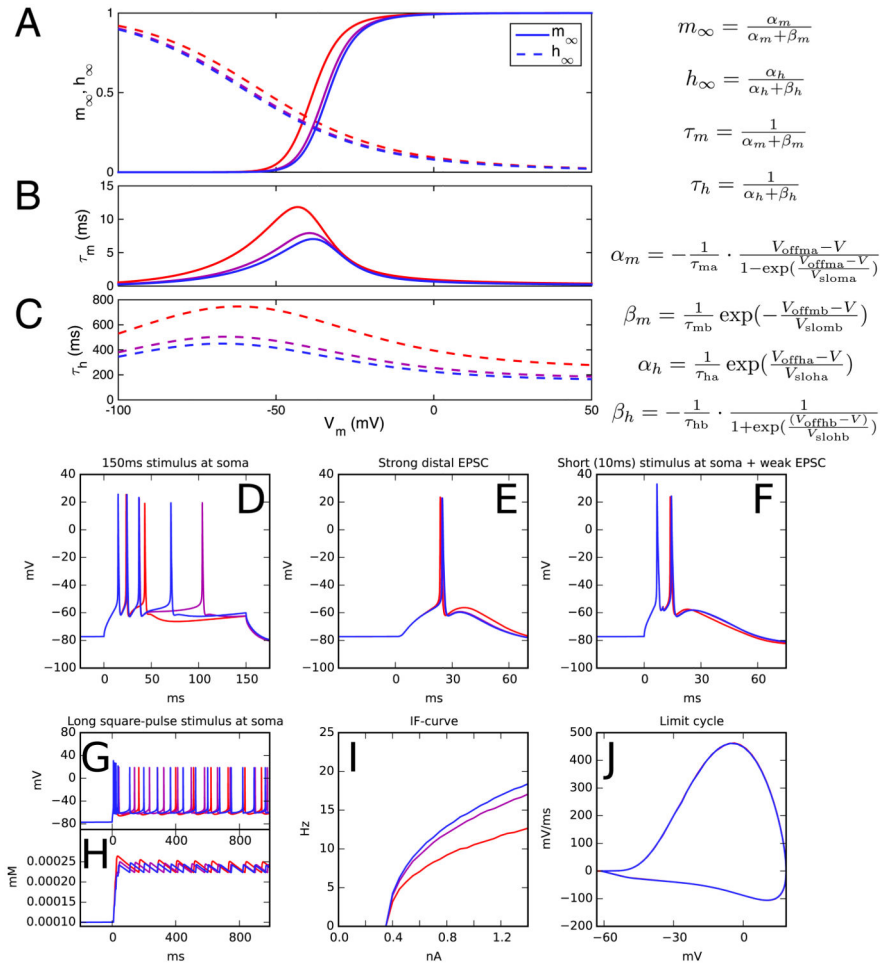
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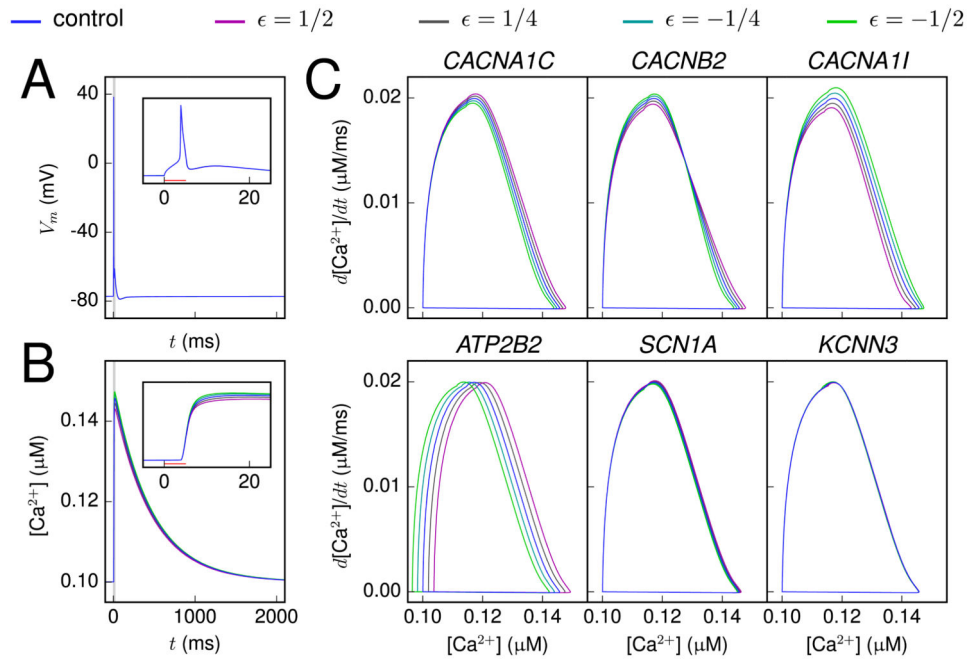
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**Figure 1.**

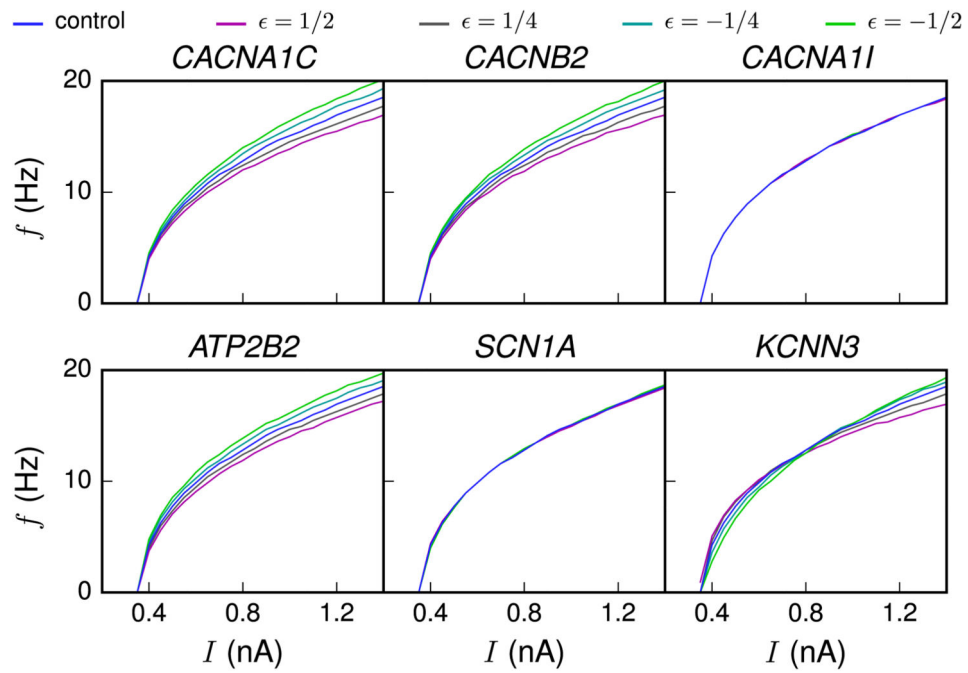
An example of downscaling with a variant (46) of the gene *CACNB2*. Link et al. (46) transfected different variants of *CACNB2* DNA into HEK293 cells together with CaV1.2 subunit DNA and measured the activation and inactivation curves of the calcium currents through the cell membrane. The cells transfected with different variants showed different values of midpoints and time constants for channel activation and inactivation: the midpoints of activation and inactivation varied by  $\pm 4.9$  and  $\pm 5.1$  mV among the variants, respectively, while the time constants of activation and inactivation varied from  $-40\%$  to  $+68\%$  and  $-40\%$  to  $+66\%$ , respectively. Here, we illustrate a variant representing one possible combination of the end points of these ranges. **(A)** The voltage dependence of steady-state activation ( $m_\infty$ ) and inactivation ( $h_\infty$ ) according to the neuron model (11). Red curves represent the unscaled variant: activation parameters  $V_{offma}$  and  $V_{offmb}$  are changed by  $\eta_1 = -4.9$  mV, inactivation parameters  $V_{offha}$  and  $V_{offhb}$  are changed by  $\eta_2 = 5.1$  mV, activation time constants  $\tau_{ma}$  and  $\tau_{mb}$  are increased by 68% ( $\eta_3 = 1.68$ ), and inactivation time constants  $\tau_{ha}$  and  $\tau_{hb}$  are increased by +66% ( $\eta_4 = 1.66$ ). Purple curves show the downscaled  $\varepsilon = 1/2$  variant (see below), and blue curves show the control neuron activation properties. **(B, C)** The voltage dependence of time constants for activation **(B)** and inactivation **(C)**. **(D-J)** Illustration of the scaling conditions. The downscaling is based on

five conditions (Supplement 1), none of which should be violated in the downscaled variant. In panels **(D-F)** (conditions with different scalings I-III), the variant neuron should respond with the same number of spikes as the control neuron (blue). In panel **(I)**, the integrated difference between the variant and control f-I curves (firing frequency as a function of direct current amplitude) should not exceed 10% of the integral of the control curve (condition IV). In panel **(J)**, the difference between the limit cycles should not exceed a set limit (condition V). It can be observed that the unscaled variant (red) violates the conditions I and IV. To scale down the variant, each applied parameter change is brought to a fraction  $c < 1$ , all in proportion, until the threshold of violating/nonviolating variant is found. In this example, the threshold variant corresponds to parameter changes  $c \times \tau_1$ ,  $c \times \tau_2$ ,  $\tau_3^c$ , and  $\tau_4^c$ , where  $c = .4574$ . Any nonnegative value below the threshold value  $c$  yields a downscaled variant that obeys conditions I-V. The purple curves represent the variant corresponding to parameter changes  $c\varepsilon \times \tau_1$ ,  $c\varepsilon \times \tau_2$ ,  $\tau_3^{c\varepsilon}$ , and  $\tau_4^{c\varepsilon}$ , where  $c = .4574$  and  $\varepsilon = 1/2$ . The blue curves show the control neuron firing behavior. **(D)** Somatic membrane potential as a response to a 150-ms somatic square-pulse stimulus (.696 nA). **(E)** Somatic response to a distal apical synaptic conductance with maximum .0612  $\mu$ S. **(F)** Somatic response to a combination of somatic square-pulse current (10 ms  $\times$  1.137 nA) and a distal apical synaptic conductance (maximum .100  $\mu$ S). **(G)** Somatic membrane potential as a response to a long somatic square-pulse stimulus current (1.0 nA). **(H)** Somatic calcium concentration as a response to the stimulus used in **(G)**. **(I)** Spiking frequency as a function of amplitude of the somatic direct current. **(J)** The membrane potential limit cycle corresponding to the late phase of **(G)**. The x-axis represents the somatic membrane potential and y-axis its time derivative. EPSC, excitatory postsynaptic current.

**Figure 2.**

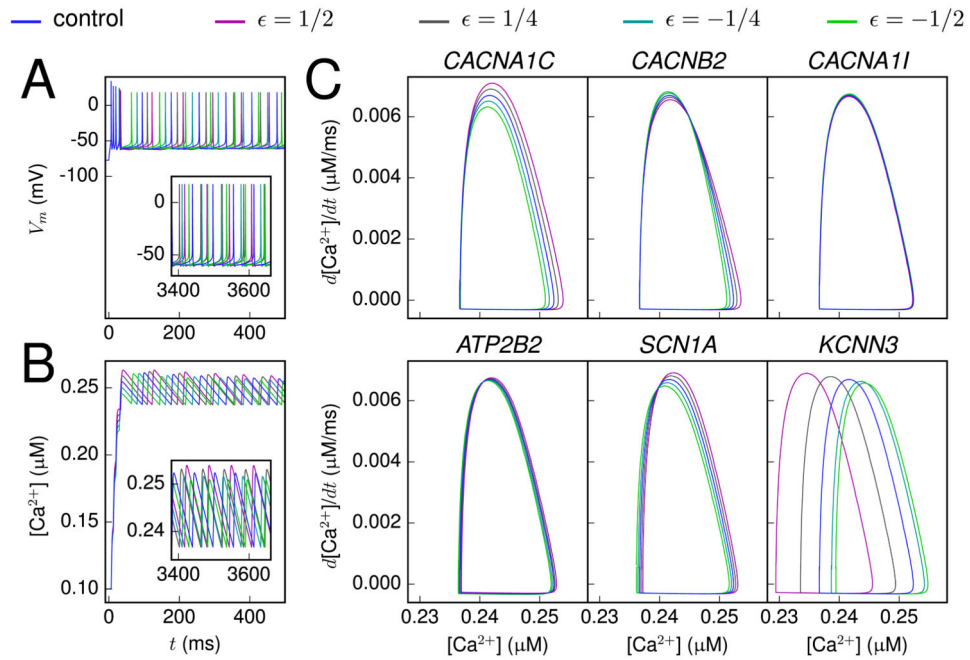
Variants show altered  $[Ca^{2+}]$  response to a short, somatic stimulus. **(A, B)** The membrane potential **(A)** and calcium ( $Ca^{2+}$ ) concentration **(B)** time courses, recorded at soma, as a response to a somatic 5-ms, 1.626 nA square-pulse stimulus. The blue curve shows the control neuron behavior, while the other curves show the behavior of a *CAC-NAII* variant (41) with different scalings (magenta:  $\epsilon = 1/2$ , dark gray:  $\epsilon = 1/4$ , cyan:  $\epsilon = -1/4$ , green:  $\epsilon = -1/2$ ). Inset: Zoomed-in view around the time of the spike. No notable differences between the variants and the control can be observed in the membrane potential time course. **(C)** The  $[Ca^{2+}]$  response plotted in the phase plane for different variants. The behavior of variants of *CACNA1C* (43), *CACNB2* (46), *CACNA1I* (41), *ATP2B2* (52), *SCN1A* (54), and *KCNN3* (62) genes are shown with similar scaling as in **(A)** and **(B)**.





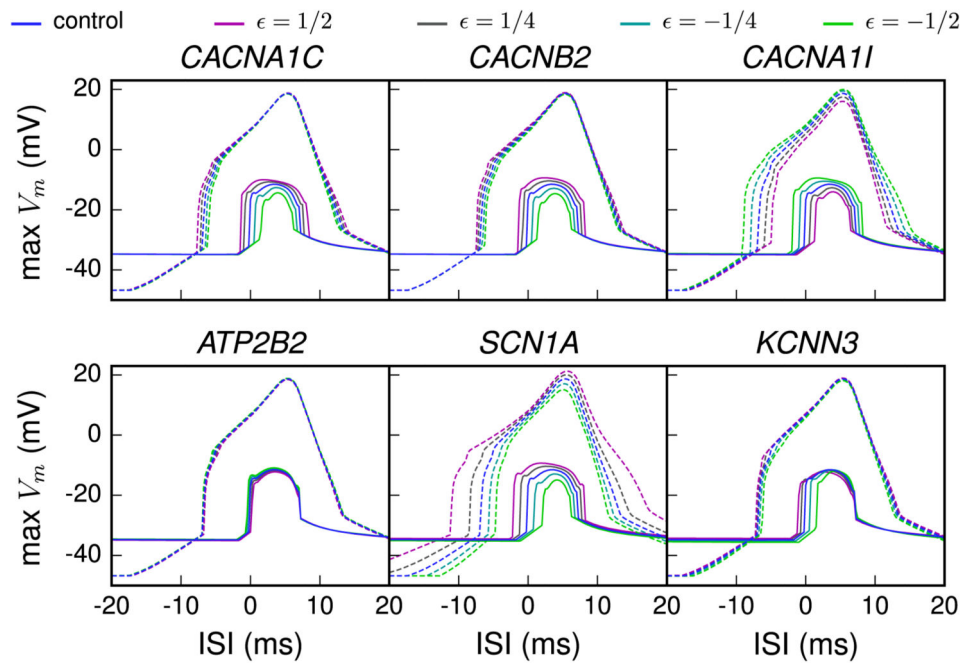
**Figure 3.**

The variant neurons show modulated gain. f-I curves (firing frequency as a function of direct current amplitude) are shown for different scalings of different variants (magenta:  $\epsilon = 1/2$ , dark gray:  $\epsilon = 1/4$ , cyan:  $\epsilon = -1/4$ , green:  $\epsilon = -1/2$ ). Differences in gain are visible for *CACNA1C*, *CACNB2*, *ATP2B2*, and *KCNN3* variants.



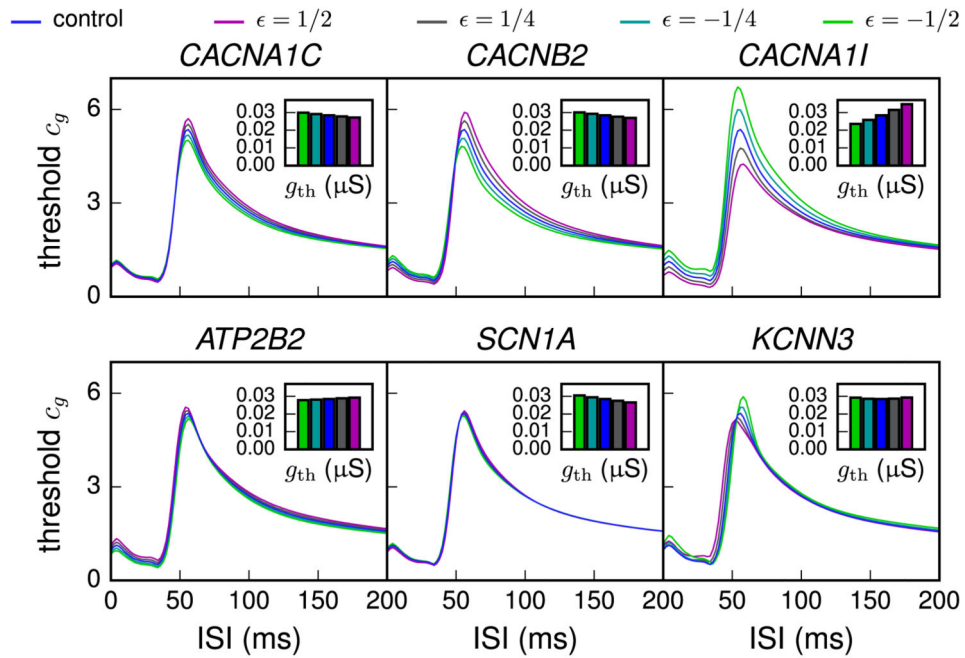
**Figure 4.**

Differences in  $[Ca^{2+}]$  limit cycles of the variants. **(A, B)** The membrane potential **(A)** and calcium ( $Ca^{2+}$ ) concentration **(B)** time courses at soma as a response to a direct current with an amplitude of 1.2 nA. The different colors represent different scalings of a *CACNA1C* variant (43) (scaling as in Figure 2). Inset: The time course shown 3.4 seconds since the beginning of the stimulus. **(C)** The intracellular  $[Ca^{2+}]$  phase plane during steady firing caused by a direct current applied to the soma. The variants and their scalings were chosen as in Figure 2C.



**Figure 5.**

Sensitivity of variant neurons to interstimulus interval (ISI) between somatic and distal apical stimuli during up and down states. The magnitude of the calcium spikes was assessed by measuring the simulated membrane potential near the bifurcation point of the apical dendrite, i.e., at a distance of 620  $\mu\text{m}$  from the soma. The dashed lines show the temporal maximum of the model response membrane potential during a down state and the solid lines during an up state. The x-axis represents the ISI between the somatic and apical stimuli, positive values denoting cases where the apical stimulus was applied after the somatic stimulus. The variants shown are the same as in Figure 2. In the up state paradigm, the neuron was first given a depolarizing square-pulse current of  $.42 \text{ nA} \times 200 \text{ ms}$  at the proximal apical dendrite, 200  $\mu\text{m}$  from the soma. In the middle of this period, a somatic square-pulse current of  $.5 \text{ nA} \times 5 \text{ ms}$  was applied, and after a time defined as the ISI, an alpha-shaped current injection with rise time  $.5 \text{ ms}$ , decay time  $5 \text{ ms}$ , and maximal amplitude of  $.5 \text{ nA}$  was applied. The down state paradigm was otherwise equal, but the long depolarizing current was absent, and to compensate for this, the short somatic square-pulse current had amplitude of  $1.8 \text{ nA}$ .



**Figure 6.**

Threshold conductance factor for inducing a second spike as a function of interstimulus interval (ISI). The neuron was first made to spike by applying 3000 simultaneously activated synaptic conductances that were uniformly distributed along the apical dendrite at a distance (300  $\mu m$ , 1300  $\mu m$ ) from the soma. All synaptic conductances were alpha-shaped with  $\tau = 5$  ms and maximal conductance  $1.1 g_{th}$ , where  $g_{th}$  was the threshold conductance value for inducing a spike in resting state. These values were variant-specific (and scaling-specific), and they are plotted in the inset of each panel. After the specified ISI, the synapses were activated again. The y-axis shows the factor  $c_g$ , chosen such that  $c_g \times g_{th}$  was the threshold conductance of the second stimulus for generating an extra spike. Curves above the control curve (blue) represent strengthened prepulse inhibition, while curves below it represent weakened prepulse inhibition. The effect of the *KCNN3* variant ( $\epsilon = 1/2$ , magenta) was bilateral: for  $ISI \lesssim 50$  ms, the effect was strengthening, whereas for  $ISI \gtrsim 50$  ms, the effect was weakening.

**Table 1**  
**Table of the Genetic Variants Used in This Study**

Gene	References	Type of Variant	Cell Type
<i>CACNA1C</i>	(43)	L429T, L434T, S435T, S435A, S435P	TSA201
<i>CACNA1C</i>	(43)	L779T, I781T, I781P	TSA201
<i>CACNB2</i>	(44)	T11I-mutant	TSA201
<i>CACNB2</i>	(45)	A1B2 vs A1 alone	HEK293
<i>CACNB2</i>	(46)	N1 vs N3 vs N4 vs N5	HEK293
<i>CACNA1D</i>	(47,48)	42A splices transfected	TSA201/HEK293
<i>CACNA1D</i>	(47,48)	43S splices transfected	TSA201/HEK293
<i>CACNA1D</i>	(42,49)	Homozygous knockout	AV node cells / chromaffin cells
<i>CACNA1I</i>	(41)	Alternative splicing of exons 9 and 33	HEK293
<i>ATP2A2</i>	(50)	Heterozygous null mutation	Myocytes
<i>ATP2B2</i>	(51)	Heterozygous knockout	Purkinje cells
<i>ATP2B2</i>	(52)	Homozygous knockout	Purkinje cells
<i>ATP2B2</i>	(53)	G283S-, G293S-mutant	CHO cells
<i>SCN1A</i>	(54)	FHM mutation Q1489K	Cultured neocortical cell
<i>SCN1A</i>	(55)	FHM mutation L1649Q	TSA201
<i>SCN9A</i>	(56)	I228M NaV1.7 variant	HEK293
<i>SCN9A</i>	(57)	A1632E NaV1.7 mutation	HEK293
<i>SCN9A</i>	(58)	L858F NaV1.7 mutation	HEK293
<i>SCN9A</i>	(59)	F1449V NaV1.7 mutation	HEK293
<i>KCNS3</i>	(60)	hKv2.1-(G4S)3-hKv9 fusion inserted	HEK293
<i>KCNB1</i>	(61)	T203K, T203D, S347K, S347D, T203W, S347W	Ltk-
<i>KCNN3</i>	(62)	hSK3_ex4 isoform	tsA
<i>HCN1</i>	(63)	D135W, D135H, D135N mutants	HEK293

For more details, see Table 2.

AV, atrioventricular; CHO, Chinese hamster ovary; FHM, familial hemiplegic migraine; HEK293, human embryonic kidney 293.

**Table 2**  
**List of Variants and Their Threshold Effect Coefficients c**

Gene	Variant Effect on Model Parameters	Threshold Effect
<i>CACNA1C</i>	$V_{\text{offm}^*,\text{CaHVA}}: -25.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -27.0 \text{ mV}$	$c = .094$
<i>CACNA1C</i>	$V_{\text{offm}^*,\text{CaHVA}}: -37.3 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -30.0 \text{ mV}$	$c = .060^a$
<i>CACNB2</i>	$V_{\text{offh}^*,\text{CaHVA}}: -5.2 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .69$	$c = .582$
<i>CACNB2</i>	$\tau_{\text{h}^*,\text{CaHVA}}: \times 1.7$	$c = 2.000$
<i>CACNB2</i>	$V_{\text{offm}^*,\text{CaHVA}}: -4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times .6; \tau_{\text{h}^*,\text{CaHVA}}: \times .6$	$c = .310$
	$V_{\text{offm}^*,\text{CaHVA}}: +4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times .6; \tau_{\text{h}^*,\text{CaHVA}}: \times .6$	$c = .154$
	$V_{\text{offm}^*,\text{CaHVA}}: -4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: +5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times .6; \tau_{\text{h}^*,\text{CaHVA}}: \times .6$	$c = .205$
	$V_{\text{offm}^*,\text{CaHVA}}: +4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: +5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times .6; \tau_{\text{h}^*,\text{CaHVA}}: \times .6$	$c = .675^b$
	$V_{\text{offm}^*,\text{CaHVA}}: -4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times 1.68; \tau_{\text{h}^*,\text{CaHVA}}: \times .6$	$c = .814$
	$V_{\text{offm}^*,\text{CaHVA}}: +4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times 1.68; \tau_{\text{h}^*,\text{CaHVA}}: \times .6$	$c = .057$
	$V_{\text{offm}^*,\text{CaHVA}}: -4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: +5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times 1.68; \tau_{\text{h}^*,\text{CaHVA}}: \times .6$	$c = .517$
	$V_{\text{offm}^*,\text{CaHVA}}: +4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: +5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times 1.68; \tau_{\text{h}^*,\text{CaHVA}}: \times .6$	$c = .078$
	$V_{\text{offm}^*,\text{CaHVA}}: -4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times .6; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.66$	$c = .275$
	$V_{\text{offm}^*,\text{CaHVA}}: +4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times .6; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.66$	$c = .188$
	$V_{\text{offm}^*,\text{CaHVA}}: -4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: +5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times .6; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.66$	$c = .190$
	$V_{\text{offm}^*,\text{CaHVA}}: +4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: +5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times .6; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.66$	$c = 1.687$
	$V_{\text{offm}^*,\text{CaHVA}}: -4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times 1.68; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.66$	$c = .707$
	$V_{\text{offm}^*,\text{CaHVA}}: +4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: -5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times 1.68; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.66$	$c = .061$
	$V_{\text{offm}^*,\text{CaHVA}}: -4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: +5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times 1.68; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.66$	$c = .457^{a,c}$
	$V_{\text{offm}^*,\text{CaHVA}}: +4.9 \text{ mV}; V_{\text{offh}^*,\text{CaHVA}}: +5.1 \text{ mV}; \tau_{\text{m}^*,\text{CaHVA}}: \times 1.68; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.66$	$c = .084$
<i>CACNA1D</i>	$V_{\text{offm}^*,\text{CaHVA}}: -10.9 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times .73; V_{\text{offh}^*,\text{CaHVA}}: -3.0 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .81; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.25$	$c = .118$
	$V_{\text{offm}^*,\text{CaHVA}}: -10.9 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times .73; V_{\text{offh}^*,\text{CaHVA}}: +3.5 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .81; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.25$	$c = .106$
<i>CACNA1D</i>	$V_{\text{offm}^*,\text{CaHVA}}: -10.6 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times .8; V_{\text{offh}^*,\text{CaHVA}}: -5.3 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .66; \tau_{\text{h}^*,\text{CaHVA}}: \times .72$	$c = .114$
	$V_{\text{offm}^*,\text{CaHVA}}: +3.4 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times .8; V_{\text{offh}^*,\text{CaHVA}}: -5.3 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .66; \tau_{\text{h}^*,\text{CaHVA}}: \times .72$	$c = 1.962$
	$V_{\text{offm}^*,\text{CaHVA}}: -10.6 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times 1.12; V_{\text{offh}^*,\text{CaHVA}}: -5.3 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .66; \tau_{\text{h}^*,\text{CaHVA}}: \times .72$	$c = .131$
	$V_{\text{offm}^*,\text{CaHVA}}: +3.4 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times 1.12; V_{\text{offh}^*,\text{CaHVA}}: -5.3 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .66; \tau_{\text{h}^*,\text{CaHVA}}: \times .72$	$c = .601$
	$V_{\text{offm}^*,\text{CaHVA}}: -10.6 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times .8; V_{\text{offh}^*,\text{CaHVA}}: +1.2 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .66; \tau_{\text{h}^*,\text{CaHVA}}: \times .72$	$c = .100$
	$V_{\text{offm}^*,\text{CaHVA}}: +3.4 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times .8; V_{\text{offh}^*,\text{CaHVA}}: +1.2 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .66; \tau_{\text{h}^*,\text{CaHVA}}: \times .72$	$c = .645$
	$V_{\text{offm}^*,\text{CaHVA}}: -10.6 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times 1.12; V_{\text{offh}^*,\text{CaHVA}}: +1.2 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .66; \tau_{\text{h}^*,\text{CaHVA}}: \times .72$	$c = .116$
	$V_{\text{offm}^*,\text{CaHVA}}: +3.4 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times 1.12; V_{\text{offh}^*,\text{CaHVA}}: +1.2 \text{ mV}; V_{\text{sloh}^*,\text{CaHVA}}: \times .66; \tau_{\text{h}^*,\text{CaHVA}}: \times .72$	$c = 1.117$
<i>CACNA1D</i>	$V_{\text{offm}^*,\text{CaHVA}}: +6.6 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times .75; \tau_{\text{h}^*,\text{CaHVA}}: \times .5$	$c = .104$
	$V_{\text{offm}^*,\text{CaHVA}}: +6.6 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times 1.19; \tau_{\text{h}^*,\text{CaHVA}}: \times .5$	$c = .068$
	$V_{\text{offm}^*,\text{CaHVA}}: +6.6 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times .75; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.12$	$c = .115^b$
	$V_{\text{offm}^*,\text{CaHVA}}: +6.6 \text{ mV}; V_{\text{slo}^*,\text{CaHVA}}: \times 1.19; \tau_{\text{h}^*,\text{CaHVA}}: \times 1.12$	$c = .072$

Gene	Variant Effect on Model Parameters	Threshold Effect
<i>CACNA1H</i>	$V_{\text{offma,CaLVA}}: +1.3 \text{ mV}; V_{\text{offha,CaLVA}}: +1.6 \text{ mV}; \tau_{\text{m}^*,\text{CaLVA}}: \times .87; \tau_{\text{h}^*,\text{CaLVA}}: \times .8$	c 2.000
	$V_{\text{offma,CaLVA}}: +1.3 \text{ mV}; V_{\text{offha,CaLVA}}: +1.6 \text{ mV}; \tau_{\text{m}^*,\text{CaLVA}}: \times 1.45; \tau_{\text{h}^*,\text{CaLVA}}: \times .8$	c 2.000 <sup>a,b</sup>
<i>ATP2A2</i>	$\gamma_{\text{CaDynamics}}: \times .6$	c = .093 <sup>b</sup>
<i>ATP2B2</i>	$\tau_{\text{decay,CaDynamics}}: \times 1.97$	c = .218
<i>ATP2B2</i>	$\tau_{\text{decay,CaDynamics}}: \times 1.5; c_{\text{min,CaDynamics}}: \times 1.4$	c = .215 <sup>a</sup>
<i>ATP2B2</i>	$\tau_{\text{decay,CaDynamics}}: \times 4.45$	c = .099
<i>SCN1A</i>	$V_{\text{offin,Nat}}: -3 \text{ mV}; V_{\text{offh,Nat}}: +5 \text{ mV}; V_{\text{sloin,Nat}}: \times 1.15; V_{\text{sloh,Nat}}: \times 1.23$	c = .056 <sup>a</sup>
<i>SCN1A</i>	$V_{\text{offin,Nat}}: +2.8 \text{ mV}; V_{\text{offh,Nat}}: +9.6 \text{ mV}; V_{\text{sloin,Nat}}: \times .984; V_{\text{sloh,Nat}}: \times 1.042$	c = .069
<i>SCN9A</i>	$V_{\text{offh}^*,\text{Nap}}: +6.8 \text{ mV}$	c 2.000
<i>SCN9A</i>	$V_{\text{offh}^*,\text{Nap}}: +3.5 \text{ mV}; V_{\text{sloh,Nap}}: \times .55; V_{\text{offin,Nat}}: -7.1 \text{ mV}; V_{\text{offh,Nat}}: +17.0 \text{ mV}; V_{\text{sloh,Nat}}: \times .69$	c = .026
<i>SCN9A</i>	$V_{\text{offin,Nat}}: -9.1 \text{ mV}; V_{\text{offh,Nat}}: +3.1 \text{ mV}$	c = .043
<i>SCN9A</i>	$V_{\text{offin,Nat}}: 27.6 \text{ mV}; V_{\text{offh,Nat}}: +4.3 \text{ mV}$	c = .043
<i>KCNS3</i>	$\tau_{\text{m}^*,\text{Kp}}: \times 2.0; \tau_{\text{h}^*,\text{Kp}}: \times 2.5; V_{\text{sloh,Kp}}: \times .5$	c 2.000
<i>KCNB1</i>	$V_{\text{offin,Kp}}: +5 \text{ mV}; V_{\text{offh,Kp}}: 13 \text{ mV}; V_{\text{sloin,Kp}}: \times 1.11; V_{\text{sloh,Kp}}: \times .86; \tau_{\text{m}^*,\text{Kp}}: \times .5; \tau_{\text{h}^*,\text{Kp}}: \times .53$	c 2.000
<i>KCNB1</i>	$V_{\text{offin,Kp}}: +1 \text{ mV}; V_{\text{offh,Kp}}: -6 \text{ mV}; V_{\text{sloin,Kp}}: \times 1.22; V_{\text{sloh,Kp}}: \times 1.0; \tau_{\text{m}^*,\text{Kp}}: \times .89; \tau_{\text{h}^*,\text{Kp}}: \times 1.13$	c 2.000
<i>KCNB1</i>	$V_{\text{offin,Kp}}: +6 \text{ mV}; V_{\text{offh,Kp}}: -8 \text{ mV}; V_{\text{sloin,Kp}}: \times 1.33; V_{\text{sloh,Kp}}: \times 1.0; \tau_{\text{m}^*,\text{Kp}}: \times .5; \tau_{\text{h}^*,\text{Kp}}: \times .87$	c 2.000
<i>KCNB1</i>	$V_{\text{offin,Kp}}: -28 \text{ mV}; V_{\text{offh,Kp}}: -27 \text{ mV}; V_{\text{sloin,Kp}}: \times 1.11; V_{\text{sloh,Kp}}: \times .71; \tau_{\text{m}^*,\text{Kp}}: \times 1.13; \tau_{\text{h}^*,\text{Kp}}: \times 2.27$	c 2.000
<i>KCNB1</i>	$V_{\text{offin,Kp}}: +14 \text{ mV}; V_{\text{offh,Kp}}: -21 \text{ mV}; V_{\text{sloin,Kp}}: \times 2.0; V_{\text{sloh,Kp}}: \times 1.0; \tau_{\text{m}^*,\text{Kp}}: \times .39; \tau_{\text{h}^*,\text{Kp}}: \times 1.2$	c 2.000
<i>KCNB1</i>	$V_{\text{offin,Kp}}: -13 \text{ mV}; V_{\text{offh,Kp}}: -13 \text{ mV}; V_{\text{sloin,Kp}}: \times 1.33; V_{\text{sloh,Kp}}: \times .71; \tau_{\text{m}^*,\text{Kp}}: \times .95; \tau_{\text{h}^*,\text{Kp}}: \times 5.13$	c 2.000
<i>KCNN3</i>	$C_{\text{off,SK}}: \times .86; C_{\text{slo,SK}}: \times 1.24$	c = 1.715 <sup>a,b</sup>
<i>HCN1</i>	$V_{\text{offin}^*,\text{Ih}}: -26.5 \text{ mV}; V_{\text{sloin}^*,\text{Ih}}: \times .64$	c = .296

The variants are ordered as in Table 1, but the variants where several combinations of parameter range end points were considered are listed as separate variants (Supplement 1).

CaHVA, high-voltage activated  $\text{Ca}^{2+}$ ; CaLVA, low-voltage activated  $\text{Ca}^{2+}$ ; Ih, nonspecific cation current; Kp, slow inactivating  $\text{K}^+$ ; Nap, persistent  $\text{Na}^+$ ; Nat, fast inactivating  $\text{Na}^+$ ; SK, small-conductance  $\text{Ca}^{2+}$ -activated  $\text{K}^+$ .

<sup>a</sup>The variants were used in Figures 2–6.

<sup>b</sup>The variants were used in Figure S3 in Supplement 1.

<sup>c</sup>The variant was used in Figure 1.