Dampening of hyperexcitability in CA1 pyramidal neurons by polyunsaturated fatty acids acting on voltage-gated ion channels

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Running head: Fatty acid modulation of ion channels
Abstract

A ketogenic diet is an alternative treatment of epilepsy in infants. The diet, rich in fat and low in carbohydrates, elevates the level of polyunsaturated fatty acids (PUFAs) in plasma. These substances have therefore been suggested to contribute to the anticonvulsive effect of the diet. PUFAs modulate the properties of a range of ion channels, including K and Na channels, and it has been hypothesized that these changes may be part of a mechanistic explanation of the ketogenic diet. Using computational modelling, we here study how experimentally observed PUFA-induced changes of ion channel activity affect neuronal excitability in CA1, in particular responses to synaptic input of high synchronicity. The PUFA effects were studied in two pathological models of cellular hyperexcitability associated with epileptogenesis. We found that experimentally derived PUFA modulation of the A-type K (K_A) channel, but not the delayed-rectifier K channel, restored healthy excitability by selectively reducing the response to inputs of high synchronicity. We also found that PUFA modulation of the transient Na channel was effective in this respect if the channel’s steady-state inactivation was selectively affected. Furthermore, PUFA-induced hyperpolarization of the resting membrane potential was an effective approach to prevent hyperexcitability. When the combined effect of PUFA on the K_A channel, the Na channel, and the resting membrane potential, was simulated, a lower concentration of PUFA was needed to restore healthy excitability. We therefore propose that one explanation of the beneficial effect of PUFAs lies in its simultaneous action on a range of ion-channel targets. Furthermore, this work suggests that a pharmacological cocktail acting on the voltage dependence of the Na-channel inactivation, the voltage dependences of K_A channels, and the resting potential can be an effective treatment of epilepsy.
Introduction

Epilepsy is a severe neurological disorder which is characterized by spontaneous recurrent seizures. Many factors have been linked to the etiology, among them ion channels. Voltage-gated ion channels are crucial for generating and regulating neuronal excitability. Their pivotal importance is evidenced by multiple channel mutations inducing hyperexcitability and epilepsy in humans [1,2]. As a rule of thumb, opening of voltage-gated sodium (Na) channels increases excitability while opening of voltage-gated potassium (K) channels reduces excitability. Thus, several gain-of-function mutations in Na channels [1] as well as loss-of-function mutations in both delayed rectifier (K\textsubscript{DR}) and A-type K (K\textsubscript{A}) channels [3-7] are associated with epilepsy.

The strong connection between voltage-gated ion channels and neuronal activity makes ion channels an attractive pharmacological target for anticonvulsive substances. The traditional pharmacological strategy is to reduce excitatory Na currents by targeting the ion conducting pore [8-10]. However, despite the great number of antiepileptics on the market, about 20-30% of patients with epilepsy respond incompletely to drug treatment [11]. Lack of therapeutic effects in many patients in combination with adverse effects [12,13] motivates the search for new antiepileptic drugs, new targets, and new pharmacological mechanisms.

An epileptic seizure has a rich repertoire of events. One pronounced feature during epileptogenesis is highly synchronized neuronal activity [14-17]. Synchronous input is very powerful in activating neurons [18-20], and therefore an enhanced neuronal response can be part of the pathology. In a previous study [21], we showed that highly synchronized activity is suppressed by the K\textsubscript{A} channel in dendrites, which therefore may function as a protective mechanism against hyperexcitability. In epilepsy, the K\textsubscript{A} current may not be strong enough to compensate for the excitability changes due to the pathology. Thus, substances changing the activity of channels involved in suppressing cellular responses to synchronicity may be a powerful way to prevent epileptic seizures.

Polyunsaturated fatty acids (PUFAs) are suggested as important antiepileptic substances in the fat-rich ketogenic diet used as an alternative epilepsy treatment in children [22-25]. The mechanism of the ketogenic diet is largely unknown, but PUFAs directly target a wide range of ion channels, including Na and K channels [e.g. 24,26]. The proposed mechanism for the modification suggests that the partly negatively charged lipophilic molecules target the lipid bilayer close to the positively charged voltage sensor of ion channels and electrostatically activate the channel and open the ion-conducting pore (Fig. 1A) [27-29]. This interaction with the voltage sensor leads to a modified voltage dependence, so that the channel opens at more negative voltages (Fig. 1B). Figure 1C shows the concentration dependence of the PUFA-induced shift of K channel activation (see figure legend for further details). In Na channels PUFAs instead shift the voltage dependence of channel inactivation.

The objective of this study was to investigate PUFAs as reducers of excessive neuronal responses to synchronous input, and thereby as possible anticonvulsive substances. To experimentally test the importance and contribution of each PUFA-induced channel modulation separately is however very difficult as specific antagonists of the modulation are unavailable. The evaluation is further complicated by some effects being predicted to oppose effects of other [24,26]. In this study we therefore turned to a computational model of a pyramidal neuron in hippocampal area CA1 (see Fig. 1D), a brain region commonly involved
in epileptic seizures [17,30]. During a simulation we activated five synaptic inputs contacting five oblique dendrites of the CA1 neuron. In different runs, the synchronicity between the inputs varied from completely simultaneous (i.e. synchronized) to relatively desynchronized. The number of produced action potentials was used as a measure of excitability.

To study hyperexcitability we implemented two cases of ion-channel alterations associated with epileptogenesis. The first case comprised enhanced Na currents [31] and the second case a rundown in the K_A conductance [3,32]. In the two pathological models we subsequently investigated how PUFAs modulate the cellular response by altering the activity of voltage-gated ion channels. We focused on three prominent channels: a Na channel, an A-type K (K_A) channel, and a delayed-rectifier K (K_{DR}) channel. PUFA modulation of the Na channel was implemented as a hyperpolarizing shift of the steady-state inactivation curve as reported in human and rat cardiac Na channels and rat CA1 Na channels [33-35]. This leads to a reduced Na current [35,36]. We implemented PUFA modulation of K_A as a hyperpolarizing shift of the steady-state activation curve [24], leading to an increased K current, and assumed K_{DR} to be similarly affected by PUFAs [37-39] and possibly also by other ketogenic diet agents [40,41] that may act on ion channels such as the ATP-dependent K channel, the h-channel, or the K-conducting M-channel. We therefore also tested the impact of altered resting potential on excitability. In the second part of this study we investigated the effect on the pathological cell models when simultaneously shifting the steady-state activation and inactivation curves equally much or when simultaneously affecting several ion channels.

The goal was to restore a physiological, non-epileptogenic, neuronal response. The studies showed that modifications of the Na channel, the K_A channel, or the resting potential were effective ways to normalize a pathological behaviour. Thus, PUFAs and substances acting through similar mechanisms could be potent anticonvulsive substances by modulating the activity of multiple ion channels and thereby reducing the neural response to synchronous input.
Results

The cellular response to highly synchronized input is suppressed
To study the effects of PUFAs on neuronal excitability, we performed computer simulations using the CA1 pyramidal neuron model by Migliore et al. [42]. The cell model, based on a morphologically reconstructed neuron, includes a transient Na current, a persistent K$_{DR}$ current, a transient K$_A$ current, and a pacemaker h current. CA1 is one of the key regions implicated in temporal lobe epilepsy and the pyramidal neuron is proposed to be the primary neuron type generating fast ripples [15], a highly synchronized ensemble activity associated with seizures. As epilepsy ultimately is a network phenomenon, and because communication between neurons is performed via spikes, measuring spike production is one way of addressing excitability on a single neuron level. In this work, we therefore measured the number of spikes produced and used this as a measure of responsiveness to the input provided.

The timing of arrival of each input was adjusted to obtain stimuli of different synchronicity (Fig. 1D, Fig. 2A) where a larger jitter of the inputs (Fig. 2A right) represents less synchronized input, and zero time difference (Fig. 2A left) corresponds to perfectly synchronized input. Passive biophysics would predict that highly synchronized input should be the most effective input to activate a neuron. However, this is not the case here since highly synchronized input (<2 ms) is suppressed (Fig. 2B-D). This is consistent with our previous results using a single-input site on a medial location of the apical dendrite [21]. Figure 2E shows the spike probability when the input jittering was randomly sampled from a normal distribution rather than sampled uniformly as in Figure 2D. The main difference between the two sampling methods was that the random sampling produced a smoother curve while qualitative characteristics were similar, in particular for the highly synchronized inputs which are our main interest.

Pathological models produce increased cellular response for highly synchronized input
A range of ion channels have been linked to epileptogenesis, particularly Na and K channels. Several Na channel gene mutations causing an increased open probability have been linked to epilepsy [1], and a study on kindling epileptogenesis observed a 22% increase in the peak Na current [31]. Thus, to implement a Na channel pathology, we increased the Na conductance by 22%. This model is referred to as the “increased Na current pathology”. The involvement of the K$_A$ channel in epileptogenesis has been studied in both animal models and in human tissue [3,32,43,44]. Some of these studies show a downregulation of the gene expression or the current of K$_A$ [44]. To implement a K$_A$-channel pathology, we decreased the K$_A$ conductance by 50% [3,32]. This model is referred to as the “decreased K$_A$ current pathology”.

Figure 3A shows the output spike responses of the control model (blue lines) and the two pathological models (red lines) for different degrees of input synchronicity. Both pathological models clearly displayed hyperexcitability which is consistent with our previous study [45]. For inputs of low synchronicity (>6 ms; large jitter values), the decreased K$_A$ current model showed particularly large increases in output excitability. The low-synchronicity input may be regarded as background input and provides a measure of general excitability. More importantly, highly synchronized inputs (<2 ms; small jitter values), with particular interest for epileptogenesis (see Discussion), led to increased responses in both pathological models. Figure 3B shows the spike probability using random sampling of the input jittering and the
result is similar as for the deterministic model in Figure 3A. As random sampling requires a great number of simulations to be representative, we will in the following use the deterministic uniformly sampled distribution.

**Pharmacological correction of pathological models**

The pathological models of the pyramidal neuron in CA1 generate excess spiking when input of different synchronicity is provided (Fig. 3). The main objective of this study was to correct the pathologies by reducing the excess spiking for high-synchronicity levels (<2 ms). To correct the pathologies we simulated the experimental effects of PUFAs on the Na, K$_{DR}$, and K$_A$ channels. We also studied the effect of an altered resting membrane potential. For more information of the modulation of the ion channels and the resting membrane potential see the Models section. The correction strategy is exemplified in Figure 4, illustrating three attempts to correct the increased Na current pathology by shifting the steady-state activation curve of the K$_A$ channel by $-1$, $-2$, or $-3$ mV along the voltage axis. The increased K$_A$ channel activity, caused by the negative shift, reduced the number of spikes. For shifts of $-1$ and $-2$ mV, the reduction in the number of spikes is selective for the highest and lowest levels of synchronicity. A shift of $-3$ mV abolished output activity for all synchronicity levels. Thus, a shift between $-1$ mV and $-2$ mV generates to best correction to the original, healthy neuron.

In the following we will define the pathological model as functionally corrected if zero spikes are generated for completely synchronized input (0 ms) and the maximum number of spikes (15) at synchronicity level 2 ms. Thus, for each correction there exists a therapeutic interval where these criteria are satisfied.

**Correcting the increased Na pathology**

Figure 5 shows the increased Na pathology model corrected by the experimentally described PUFAs modulation of the K$_A$ channel, the Na channel, and the resting membrane potential, respectively. Modulation of the K$_A$ channel by shifting the steady-state activation curve by $-1.6$ mV (Fig. 5A) or the resting potential by $-1.1$ mV (Fig. 5C) restored the spike activity for both high (<2 ms) and low (>6 ms) synchronicity levels, while modulation of the Na channel by shifting the steady-state inactivation curve by $-6.7$ mV (Fig. 5B) suppressed the spike activity for high and low synchronicity levels but also suppressed the activity for some intermediate (5-6 ms) synchronicity levels. PUFAs-induced shifts of the K$_{DR}$ channel steady-state activation curve (data not shown) were unable to correct the pathological model. When the shift was large enough (~27.2 mV) to abolish spikes for high synchronicity levels (0 ms and 1 ms) it was not able to produce 15 spikes at 2-ms synchronicity. A functionally corrected model should be robust in the sense that small alterations of parameter values (here PUFAs-induced shifts of the channels’ voltage dependences or the resting potential) should be tolerated without reintroducing pathological spiking. But, at the same time, the model should not be totally insensitive to changes of a parameter presumed to play a role in correcting the pathology. The models were therefore analyzed further by measuring the parameter interval within which the model was classified as functionally corrected, that is no spikes at 0-ms synchronicity and 15 spikes at 2-ms synchronicity (Fig. 5D). Smaller shifts were needed for the K$_A$ channel ($-0.9$ to $-1.8$ mV) than for the Na channel ($-4.3$ to $-7.2$ mV).

**Correcting the decreased K$_A$ current pathology**

Figure 6A-C shows the best correction of the decreased K$_A$ pathology for PUFAs modulation of the K$_A$ channel, the Na channel, and the resting membrane potential, respectively. PUFAs modulation of the K$_A$ channel perfectly restored normal output activity, while modulation of the Na channel and the resting potential were not able to compensate the output activity at both low- and high-synchronicity at the same time; when functional corrections were
obtained at synchronicity levels of 0-2 ms, the correction was too large at intermediate synchronicity of 4-6 ms. As for the increased Na pathology model, shifting the voltage dependence of the K_{DR} channel did not correct the K_A pathology (data not shown). Figure 6D shows the interval of the modulatory shift within which output was classified as functionally corrected. Also for the decreased K_A pathology model, the smallest shifts required to correct the model were found for K_A and the resting membrane potential.

**Impact of shifting both steady-state activation and inactivation curves simultaneously**

It is known that PUFAs affect both steady-state activation and inactivation of some Na channels [46]. Furthermore, inactivation of K_A is tightly coupled to activation [47], meaning that a shift in the channel’s steady-state activation curve is expected to result in an equally large shift of the steady-state inactivation curve. In this section we consequently extended the study by allowing PUFAs to shift both steady-state activation and inactivation curves by equal amounts for Na and K_A channels. These simulations were not applicable to the K_{DR} channel since it is not capable of fast inactivation. If both the steady-state activation and inactivation curves of the Na channel were shifted, the Na current increased and as a consequence neither of the pathological models could be functionally corrected (data not shown). Conversely, both pathological models could be functionally corrected when the two steady-state curves of the K_A channel were equally shifted (Fig. 7).

**The combined effect of PUFA on several ion channels**

Above, we have studied the effects of modulating one ion channel at a time in two different pathological models. In the next step we explored the three-dimensional space of the combined effect of PUFA modulation of K_A and Na channels, and the resting membrane potential. Figure 8 shows three slices of the explored three-dimensional space for both the increased Na pathology and the decreased K_A pathology models. PUFA was modelled as 1) a shift of the steady-state inactivation curve of the sodium channel, 2) combined shifts of steady-state activation and inactivation curves of the K_A channel, and 3) an alteration of the membrane potential. The solutions of functionally corrected models are found in the blue areas. Interestingly, but not surprisingly, smaller effects on the targets are needed for functional correction if all three effects are included simultaneously.

**Therapeutic intervals**

Based on experimentally obtained dose-response curves described in the Models section (Eq. 2) it is possible to calculate the therapeutic concentration intervals for effects on the steady-state Na inactivation and K_A activation curves. The dose-response curve given for Na inactivation is from CA1 neurons (explored in the present investigation). In contrast, the dose-response curve for the K_A activation is from the Shaker K channel expressed in *Xenopus* oocytes, which is known to underestimate the efficacy of pharmacological substances [48]. Therefore, in the quantitative evaluation below we will give alternative therapeutic intervals assuming a 10 times higher affinity in CA1 neurons than in the *Xenopus* oocytes (referred to as CA1 affinity). The therapeutic interval for PUFA affecting the resting potential is not possible to calculate because of limited experimental data available. Instead, we will estimate the upper limit of the therapeutic interval (see Models section for further details). Table 1 summarizes the therapeutic interval needed to restore the pathological models.

For the increased Na pathology model, the therapeutic interval for PUFA when only K_A activation is affected is 8.2-18 µM (0.82-1.8 µM assuming CA1 affinity, see above). If only Na inactivation is affected, the interval is 1.7-2.8 µM. The therapeutic interval for PUFA acting on the resting potential is <3.9 µM. Moreover, if PUFA acts on both steady-state
inactivation and activation curves for the $K_A$ channel the therapeutic interval is 15-45 μM (1.5-4.5 μM assuming CA1 affinity). Interestingly, this means that the therapeutic interval widens from 8.2-18 μM (factor of 2.2) to 15-45 μM (factor of 3), resulting in a more stable correction even though higher concentrations are required. Furthermore, if PUFA acts simultaneously on the steady-state Na inactivation, steady-state $K_A$ activation and inactivation, and the resting potential, then the therapeutic interval is <1.5 μM (<1.1 μM assuming CA1 affinity).

For the decreased $K_A$ pathology model, the therapeutic interval for PUFA when only $K_A$ activation is affected is 79-110 μM (7.9-11 μM assuming CA1 affinity). If both the steady-state inactivation and the activation curves are modulated it is above 680 μM (>68 μM assuming CA1 affinity). When PUFA only acts on Na inactivation, modulatory effects are not sufficient and no correction is obtained. Furthermore, the therapeutic interval for PUFA acting on the resting potential is ~10 μM. Moreover, if PUFA acts simultaneously on the steady-state Na inactivation, steady-state $K_A$ activation and inactivation, and the resting potential, then the therapeutic interval is <4.8 μM (<3.5 μM assuming CA1 affinity). Thus, strikingly low PUFA concentrations are in several cases adequate to rescue the pathologies. The amount of PUFA required is well within the estimated PUFA concentration range in cerebrospinal fluid during ketogenic diet treatment [22,24]. This highlights the pharmacological potency of shifts in channel voltage-dependencies or the cellular resting potential.
Discussion

In this work we have modelled effects of PUFAs on excitability in a CA1 pyramidal cell. The rationale behind this was to explore a possible mechanism for the ketogenic diet used in the treatment of epilepsy. One important factor contributing to its anticonvulsive effect is believed to be direct ion-channel effects due to elevations in PUFA levels. We have investigated whether experimentally reported PUFA-induced changes in ion channel characteristics can reverse epileptogenic hyperexcitability and thereby normalize neuronal excitability. Altogether our results show that small alterations either from shifting steady-state curves or resting membrane potential is effective in reducing hyperexcitability, particularly for high synchronicity input. Table 1 summarizes the findings of the present investigation.

Selective modulation of the response to synchronous input may minimize adverse effects

Our study has focused on suppression of highly synchronized input as an effective means to reduce excitability related to epileptogenesis. Epileptogenic activity, and in particular fast ripples, is characterized by spiking at high synchronicity levels [14-17]. Such activity will at target cells produce synchronous compound EPSPs characterized by fast rise times, large amplitudes and short duration. In vivo, action potentials most commonly arise from brief dendritic depolarisations of this kind [19,49]. Moreover, synchronous input in vitro is the most effective input to produce spikes [18-20]. We therefore argue that selective reduction of the response to synchronous input might be more beneficial than reduction of spiking activity in general which might be causing the sedative adverse reactions commonly reported for anticonvulsive agents. To us it was an unexpected finding that a shift of the resting potential reduced the pathological excitability without leading to a decrease of the general excitability. Our interpretation is that the selective effect is mediated indirectly by the voltage dependence of K_A, in particular its steady-state inactivation.

The best candidate targets

To rescue the increased Na pathology, there are a number of reasonable target candidates. Acting on Na inactivation, K_A activation and inactivation, or resting membrane potential requires roughly 1-5 μM of PUFA (assuming CA1 affinity). Combined effects on all these targets lower the concentration even further. To rescue the decreased K_A pathology we are limited to effects on the resting potential or combinations including effects on the resting membrane potential, given that we are not able to act separately on the K_A activation and if we believe that the increase in PUFA concentrations during the ketogenic diet does not exceed 30 μM [see ref. 24]. Combined effects on all three targets lower the PUFA concentration required to <4 μM. In contrast to these successful modulations, PUFA modulation of the K_Dr channel was unable to correct any of the two pathologies. Our interpretation is that for an ion channel to contribute significantly to the occurrence of a spike at high synchronicity levels, it has to be fast enough to affect the EPSP before it reaches the spike threshold. K_Dr may nevertheless have effects on the spike threshold and slower depolarisations.

Combined effects on several ion channels as future pharmacological approach?

Combined effects on several ion channels means that beneficial effects are achieved with lower concentrations of the modulator. These simulations highlight the possible usefulness of a cocktail of pharmacological compounds in epilepsy treatment, each with high specificity and affecting different ion channel targets. For instance could M-channel openers like ZnPy
[50], retigabine [51], acrylamide (S)-2 [52] and NH29 [53], which are expected to hyperpolarize the resting potential, potentially be combined with compounds targeting $K_A$ channel voltage dependence or Na channel inactivation for a more effective antiepileptic effect. Our simulations further imply that $K_A$ modulating drugs would act beneficial even when steady-state inactivation is affected similarly as steady-state activation, which would be the most biological relevant modulation of $K_A$ [47]. In contrast, for Na channels a more selective modulation of only the inactivation is preferable. Indeed, the modulation pattern generally seen for PUFAs on Na channels is dominated by the effect on Na channels inactivation.
Models

In this study we investigated the effect of PUFAs on hyperexcitable pathological neurons using computational methods. The reason for using a computational strategy is that the effects on different ion channels can be isolated and evaluated on its own, and that the effect of synaptic input with different levels of synchronicity can be studied in detail.

The cell model

All simulations were performed using the simulator NEURON [54]. The neuron model was based on the work by Migliore et al. [42]. It is a detailed compartmental model of a CA1 pyramidal cell with 474 compartments. The cell model includes a transient Na current, a persistent $K_{DR}$ current, a transient $K_A$ current, and a pacemaker $h$ current. The ion channels were described by Hodgkin-Huxley dynamics. To this published model we added synaptic input of different synchronicity levels. We used five synaptic inputs at five dendritic branches located at distances of 424, 500, 544, 635, and 715 µm from the soma (Figure 1D). Each synapse was stimulated with a frequency of 12 Hz. The simulation was run for 1500 ms, with an initial delay of 300 ms, leading to a maximum of 15 spikes (=action potentials). The postsynaptic conductance, $G$, is described by

$$G = G_{\text{max}} \cdot A (\exp(-t/\tau_2) - \exp(-t/\tau_1)),$$  

where $G_{\text{max}}$ is the maximal conductance of the synapse, $A$ is set to 1.72 so that the synaptic peak conductance equals $G_{\text{max}}$ in the present case. $\tau_1$ is the rise time constant (=0.5 ms), $\tau_2$ is the decay time constant (=3 ms). The synaptic conductances were set to increase linearly with the distance from the soma [55,56] and thus $G_{\text{max}}$ was set to 2.0, 2.3, 2.5, 3.0, and 3.4 nS respectively. To obtain different degrees of synchronicity we used a temporal normal distribution for either 1) a deterministic model, or 2) a stochastic model. The stochastic simulation was repeated 50 times to generate stable mean values. However, the stochastic simulations were relatively time demanding, and because the results were similar for the two models, we used the deterministic model in this study except when otherwise noted.

Pathological models of epilepsy

To generate pathological models, we either increased the Na conductance by 22% [31] (the “Increased Na current pathology model”), or decreased the $K_A$ conductance by 50% [3] (the “Decreased $K_A$ current pathology model”).

PUFA modulation of voltage-gated ion channels

As mentioned in the Introduction, PUFAs modulate voltage-gated ion channels by shifting the steady-state activation and/or inactivation curves in negative direction along the voltage axis (Fig. 1B). To convert shifts, $\Delta V$, to concentrations, $c$, and vice versa we used the general dose-response curve:

$$\Delta V = \Delta V_{\text{max}} / (1 + (K_d/c)^{n_H}),$$  

where $\Delta V_{\text{max}}$ is the maximal shift, $K_d$ is the dissociation constant, and $n_H$ is the Hill coefficient. In Na channels, PUFAs mainly affect the steady-state inactivation curve, leading to a reduced Na current, and in K channels PUFAs mainly affect the steady-state activation curve, leading to an increased K current. To implement the effects on the Na channel we used the data from rat CA1 Na channels [35], where $\Delta V_{\text{max}} = -11.2$ mV, $K_d = 2.1$ µM, and $n_H =$
2.0. To implement the effects on the $K_A$ channel we used the data from the Shaker $K$ channel expressed in *Xenopus* oocytes [24], where $\Delta V_{\text{max}} = -9.6$ mV, $K_d = 79$ µM, and $n_H = 1$. The *Xenopus* oocyte expression system is known to underestimate the efficacy of pharmacological substances [48]. Therefore, in the quantitative evaluation we will also give alternative therapeutic intervals assuming a 10 times higher affinity in CA1 neurons than in the *Xenopus* oocytes (referred to as CA1 affinity). In some of the simulations we shifted steady-state inactivation and activation curves equally much. The dynamics of $K_A$ is different in proximal dendrites compared with distal dendrites [57]. When implementing the PUFA shift the distal and proximal $K_A$ steady-state gates were shifted equally. PUFAs (docosahexaenoic acid, eicosapentaenoic acid, or arachidonic acid) hyperpolarize the resting membrane potential of excitable cells [37-39]. We implemented the modulation of the resting membrane potential as a hyperpolarizing shift of the resting potential of up to $-4$ mV by increasing the conductance of a leak channel. A concentration of about 10 µM PUFA shifts the resting potential in negative direction with about $-4$ mV (shift range from $-1.5$ to $-5$ mV), but no dose-response curve is given. However, assuming a Hill coefficient of 1 (Eq. 2), the maximum concentration ($C_{\text{max}}$) required for a shift in resting potential ($\Delta V_M$) is $C_{\text{max}} = -10 \Delta V_M/4$. For example, the maximum concentration to alter the resting potential with $-1$ mV is 2.5 µM PUFA.

**Functional correction of the pathological models**

We defined the pathological model as functionally corrected if it generated zero spikes for completely synchronized input (0 ms) and the maximum number of spikes (15) at synchronicity level 2 ms. The interval where these two conditions were satisfied is referred to as the therapeutic interval. The interval was estimated by binary search which terminated when the interval was less than 0.1 mV.

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References


Figure legends

Figure 1. Conceptual background to the computational approach. (A) Schematic illustration of PUFA modulation of ion channel voltage dependence. (B) The continuous line schematically illustrates the conductance versus voltage curve for a voltage-gated ion channel. The dashed line illustrates a PUFA-induced negative shift of the voltage dependence. (C) Dose-response curve for the PUFA (docosahexaenoic acid = DHA) induced shift of the voltage dependence of the Shaker channel expressed in Xenopus oocytes. pH = 7.4. Figure modified after Xu et al. (2008) (D) Schematic figure of the CA1 pyramidal cell and the placement of the synaptic inputs used in the simulations. The five medial to distal oblique dendrites were located 424, 500, 544, 635, and 715 µm from the soma. On each oblique dendrite, five synaptic inputs were placed (arrows). The first input cycle was set to have a midpoint at 300 ms, providing an initial period of baseline membrane potential. To represent inputs of different synchronicity levels, input arrival times were jittered using normal distributions of different widths (standard deviations).

Figure 2. Spike-response diagram for the control model for inputs of different synchronicity. (A–C) The left column shows simulation results with synchronicity of 0 ms, middle column of 4 ms, and right column of 8 ms. The first input cycle was set to have a midpoint at 300 ms, providing an initial period of baseline membrane potential. (A) Temporal distribution of synaptic inputs. Each bar corresponds to one synaptic input. The 5 inputs are superimposed in the left panel. (B) The membrane potential (EPSP) at one of the input sites. (C) The membrane potential in the soma. (D) Number of spikes generated for different synchronicity levels of the input. Closed boxes correspond to synchronicity levels and spike counts shown in C. Note the absence of spikes for high synchronicity levels (0 and 1 ms). (E) Spike probability when the input was chosen from a randomly sampled normal distribution. Error bars indicate the standard deviation from 50 simulations.

Figure 3. Pathological ion channel alterations generate hyperexcitable cell models. The figure shows the number of spikes generated when different levels of synchronized input were provided to the cell. The left column shows the increased Na current pathology (22% increase in Na conductance). The right column shows the decreased K_A current pathology (50% decrease in K_A conductance). (A) Deterministic sampling models and (B) random sampling models. Error bars indicate the standard deviation from 50 simulations. The spike counts for the control model (blue) and pathological models (red) are shown. Note that the pathological models do not reduce the cellular response to highly synchronized inputs (0–1 ms). Furthermore the decreased K_A pathology shows hyperexcitability also for lower synchronicity levels (7–24 ms).

Figure 4. Modulation of K_A channels reduces the spike activity in the increased Na current pathology model. Spike activity for different synchronicity levels is reduced when the steady-state activation curve of the K_A channel is shifted in the negative direction along the voltage axis. When the shift was −3 mV (or higher) the cell did not produce any spike at any level of synchronized input and was therefore regarded as pathological.

Figure 5. Effects of PUFA on the increased Na current pathology model. (A–C) The spike activity for the functionally corrected model when (A) K_A is modulated by shifting the steady-state activation curve (−1.6 mV), (B) Na is modulated by shifting the steady-state inactivation curve (−6.7 mV) and (C) the resting membrane potential was shifted (−1.1 mV).
(D) The interval of the shift of the steady-state curves or resting membrane potential where the model was functionally corrected (generated zero spikes for synchronicity level 0 ms and 15 spikes for synchronicity level 2 ms).

**Figure 6. Effect of PUFA on the decreased K\textsubscript{A} current pathology model.** (A-C) The spike activity for the functionally corrected model when (A) K\textsubscript{A} is modulated by shifting the steady-state activation curve (−5.4 mV), (B) Na is modulated by shifting the steady-state inactivation curve (−15.4 mV) and (C) the resting membrane potential was shifted (−4.2 mV). (D) The intervals of the shift of the steady-state curves or resting membrane potential where the model was functionally corrected (generated zero spikes for synchronicity level 0 ms and 15 spikes for synchronicity level 2 ms).

**Figure 7. Effect of PUFA when the steady-state activation and inactivation of K\textsubscript{A} is equally modulated.** (A) Spike activity for the functionally corrected model of the increased Na pathology model. (B) Spike activity for the functionally corrected model of the decreased K\textsubscript{A} pathology model. To functionally correct the two pathologies the steady-state activation and inactivation curves of K\textsubscript{A} were shifted −3.2 mV (increased Na pathology) and −11 mV (decreased K\textsubscript{A} pathology). (C) Shift regions for the K\textsubscript{A} steady-state curves where the pathologies were functionally corrected (generated zero spikes for synchronicity level 0 ms and 15 spikes for synchronicity level 2 ms).

**Figure 8. Regions of functionally corrected models when PUFA is affecting the Na and K\textsubscript{A} channels, and the resting membrane potential simultaneously.** These nomograms indicate the areas (blue) where the pathological models are functionally corrected for different degrees of resting potential adjustments. PUFA was modelled as a shift of the steady-state inactivation curve of the Na channel, shifts of steady-state activation and inactivation curves of the K\textsubscript{A} channel, and a hyperpolarizing shift of the membrane potential. The black lines are solutions for Eq. 2 in the main text for the cases of the K\textsubscript{A} channel and the Na channel. Each point in the line represents the shifts for Na inactivation and K\textsubscript{A} activation respectively for a particular concentration. The dashed line is the corresponding line for CA1-corrected K\textsubscript{A} data.
Table 1. The therapeutic intervals of the concentration needed to functionally correct two pathological models.

<table>
<thead>
<tr>
<th></th>
<th>Increased Na current pathology</th>
<th>Decreased K_A current pathology</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Shift (mV)</td>
<td>Concentrations (µM)</td>
</tr>
<tr>
<td>Na_inact.</td>
<td>4.3-7.2</td>
<td>1.7-2.8</td>
</tr>
<tr>
<td>K_A,act.</td>
<td>0.9-1.8</td>
<td>8.2-18 (0.82-1.8*)</td>
</tr>
<tr>
<td>K_D,act.</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>ΔV_m</td>
<td>0.6-1.5</td>
<td>&lt;3.9</td>
</tr>
<tr>
<td>Na_act. + Na_inact.</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>K_A,act. + K_A,inact.</td>
<td>1.5-3.5</td>
<td>15-45 (1.5-4.5*)</td>
</tr>
<tr>
<td>Na_inact. + K_A,act. + K_A,inact. + ΔV_m</td>
<td>Varies</td>
<td>&lt;1.5 (&lt;1.1*)</td>
</tr>
</tbody>
</table>

* denotes CA1 affinity
Figures

Fig. 1
Fig. 3

A

Increased Na current pathology

B

Decreased $K_A$ current pathology

Control

Pathology
Increased Na current pathology

- 1 mV shift
- 2 mV shift
- 3 mV shift

Control  Pathology
Fig. 5

Increased Na current pathology

A

B

C

D

Number of spikes

Synchronicity (ms)

Shift (mV)

Control Pathology $K_A$ act. Na inact. $\Delta V_m$
Fig. 6

Decreased $K_A$ current pathology

A

B

C

D

Number of spikes vs. Synchronicity (ms)

Shift (mV)

Control  Pathology  $K_A$ act.  Na inact.  $\Delta V_m$
Fig. 7

Increased Na current pathology

Decreased $K_A$ current pathology

Shift (mV)

Inc. Na
Dec. $K_A$

Control  Pathology  $K_A$ act. & inact.
Fig. 8

**Increased Na current pathology**

Shift of the steady–state activation and inactivation curves of $K_A$ (mV)

- $\Delta V_m = 0$ mV
- $\Delta V_m = -0.5$ mV
- $\Delta V_m = -1$ mV
- $\Delta V_m = -2$ mV

**Decreased $K_A$ current pathology**

Shift of the steady–state inactivation curve of the Na channel (mV)

- $\Delta V_m = 0$ mV
- $\Delta V_m = -1$ mV
- $\Delta V_m = -2$ mV

Increased Na current pathology

Decreased $K_A$ current pathology