Hyperpolarization-Activated Cation Current Contributes to Spontaneous Network Activity in Developing Neocortical Cultures

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Introduction

Synchronized spontaneous burst activity of developing cortical networks has been observed in vivo, in isolated neocortical slices and in primary culture [1–3]. The underlying mechanisms, however, are still elusive. It has been proposed that SBA in cortical networks may be due to activity-dependent changes in the balance of excitation and inhibition [1, 4]. Alternatively, neurons could have intrinsic mechanisms that potentially can generate pacemaker activity [5, 6]. Among potential mechanisms, the I_h current provides a depolarizing current upon hyperpolarization, thereby shaping patterns of rhythmic action potential discharge [7]. Moreover, I_h has been shown to contribute to the resting potential, presynaptically facilitates neurotransmitter release and regulates shaping of synaptic potentials during dendritic integration [8–11].

As a hallmark of I_h, a voltage sag upon hyperpolarizing current injections has been observed in various types of neurons [12–14]. The presence of I_h has also been documented for putative GFP-positive interneurons in subcoeruleus and pontine nucleus oralis of GAD67 knock-in mice [15]. The ion channels mediating I_h are the hyperpolarization-activated cyclic nucleotide-gated cation (HCN) channels. Expression of HCN1 and HCN2 were the prevalent isoforms of HCN channels expressed in L-INs and PNs. These results suggest an important contribution of HCN channels to the maintenance of SBA in embryonic corticocortical cultures.
HCN1 channels as early as the first postnatal week critically contributes to giant depolarizing potentials (GDPs) in the developing rat hippocampus [14]. In this preparation, interneuronal HCN channels have been shown to either generate a depolarizing drive or synchronize firing of principal neurons [12, 16] and determine the integrative properties of fast-spiking basket interneurons [17]. The reports of SBA in cortical preparations [1–3] point to a comparable role of Ih in cortical development and prompted us to examine whether Ih current is present in cultured embryonic cortical interneurons and PNs of GAD67-GFP mice, and to characterize its properties. Subsequently, we attempted to answer the question whether Ih current contributes to shaping of SBA, and which isoforms of HCN channels are expressed in cultured cortical neurons. We found that HCN channels are functionally expressed at pre- and postsynaptic locations and may contribute to the shaping and maintenance of SBA in developing cortical cultures.

Materials and Methods

Cell Culture

Cortical cultures were prepared from embryonic brains of GAD67-GFP (Aneo) mice (in the following: GAD67-GFP mice [18]). Details of the culture protocol are described previously [19]. Briefly, 3–5 mouse brains (taken at embryonic day 16) emitting green fluorescence were used for every preparation. The dorsal parts of the telencephalic vesicles were dissociated with trypsin/EDTA and seeded onto the top of a confluent astroglial feeding layer at a density of 1,300 cells/mm² in N2 medium [75% DMEM, 25% Ham’s F12, N2 supplements (Life Technologies, Karlsruhe, Germany), supplemented with 20 ng/ml triiodo-L-thyronine and 0.2 µg/ml corticosterone (Sigma-Aldrich, Germany)]. Half of the medium was then changed once a week. All cultures were maintained at 36°C in a humidified atmosphere (95% air and 5% CO₂). Most characterizations were performed on mature cultures at 12–29 days in vitro (DIV), unless stated otherwise.

Astrogial subcultures were prepared from cortex of P0 to P2 C57/B6 mice, plated at a density of 500 cells/mm² and cultivated to confluent monolayer for one week before plating of neurons.

All experimental procedures were approved by local government (Landesverwaltungsamt Halle, Germany).

Immunocytochemistry

For immunocytochemical double staining of HCN 1–4 channel isoforms and GFP, cell cultures were first fixed for 30 min at 36°C in 4% paraformaldehyde in phosphate buffer (PFA, 0.1 M, pH 7.4). After washing with PBS (0.01 M, pH 7.4), cultures were briefly exposed to ice-cold methanol and then incubated in 0.25% Triton X-100, 10% goat serum, 5% sucrose and 2% BSA in PBS for 15 min at 4°C. Cultures were then incubated overnight with one of the following rabbit polyclonal antibodies: anti-HCN1, anti-HCN2, anti-HCN3 and anti-HCN4 (Alomone Labs, Jerusalem, Israel; 1:400). HCN antibody binding was visualized with a secondary antibody (goat anti-rabbit indocarbocyanine 3 (Cy3), 1:400; Dianova, Hamburg, Germany). In control experiments, each primary antibody was preincubated with control peptide antigen (1:1) for 1 h at room temperature, and no specific immunofluorescence for the preincubated antibodies was detected. After the second fixation, cultures were stained for GFP (monoclonal anti-GFP 1:100; Molecular Probes, Goettingen, Germany). The secondary antibody used was goat anti-mouse indocarbocyanine 2 (Cy2, 1:400; Dianova). Cultures were mounted with fluoromount (Serva, Heidelberg, Germany) for further fluorescence microscopy.

Colocalization of HCN1–4 channels with GFP was analysed in randomly chosen fields from three to seven coverslips from 2–4 preparations. Images were taken on an upright epifluorescence microscope equipped with a CCD camera (Axioptoph; Zeiss, Oberkochen, Germany). At each position, one image was taken with a filter set optimized for Cy2 and the second image was taken with a filter set optimized for Cy3 and the third was a DIC image. Images were then superimposed in MetaMorph software (version 6.01; Universal Imaging Corp., West Chester, Pa., USA). No contrast or brightness correction was used.

Western Blot

Protein was extracted from cultured neurons according to standard procedures using RIPA-Lysis-Buffer [50 mM Tris, pH 8.0, 150 mM NaCl, 1% Igepal, 0.5% deoxycholate, 0.1% SDS, 10% glycerol, 1 mM PMSF, 4% protease inhibitor cocktail (Complete, Roche)]. Protein concentration was determined by the bicinchoninic acid assay (BCA, Pierce, Thermo Scientific). Equal amounts of protein homogenate were separated on SDS-PAGE (10–20 µg per lane) and transferred to nitrocellulose membranes (Optitran BA-S; Schleicher & Schuell). Membranes were blocked with 5% non-fat dry milk, 1% normal goat serum in PBS-Tween (Optitran BA-S; Schleicher & Schuell). Membranes were blocked with 5% non-fat dry milk, 1% normal goat serum in PBS-Tween 20 (0.1%) for 1 h and incubated in primary antibody [HCN1 1:200, HCN2 1:300 (Alomone Labs), GAPDH 1 µg/ml (Covance, Berkeley, Calif., USA) or β-actin 0.5 µl/ml (Sigma)], dissolved in 1% non-fat dry milk, 1% normal goat serum in PBS-Tween 20 (0.1%). Blots were then washed 3 × 10 min in PBS-Tween 20 (0.1%) and placed in HRP-conjugated anti-rabbit secondary antibody (1:2,000 for HCN1 and HCN2; DAKOcytromab, Hamburg, Germany) or HRP-conjugated anti-mouse antibody (1:10,000 for β-actin and GAPDH; DAKOcytromab). After the final washes (three times in PBST), the immunoreactive bands were visualized using Super-Fluo West Dura (Thermo Scientific) reagents and captured by Gene Gnome, using Gene Snap Software (Syngene). Digital images were quantified using MetaMorph software. To determine both HCN1 and HCN2 amounts in the same gels, the nitrocellulose blots were stripped (Restore™ Western Blot Stripping Buffer; Thermo Scientific), after images were taken from the first reaction (either HCN1 or HCN2), re-probed for the second antibody, and quantified again as described above. Optical densities for bands were determined relative to baseline values immediately above or below the bands in the same lanes. The HCN protein bands were normalized to the actin bands, which served as controls for loading equivalence. For display purposes, the normalized levels were then expressed as a fraction of the mean values of all lanes in single gels. Data were taken from gels of 4 independent developmental series, each derived from one preparation. Whole protein extracts from 5 sister cultures were pooled for each age and experiment. Statistical significance of differences
among different ages was tested with one-way ANOVA or ANOVA on Ranks (Sigma Stat, version 3.5; Systat Software, Inc., Chicago, Ill., USA).

**Electrophysiology**

An acrylic insert was glued to the culture dish with silicon resulting in a recording chamber with a volume of 1–1.5 ml. The chamber was mounted on the stage of an inverted microscope and constantly perfused with ACSF at 1.5–2 ml/min. All recordings were performed at 27–29°C. Whole-cell current-clamp and voltage-clamp recordings were obtained using a patch-clamp amplifier (EPC-7; HEKA Elektronik, Lambrecht/Pfalz, Germany). Patch pipettes were pulled from borosilicate glass (GC150TF-10; Harvard Apparatus Ltd., London, UK) with tip resistances of 2.5–4 MΩ. Characterization of basic properties of I\textsubscript{h} and the effects of I\textsubscript{h} blockers were performed with the following pipette solution (in mM): 95 potassium gluconate, 20 potassium citrate, 10 NaCl, 1 MgCl\textsubscript{2}, 0.5 CaCl\textsubscript{2}, 10 HEPES, 3 potassium BAPTA, 3 Mg-ATP, 0.5 Na-GTP. The resting membrane potential was measured directly after establishment of the whole-cell configuration, and then sets of hyper- and depolarizing currents were applied from a potential of −60 mV. The resting membrane potential, input resistance, rebound depolarization and sag amplitude were measured in the presence of antagonists of glutamate receptors 6-cyano-7-nitroquinoxaline-2,3-dione disodium salt (CNQX, 5 μM) and D-2-amino-5-phosphonopentanoic acid (APV, 12.5 μM; Tocris, Bristol, UK) and antagonist of GABA\textsubscript{A} receptors (10 μM bicusculine methiodide (BML), Tocris) to block SBA.

Spontaneous and miniature PSCs were recorded with a pipette solution containing (in mM): 117 caesium gluconate, 13 CsCl, 1 MgCl\textsubscript{2}, 0.07 CaCl\textsubscript{2}, 10 HEPES, 0.1 EGTA, 0.5 Na-GTP. Quantitative analysis of I\textsubscript{h} was performed by digital subtraction of traces in the absence and presence of 30 μM ZD7288 (Tocris) or 1 mM CsCl (Cs\textsuperscript{+}). All measurements were corrected for a liquid junction potential of 10 mV. The fully activated current versus voltage (I–V) relationship was obtained by a test pulse to −120 mV (duration 1,200 ms) followed by steps to different test potentials from −60 to −110 mV in 5-mV increments. The recorded tail current amplitudes, measured 50 ms after the beginning of test pulse, were plotted against each test potential to construct the fully activated I–V. The reversal potential was extrapolated by linear regression. The steady-state activation curve of I\textsubscript{h} was measured by stepping from a holding potential of −50 to −130 mV in 10-mV increments for 800–2,200 ms followed by a repolarizing step to −60 mV (fig. 2b). To increase the stability of whole-cell recordings with increasing hyperpolarization, pulse duration was decreased in each step by 200 ms. Steady-state activation of I\textsubscript{h} was constructed by normalizing the tail current amplitudes, 50 ms after onset of tail currents, using the following equation p(O) = (I – I\textsubscript{min})/(I\textsubscript{max} – I\textsubscript{min}), with I\textsubscript{max} – current amplitude for the voltage step at −130 mV. The resulting data were fitted by a Boltzmann function of the form:

\[ p(O) = \frac{[1 + \exp((V_m – V_{1/2})/\kappa)]^{-1}} \]

from which the values of half-maximal activation (V\textsubscript{1/2}) and the slope factor (κ) were obtained.

GABAergic PSCs were recorded at a holding potential of 0 mV as outwardly directed currents in the presence of glutamate receptor antagonists APV (12.5 μM) and CNQX (5 μM). Miniature GABAergic postsynaptic currents (mGABAPSCs) were recorded in the presence of 1 μM TTX (Tocris). Signals were identified as GABAergic currents by block with BM1 (10 μM) at the end of some experiments (n = 3). Glutamatergic sPSCs were recorded at a holding potential of −60 mV as inwardly directed currents. All drugs were bath-applied. Detection of synaptic currents with subsequent analysis was performed using MiniAnalysis software (version 6.0.3; Synaptosoft, Inc., Decatur, Ga., USA).

Data were filtered at 3 kHz and sampled at 6.3 kHz in current clamp experiments and at 3.2–5 kHz in voltage clamp experiments using Pulse software (v8.77; HEKA Elektronik). Effects of ZD7288 and Cs\textsuperscript{+} were measured 5–10 min after application of the substance.

All data are expressed as means ± SEM. Statistical significance of differences before and after application of ZD7288 or Cs\textsuperscript{+} was tested with the Wilcoxon signed-rank test (Graph Pad Prism 3.02; GraphPad Software Inc., Calif., USA) or with the one-sample t test if the distribution of the data were Gaussian.

**Calcium Imaging**

The spontaneous network activity of 13- to 16-day-old mouse cultures was recorded by using Fluo-3 Calcium Indicator ( Molecular Probes, Invitrogen). Each culture was incubated with 5 μM Fluo-3 for 1 h followed by several washes with Hepes-buffered Ringer solution (in mM: 140 NaCl, 5 KCl, 1.5 CaCl\textsubscript{2}, 0.75 MgCl\textsubscript{2}, 1.25 NaH\textsubscript{2}PO\textsubscript{4}, 20 D-glucose, 15 Hepes/NaOH, pH 7.4). After 30 min to allow deesterification of the dye, sequences of fluorescence images were recorded at 1 Hz during 4-min sessions on an inverted microscope (Axiovert; Zeiss, Oberkochen, Germany) equipped with a cooled CCD camera (CoolSNAP ES; Roper Scientific, Germany). The recorded fields (3–5 for each culture dish) were chosen randomly and marked with a diamond tool for repeated measurements. A differential interference contrast (DIC) image of each field was acquired before imaging for later identification of cells. Images were processed with MetaMorph software (version 7.0; Molecular Devices, Sunnyvale, Calif., USA). A change in [Ca\textsuperscript{2+}]\textsuperscript{−}, was considered significant when the absolute difference of grey values exceeded five times the standard deviation of background noise measured in cell-free areas [20]. Fluorescence data are expressed as ΔF/F\textsubscript{0} (background-corrected increase in fluorescence divided by resting fluorescence). Attendance is used throughout in terms of participating cells in a given frame as percent of all active cells. The analysis of the calcium activity was performed with custom-made software written in MATLAB (version 7.5; MathWorks, Natick, Mass., USA) and Excel (Microsoft Corp., Richmond, Wash., USA). Statistical tests were performed using SigmaStat (version 2.03, SPSS Inc., Chicago, Ill., USA).

**Results**

**Effects of ZD7288 and Cs\textsuperscript{+} on Membrane Properties of Cultured Cortical Neurons**

During injection of hyperpolarizing current pulses (−80 to −150 pA), a distinct voltage sag and rebound depolarization (RD) were observed in all recorded L-INs (n = 6; fig. 1a) and PNs (n = 11; fig. 1b). ZD7288 (30 μM) and D-2-amino-5-phosphonopentanoic acid (APV, 12.5 μM) and CNQX (5 μM). Miniature GABAergic postsynaptic currents (mGABAPSCs) were recorded

\[ p(O) = \frac{[1 + \exp{(V_m – V_{1/2})/\kappa}]} \]
and the rebound depolarization by ZD7288. This confirms recent observations that HCN channels as a conductor of I_h current are widely expressed in acute slices of mouse cerebral cortex at 12–29 DIV.

Another antagonist of HCN channels, Cs^+ (1 mM), showed similar effects, abolishing the voltage sag and the rebound depolarization in 5 of 6 tested L-INs and in 3 of 3 tested PNs. ZD7288 led to a distinct hyperpolarization of the L-INs by 6.8 \pm 1.6 mV (n = 6, p < 0.05) and 4.3 \pm 1.1 mV by PNs (n = 9, p < 0.05; fig. 1c). Correlated with the marked hyperpolarization induced by ZD7288, the somatic input resistance (R_i), measured at the end of a hyperpolarizing current pulse, increased to 210.2 \pm 41.7% in L-INs (n = 5, p < 0.05) and to 140.8 \pm 17.4% in PNs (n = 7, p < 0.05; fig. 1d). Effects of ZD7288 and Cs^+ were observed already 3–5 min after the onset of application. This confirms recent observations [13] that HCN channels as a conductor of I_h current are widely expressed in acute slices of mouse cerebral cortex at 12–29 DIV.

**Kinetic Properties of I_h Current in Cultured Cortical Neurons**

Using the voltage-clamp mode, we next examined the reversal potential and half-maximal activation of I_h currents in L-INs and PNs. I_h current was isolated by subtracting current traces in the presence of 1 mM Cs^+ from those under control conditions. To determine the reversal potential, I_h current was activated by a hyperpolarizing prepulse to –120 mV from a holding potential of –50 mV and measured 50 ms after the beginning of the test pulse to potentials between –110 and –60 mV for L-INs or –120 to –10 mV for PNs, respectively (fig. 2a). Instantaneous currents were plotted against the test potential. The reversal potential (E_{rev}) was analyzed by estimating the point of intersection of a linear regression line with the voltage axis. E_{rev} amounted to –48.4 \pm 2.1 mV for L-INs (n = 15; fig. 2c) and –45.7 \pm 2.6 mV for PNs (n = 5; fig. 2e).

To determine the steady-state activation curve of I_h, tail currents were measured 50 ms after repolarizing the membrane to –60 mV following voltage steps to potentials between –50 and –130 mV (fig. 2b). To minimize the time used to clamp the membrane potential at very negative potentials, we decreased step duration with increasing hyperpolarization. Fitting the data with a Boltzmann function revealed half-maximal activation of I_h in L-INs of –94.2 \pm 0.9 mV (n = 15; fig. 2d) and –89.5 \pm 0.9 mV in PNs (n = 5; fig. 2f).

**Blockade of I_h Impairs SBA in Cortical Cultures**

As has been previously shown, I_h contributes to the occurrence of developmentally relevant spontaneous activity in the immature cortex in vitro [13]. This prompted us to study the functional role of I_h in generation and maintenance of SBA in mouse cortical cultures. Spontaneous bursts were recorded under current-clamp conditions from PNs in cortical cultures as early as 5–7 DIV (data not shown). The effects of ZD7288 and Cs^+ on network activity were examined in mature cultures, characterized by a frequency of spontaneous bursts of 0.16 \pm 0.05 Hz (fig. 3a, c; online supplementary fig. 1, available at www.karger.com/doi/10.1159/000330813). ZD7288 at a concentration of 1 \mu M led to a significant decrease (p < 0.05) of burst frequency to 0.09 \pm 0.04 Hz, 5 min after application (n = 6; fig. 3b, c). Increasing concentrations of ZD7288 further reduced the frequency of spontaneous bursts in all recorded PNs (10 \mu M ZD7288: 0.05 \pm 0.02 Hz; 30 \mu M ZD7288: 0.04 \pm 0.02 Hz; n = 6; fig. 3c, d). To rule out that ZD7288 exerts its effects on burst frequency by directly blocking AMPA-type glutamate receptors [21], we repeated this experiment with 1 mM Cs^+. Shortly after applica-
tion, Cs⁺ showed similar effectiveness in reducing burst frequency to 23.2 ± 4.0% (n = 4, p < 0.05; online suppl. fig. 1b). Prolonged exposure to 1 mM Cs⁺, however, disrupted oscillatory activity so that it became difficult to quantify a characteristic oscillation frequency (online suppl. fig. 1c).

Additionally, we examined the effect of ZD7288 on spontaneous glutamatergic postsynaptic currents (sGluPSCs) recorded in five projection neurons (PNs). At the concentration used (30 μM), ZD7288 did not significantly (p > 0.05) affect the frequency or the amplitude of sGluPSCs. Mean frequency amounted to 85.7 ± 8.0% (n = 5) and mean amplitude to 107.6 ± 5.9% (n = 5) of control values (fig. 3e), confirming the absence of a direct effect of 30 μM ZD7288 on glutamatergic transmission.

Spontaneous network activity was also investigated by means of calcium imaging (fig. 4). The frequency of spontaneous calcium transients and the proportion of cells actively participating in a single transient were calculated before and after acute application of 1–30 μM ZD7288 or 1–2 mM Cs⁺. Overall, the cultures, which were examined between 13 and 16 DIV (n = 12, from two experiments), showed spontaneous transients with a frequency range between 4 and 12.5 transients/min (8.28 ± 0.38 transients/min; fig. 4a–d). Between 64 and 96% of all active neurons participated in single synchronous events (attendance: 82.2 ± 2.3%).

ZD7288 decreased spontaneous network activity in a dose-dependent manner (fig. 4a–d), but did not alter attendance during synchronous events (p = 0.4, one-way ANOVA; fig. 4g, grey columns). The number of active neurons did not change with 1 μM ZD7288 (p = 0.261) or 10 μM ZD7288 (p = 0.075) but decreased with higher concentrations of ZD7288 (30 μM: p < 0.001; fig. 4g, black columns). A total of 1,766 active neurons were examined in 12 cultures.

Prolonged treatment with Cs⁺ (>30 min) dramatically reduced or completely blocked all calcium transients. Only results of recordings made within 10–30 min after drug application were considered here. Overall, after Cs⁺ application the frequency was significantly decreased (6 cultures of 14–16 DIV from 2 experiments). However, the frequency of transients in single fields of 2 cultures was either increased or decreased. The increase in frequency was mostly due to the appearance of intra-burst transients of low-amplitude riding on top of the prolonged calcium transients. Cs⁺ significantly decreased the attendance of neurons in single transients (fig. 4h, grey columns). The overall number of active cells was not altered.

Fig. 2. Basic characteristics of Iₕ in L-INs and PNs. a Protocol (upper panel) and traces of membrane currents (lower panel) activated by a hyperpolarizing prepulse to –120 mV, followed by a test pulse to potentials between –60 and –110 for a representative L-IN (DIV 13). b Protocol (upper panel) and traces of Cs⁺-sensitive currents (lower panel) activated by a hyperpolarizing test pulse to potentials between –60 and –130 mV and measured as tail currents by repolarization of the membrane to a fixed potential of –60 mV in a DIV 13 L-IN. c Instantaneous current-voltage relationship (I–V) obtained by plotting the current 50 ms after the onset of the test pulse against test pulse potential. Lines are obtained by linear regression. The extrapolated reversal potential was –48.4 ± 2.1 mV (n = 15). d Iₕ channel activation curve of L-INs. Iₕ amplitude was measured 50 ms after stepping to a constant potential (–60 mV) and normalized to the value at –130 mV. Curves are fitted by a Boltzmann function (P(O) = [1 + exp(Vm – V 1/2 )/κ])⁻¹). Half-maximal activation was –94.2 ± 0.9 mV (n = 15). e Instantaneous current-voltage relationship (I–V) of PNs. The average reversal potential amounted to –45.7 ± 2.6 mV (n = 5). f Iₕ channel activation curve of PNs. Half-maximal activation was –89.5 ± 0.9 mV (n = 5).
after addition of 1 mM Cs⁺, but was decreased after 2-mM Cs⁺ treatment (fig. 4h, black columns).

GAD-GFP neurons’ spontaneous network activity mirrored the spontaneous activity of non-GABAergic neurons (fig. 4a–c, e inset). Similarly, the blockade of I_h reduced activity frequency of this neuronal subpopulation in a dose-dependent manner (fig. 4a–c, e inset). The number of active GAD-GFP neurons or their attendance in single events was not significantly altered after treatment with ZD7288 (p = 0.180, ANOVA on ranks, p = 0.892, one-way ANOVA, respectively).

Subcellular Localization
To define the subcellular localization of I_h, we recorded miniature and spontaneous GABAergic postsynaptic currents from PNs, the major target cells of L-INs in mouse cortical cultures. Miniature GABAPSCs (mGABAPSCs) were recorded in a total of 9 PNs by holding PNs at 0 mV, in the presence of 1 μM TTX, 12.5 μM APV and 5 μM CNQX (fig. 5a). Under control conditions, the mean frequency of mGABAPSCs was 0.37 ± 0.2 Hz (n = 9) and the mean amplitude was 21.9 ± 1.5 mV (n = 9). ZD7288 (30 μM) reduced the frequency to 81.4 ± 3.6% of the control value (p < 0.05; n = 4), whereas the amplitude was not significantly affected (93.9 ± 3.0% of the control value, p > 0.05; fig. 5b). Cs⁺ (1 mM) yielded similar effects on the frequency of mGABAPSCs (53.9 ± 15.9% of the control value; n = 5; p < 0.05), confirming the presynaptic localization of HCN channels at the L-INs. Interestingly, in the presence of Cs⁺, the amplitude of mGABAPSCs was also reduced to 80.0 ± 6.5% (n = 5; p < 0.05; fig. 5c), suggesting a potential localization of HCN channels also at postsynaptic PNs.

To test whether the effect of I_h blockers was dependent on synaptic recruitment of L-INs into network activity, we examined spontaneous GABAPSCs (sGABAPSCs) in PNs. Both compounds influenced neither the frequency nor the amplitude of sGABAPSCs (fig. 5d, e). After application of ZD7288, the mean normalized frequency of sGABAPSCs was 120.1 ± 15.2% and the amplitude 90.3 ± 2.9% of control (n = 4; fig. 5d). In the presence of Cs⁺, the frequency amounted to 108.4 ± 15.8% and the amplitude to 107.2 ± 4.8% of control values (n = 5; fig. 5e).
**Fig. 4.** Decreased spontaneous network activity after $I_h$ blockade. 

- **a–c** Calcium transients recorded in 2-week-old networks (a, b DIV 15, c DIV 14) in control Ringer (left side) and after bath application of 1 μM ZD7288 or 1 mM CsCl. Neurons in b were GAD-GFP interneurons in the same network as unlabeled neurons shown in a. 

- **d** Summary graph shows the effect of increasing concentrations of ZD7288 on the activity of 13- to 16-day-old networks (11 cultures from 2 experiments). 

- **e** The network activity is significantly decreased (significant changes compared with the anterior bar: ** p < 0.001, * p = 0.021). Inset graph shows that the activity of GABAergic neurons decreased in the same way by addition of ZD7288 (control vs. 1 μM ZD7288, ** p < 0.001; 1 μM ZD vs. 10 μM ZD, * p = 0.042). 

- **f** Network activity decreases significantly with addition of 1–2 mM Cs$. The frequency of network events is not further reduced by 2 mM compared with 1 mM Cs$ (* p = 0.021). 

- **g, h** Bar graphs show the relative number of active neurons (black bars) and the attendance of active neurons in single events (gray bars) in recordings under $I_h$ blockade with ZD7288 (g) or Cs$ (h). Contrary to ZD7288, which does not alter attendance to single bursts, Cs$ treatment significantly decreases the number of active neurons participating in single bursts (control vs. 1 mM Cs$, p < 0.015; control vs. 2 mM Cs$, p = 0.004). The total number of active neurons decreases in both cases at the higher concentrations (30 μM ZD7288, p < 0.001; 2 mM Cs$, p = 0.003). Data in d–h are normalized to the mean value of control recordings.
Thus, our results are consistent with a presynaptic localization of HCN channels on the membrane of L-INs and an additional postsynaptic localization in PNs, which both may contribute to the maintenance of SBA in mouse cortical cultures.

Developmental Expression of HCN Subunits

To examine the development of HCN expression and putative subunit composition of $I_h$ channels in L-INs and PNs, we analyzed the expression of HCN1–4 isoforms by means of immunocytochemistry (fig. 6) and Western blot (fig. 7). At 6–7 DIV, HCN2 and HCN1 immunoreactivity is conspicuous in cell bodies but also in puncta in neuropile and over dendritic surfaces. Low expression levels of HCN4 and HCN3 were also detected, mainly on cell bodies. Interestingly, although HCN4 was also expressed in interneurons, most labeled somata were non-GABAergic neurons. At older ages (14–28 DIV), specific immunoreactivity was observed to HCN1 and HCN2, but not to HCN3 and HCN4. Dual labeling of GAD-GFP and HCN1 or HCN2 revealed overlapping expression predominantly at somatic (fig. 6a–c), but also at dendritic localizations of L-INs at 21 DIV (fig. 6d, e). The colocalization of HCN1 or HCN2 (red puncta) with green dendrites of L-INs is shown at higher magnification in figure 6d, e. Note that GFP-negative neurons (PNs) also show labeled puncta along somatodendritic surface (fig. 6f–h, arrow).

Western blots confirmed the expression of HCN1 and HCN2 subunits in cortical mouse cultures, but failed to detect HCN3 or HCN4. Quantitative Western blot analyses showed that HCN1 and HCN2 isoforms undergo developmental changes in expression levels. A significant increase in HCN1 and HCN2 protein levels was detected during the first week in vitro (fig. 7a–c). HCN1 and HCN2 protein levels were low at the onset of culturing time, increased significantly between 0 and 7 DIV and showed little change during the second and third weeks in vitro (7–28 DIV) (fig. 7).

Discussion

In the present study, all L-INs and PNs showed a prominent voltage sag during hyperpolarization, indicating activation of the HCN channels. The slow deactivation kinetics of these channels create a slow depolarization of the membrane [7] which may reach a burst-generating voltage window. Both pharmacological blockade of HCN channels by ZD7288 and Cs$^+$ caused a strong reduction of the frequency of SBA. At the relatively low concentrations used in this study, ZD7288 (1–30 µM) and Cs$^+$ (1–2 mM) had no detectable effects on conductances other than $I_h$, which have been reported for higher concentrations and long application times [21, 22]. Caesium is frequently referred to as a blocker of $I_h$ at low millimolar concentrations [23]. Extracellular Cs$^+$ has been used as a selective blocker of $I_h$ up to concentrations of 5 mM in pyloric neurons of the lobster stomato-
gastric ganglion, where it does not block other currents at these concentrations [24]. More recently, extracellular Cs+ concentrations up to 1 mM have been shown to decelerate thalamic oscillations due to blockade of I_h in rat thalamic slices [25]. In slice preparations of the basolateral amygdala of rats, Cs+ (1 mM) was used to selectively block modulation of I_h by neuropeptides [26]. This does not rule out that other ion channels are affected by extracellular Cs+, as for instance the two-pore domain K+ channel TASK-1 [27], which contributes to the control of activity modes in thalamocortical neurons [28]. The low abundance of TASK-1 in neocortical neurons [29], however, argues against a major contribution of this current to spontaneous burst activity of cortical cultures at the low concentrations (1–2 mM) of extracellular Cs+ used in the present study.

Fig. 6. HCN subunit expression in cortical cultures. a–c Dual labelling of GFP (green) and HCN2 (red) reveals overlapping expression in the soma as well as in the dendrites of L-INs at 21 DIV. d Enlarged image of the boxed region in c. e Merged image of a dual labelling of GFP (green) and HCN1 (red) showing dendritic localization in L-INs at 21 DIV. f–h Dual labelling of GFP (green) and HCN2 (red) reveals overlapping expression in the soma (arrow) as well as in the dendrites of non-GABAergic neurons at 7 DIV. Scale bar for a–c, f–h shown in f = 10 μm; scale bar for d and e shown in d = 5 μm. Colors refer to the online version only.
Developmental Regulation and Subcellular Localization of HCN Subunits

HCN channels are abundant in cultured cortical networks. Quantitative Western blot analysis of HCN subunit expression showed an age-dependent upregulation of HCN1 and HCN2 isoforms in mouse cortical cultures. This result suggests that HCN1 and HCN2 are the major subunits in mature networks, which corresponds very well to the HCN channel pattern expressed by hippocampal fast-spiking basket cells obtained by single-cell RT-qPCR analysis [17]. In other neuronal populations of the hippocampal CA3 region also the HCN4 subunit is expressed, suggesting that there is a cell-type-selective and subunit-specific expression pattern of HCN channels [30].

Immunocytochemistry showed that HCN channels are readily detected in the somatodendritic surface of both GABAergic L-INs and PNs. Specific localization of HCN channels to GABAergic basket neurons has also been found in the medial septum [31], hippocampus [17] and cerebellum [32]. In these interneurons, three of the four known HCN channel isoforms (HCN1, 2, 4) seem to be expressed preferentially at somatic and axonal localizations. Our results suggest that HCN channels are also expressed in presynaptic terminals of L-INs, because ZD7288 and Cs⁺ reduced the frequency of mGABAPSCs recorded from PNs. These data confirm results from hippocampal dentate gyrus [17], where HCN channels have been found in somatodendritic and axonal compartments of fast-spiking basket-cell interneurons.

Recently, HCN1 channels were demonstrated at the active zone of mature glutamatergic synaptic terminals of cortical projection neurons, establishing contact with entorhinal cortical layer III cells, where they are involved in regulation of Ca²⁺ influx via presynaptic T-type (CaV3.2) Ca²⁺ channels [33].

What Is the Contribution of Iₜ to Early Network Activity?

Instead of Iₜ, a persistent Na⁺ current (I_Nap) has been found to account for the slow depolarization following a firing-evoked afterhyperpolarization in CA3 pyramidal cells of the immature hippocampus [34]. Moreover, recurrent burst activity is promoted by a tonically depolarizing action of GABA in immature CA3 pyramidal neurons [35]. In the present study, ZD7288 concentrations as low as 1 µM significantly reduced the SBA frequency. This correlates well with the expected increase in interburst intervals by blocking Iₜ. At this concentration, unspecific effects of ZD7288 on SBA frequency mediated by inhibition of T-type calcium currents [22] can be largely ruled out. Moreover, Iₜ-independent depression of synaptic transmission, as reported for hippocampal mossy fibers [36], seems not to contribute to the effects of low

![Quantitative Western blot analysis of HCN1 and HCN2 protein expression in mouse cortical cultures.](image)
Ih has been considered a source of tonic excitatory drive promoting the emergence of oscillations in interneuronal networks [43]. In hippocampal fast-spiking basket cells, this has been demonstrated to have stabilizing effects on firing and coherence of gamma oscillations [44]. Furthermore, medial septum GABAergic and parvalbumin/HCN-immunoreactive neurons were shown to serve as pacemakers of hippocampal theta rhythmic activity via the synchronization of hippocampal interneurons in vivo [45, 46]. Based on the present data and morphological features of L-INs [20], we suggest that L-INs are essential for the synchronization of SBA and may also contribute to the maintenance of SBA during maturation of cortical networks. During its limited expression between neurogenesis and the onset of experience-dependent plasticity, the large-scale synchronized early network oscillations are essential to neuronal survival [47], to the setup of proper long-range network wiring [48–50], and to the development of local circuits [51, 52].

According to our results, Ih plays a prominent role in increasing the pace of early SBA. Interestingly, the developmental increase of Ih, which reaches adult expression levels at the end of the second week in vivo, has also been suggested to contribute to desynchronization of early network bursting by limiting action potential firing during correlated network events [53]. However, the role of Ih might change from promoting synchronization of SBA at early stages of development to de-correlating SBA at later stages. Accordingly, developmental and activity-dependent changes in channel structure and properties [54, 55] might contribute to a shifted role of Ih in network activity.

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