Maturation of AMPAR Composition and the GABA_A Reversal Potential in hPSC-Derived Cortical Neurons

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Rodent-based studies have shown that neurons undergo major developmental changes to ion channel expression and ionic gradients that determine their excitation-inhibition balance. Neurons derived from human pluripotent stem cells theoretically offer the potential to study classical developmental processes in a human-relevant system, although this is currently not well explored. Here, we show that excitatory cortical-patterned neurons derived from multiple human pluripotent stem cell lines exhibit native-like maturation changes in AMPAR composition such that there is an increase in the expression of GluA2(R) subunits. Moreover, we observe a dynamic shift in intracellular Cl− levels, which determines the reversal potential of GABA_A-mediated currents and is influenced by neurotrophic factors. The shift is concomitant with changes in KCC2 and NKCC1 expression. Because some human diseases are thought to involve perturbations to AMPAR GluA2 content and others in the chloride reversal potential, human stem-cell-derived neurons represent a valuable tool for studying these fundamental properties.

Key words: GABA; glutamate; neurotransmitter; patch clamp; qRT-PCR; stem cells

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Introduction

The ability to generate in vitro cortical neuronal populations from human pluripotent stem cells (hPSCs) provides an experimental resource for the investigation of human cortical physiology and disease (Hansen et al., 2011). hPSCs have been used to derive human excitatory cortical neurons (hECNs) in vitro that appear to recapitulate aspects of in vivo cortical development (Johnston et al., 2007; Zeng et al., 2010; Shi et al., 2012; Espuny-Camacho et al., 2013). We have developed a protocol that enables the generation of a propagatable pool of anterior neural precursor cells (aNPCs) from human embryonic stem (ES) cell and induced pluripotent stem cell lines (Bilican et al., 2014). aNPCs can be efficiently differentiated as a monolayer of excitatory cortical neurons that are representative of neuronal populations present within native upper and lower cortical layers. Our recent study (Bilican et al., 2014) demonstrated the ability of hECNs to fire action potentials, express voltage-gated and ligand-gated ion channels, and exhibit putative synaptic activity; however, there is a need to identify specific physiologically postnatal/adult properties of such neurons to aid in vitro human studies of neurodevelopmental physiology and modeling of neurodevelopmental and adult diseases (Yang et al., 2011; Sandoe and Eggan, 2013).

Here, we demonstrate that hECNs differentiated from aNPCs exhibit native-like maturation shifts in AMPAR subunit composition and in the relative expression of the Na+−K+−Cl− cotransporter-1 (NKCC1) and K+−Cl− cotransporter-2 (KCC2) subunits to reduce intracellular chloride concentration ([Cl−]INT), which determine the nature of GABA receptor type-A (GABA_A) function (Blaesse et al., 2009). Beyond a platform for the study of maturation, these hECNs are of interest for studies of human diseases in which AMPAR and [Cl−]INT regulation are disrupted (Blaesse et al., 2009; Wright and Vissel, 2012).

Materials and Methods

In vitro hECN preparation. A detailed description of the derivation of hECNs, including immunohistochemistry protocols, can be found in Bilican et al. (2014) and is summarized in Figure 1A. The H9 (female), hereafter referred to as ES1 (WiCell) and SHEF4 (male), referred to as ES2 (UK Stem Cell Bank) human ES cell lines were obtained under full ethical/Institutional Review Board approval of the University of Edinburgh. For IPS1 (male M337V mutant line 1; Bilican et al., 2012) and IPS2 (female in-house control line) human IPS cell lines, written informed consent was obtained from each participant.

hECN culture media supplements. Forskolin (10 μM) was present during weeks 1–3 of differentiation. For AMPAR characterization, medium was supplemented with BDNF and GDNF (both 5 ng/ml) from week 3 onward. For experiments to determine [Cl−]INT, BDNF and GDNF were omitted unless otherwise stated. When cultures were maintained beyond 5 weeks, the culture medium was supplemented with IGF1 (10 ng/ml) from week 4 of differentiation.
Developmental maturation of AMPARs in hECNs

Functional AMPAR expression was assessed, at weekly intervals, by the ability of AMPA (50 mM) to elicit whole-cell currents. With gqRT-PCR. Protocols were performed as described previously (Bilican et al., 2014). Forward and reverse sequences (5‘–3’ ) of primers used were as follows: GRIA1: TGCTTTTGCGCAACTCACAGA, GGCATAGACTCCTTTGGAGAAC; GRIA2: CATTCCAGTGGACCCGACCT, GGTTGCGAAACTTGCTCCATTGA; GRIA3: ACCATCGCATAAGGGCTGG; GCTTGGTGTTGTTGTTAACTGTCAGC; GRIA4: TTCCGAGCAGGTTGCAAATA, GCA TTCGGGCGTTGGTTTATGA; SLC1A2: TG- GTGTTGCAATGGTCTAA, TCCACCTCCA TTCGACGTAC; SLC12A5: GGAAGGAAATGATAGCAGGCTGA, TCCACTCCTTCCTCCACATCT. Electrophysiology. Whole-cell patch-clamp recordings were made from hECNs using an Axon Multiclamp 700B amplifier (Molecular Devices). AMPA-evoked currents were made at a holding potential of −74 mV in the presence of picrotoxin (50 µM), strychnine (20 µM), and TTX (300 nM). GABA-evoked currents were made at a holding potential of −14 mV in the presence of CNQX (5 µM), d-APV (50 µM), strychnine, and TTX. Patch electrodes were filled with a solution containing the following (in mM): 155 K-gluconate, 2 MgCl2, 10 Na-HEPES, 5 Na-Pi-Creatine, 2 Mg2-ATP, and 0.3 Na3-GTP, pH 7.3, 300 mOsm, resistances 4–7 MΩ. Cover-slip containing hECNs were placed in the recording chamber and perfused using gravity feed with an extracellular solution composed of the following (in mM): 152 NaCl, 2.8 KCl, 10 Na-HEPES, 10 Na-Pi, 2 CaCl2, 1 glucose, pH 7.3, 320–330 mOsm. Recordings were made at 20–23°C. The liquid junction potential (LJP) was calculated to be +14 mV (JPCalc; Clampex). Perforated whole-cell recordings were performed using a gamicidin (50–100 µg/ml) supplemented patch-pipette solution containing the following (in mM): 145 KCl, 0.1 CaCl2, 1 EGTA, 10 HEPES, pH 7.3, 320–330 mOsm; the LJP was calculated to be −4.3 mV. The progress of perforation in the voltage-clamp configuration was monitored by the development of capacitive transients. Abrupt rises in transients indicated membrane rupture to the whole-cell configuration and the experiment was terminated. Once perforation stability was established, series resistances were <40 MΩ.

Data analysis. Nonstationary fluctuation analysis of slowly rising whole-cell currents evoked by AMPA were used to estimate the AMPAR unitary single-channel current. The analysis of AC and DC currents was performed as described previously (Brown et al., 1998) with the exception that the AC-coupled signal was filtered with a 1–1200 Hz band-pass frequency. All other recordings were low-pass filtered online at 2 kHz and all recordings were digitized at 10 kHz and recorded to computer hard disk using the WinEDR V2.7.6 Electrophysiology Data Recorder (J. Dempster, University of Strathclyde, United Kingdom; www.strath.ac.uk/Departments/PhysPharm/). [Cl−] INT was calculated using the Nernst equation in which extracellular [Cl−] was converted into activity using an activity coefficient.

Data are presented as mean ± SEM. The number of experimental replicates is denoted as n and N represents the number of de novo preparations of batches from which n is obtained.
time, hECNs displayed an increase in AMPAR-mediated current densities together with an increase in the proportion of hECNs that responded to application of AMPA. Five weeks after aNPC plate-down, ~90% of all cells gave currents (Fig. 1B) that could be blocked by CNQX. AMPARs are tetrameric assemblies of different combinations of GluA1, GluA2, GluA3, or GluA4 subunits. AMPAR composition in rodent and human excitatory cortical neurons is developmentally regulated; GluA1/GluA4-containing stoichiometry predominates in immature neurons, whereas mature neurons have an increased expression of GluA1 and GluA2 subunits (Monyer et al., 1991; Talos et al., 2006a, 2006b; Orlandi et al., 2011). We assessed relative AMPAR subunit mRNA expression levels at 2 and 5 weeks after aNPC plate-down by qRT-PCR (Fig. 1C) and found significant increases in the relative mRNA expression levels for each of the AMPAR subunits.

GluA2 subunits are RNA edited, resulting in a gene-encoded glutamine codon in the M2 reentrant loop region being replaced by an arginine codon in ~99% of mRNA transcripts (for review, see Wright and Vissel, 2012). Moreover, the biophysical properties of AMPARs are considerably influenced by the presence of edited GluA2 (GluA2(R)) subunits that give rise to AMPARs that possess low single-channel conductances, low Ca$^{2+}$ permeability, and insensitivity to polyamine-mediated channel block (Traynelis et al., 2010). To assess whether AMPARs expressed by hECNs show a developmental increase in the proportion of receptors containing GluA2(R) subunits, we performed nonstationary fluctuation analysis to estimate the mean unitary conductance ($\gamma$) of AMPARs at weeks 2 and 5 after aNPC plate-down. Current-variance plots constructed from AC- and DC-coupled components of whole-cell AMPAR-mediated responses were used to estimate unitary conductances (Fig. 1D,E). Figure 1F illustrates that, for the ES1 cell line, the mean unitary conductance decreases at week 2 from 10.9 ± 1.0 pS ($n = 13$, $N = 2$) to 4.5 ± 0.4 pS at week 5 ($n = 12$, $N = 3$, $p < 0.001$, unpaired $t$ test). Significant decreases in conductance over this same period were also observed in hECNs derived from a separate ES cell line (denoted ES2) and from two independent IPS cell lines (Fig. 1F). The factors that control the developmental increase in the expression of GluA2(R)-containing AMPARs remain largely unknown (Traynelis et al., 2010) and we found no evidence that this regulation is mediated by excitatory or inhibitory neurotransmitter action or is regulated by an activity-dependent process, because pharmacological blockade from weeks 1–5 of AMPAR, NMDAR, GABA$_A$R, and glycine receptors by CNQX (15 $\mu $M), d-AP5 (25 $\mu $M), bicuculline (100 $\mu $M), and strychnine (400 $\mu $M), respectively, and voltage-dependent Na$^+$ channels by TTX (1 $\mu $M) did not prevent the decrease in conductance that occurs between weeks 2 and 5 after aNPC differentiation (Fig. 1G). Furthermore, media supplements (see Materials and Methods) are not implicated in the reduction of conductance, because hECNs maintained in the absence of supplements displayed equivalent estimates of conductance at week 2 (12.7 ± 1.5; $n = 11$, $N = 3$) and week 5 (5.0 ± 1.1; $n = 5$, $N = 1$).

An increase in the proportion of AMPARs containing GluA2(R) subunits was confirmed in hECNs by assessing the polyamine sensitivity of AMPAR-mediated current to 1-naphthyl acetyl spermine (NASPM), a selective antagonist of GluA2(R)-lacking AMPARs (Koike et al., 1997). At week 2, NASPM (3 $\mu $M) strongly inhibited (77 ± 5%) AMPA-evoked steady-state whole-cell currents (Fig. 2A). The extent of this block is consistent with the expression of AMPARs that lack GluA2(R) subunits. At week 5, NASPM gave only modest inhibition (Fig. 2B,C). Overall, these data, together with a development switch in unitary conductance of AMPARs, are entirely consistent with a switch from a GluA2(R)-lacking to a GluA2(R)-containing AMPAR population.

hECNs show a developmental reduction in [Cl$^-$]$_{\text{INT}}$

Functional GABA$_A$R expression was assessed at weekly intervals by the ability of GABA (100 $\mu $M) to elicit whole-cell currents. The magnitude of steady-state GABA-evoked currents increased with increasing periods of culture and, after 4 weeks of differentiation, all cells responded to GABA (Fig. 3A) with robust currents that were blocked by picrotoxin. Fast depolarizing actions of GABA on neural precursor cells and immature cortical neuron wells are well documented and are a consequence of higher [Cl$^-$]$_{\text{INT}}$ (~25 mM) due to the higher relative expression level of NKCC1 over KCC2 (Ben-Ari et al., 2007). In both rodent and human cortex, the postnatal expression levels of KCC2 increase (Dzhala et al., 2005), leading to a net extrusion of Cl$^-$ from excitatory neurons and levels of [Cl$^-$]$_{\text{INT}}$ of 5–10 mM, which results in a net inward movement of Cl$^-$ and a hyperpolarizing response in neurons at or near their resting membrane potentials (Blaesse et al., 2009). Using gramicidin-perforated current-clamp recording to prevent disruption of [Cl$^-$]$_{\text{INT}}$ (Kyrizos and Reichling, 1995), we assessed the nature of the GABA response in hECNs when held at a potential of ~60 mV. After 1 week of culture after aNPC plate-down, GABA elicited a depolarizing response (Fig. 3B), whereas at later periods, the response was hyperpolarizing (Fig. 3C). To determine the
The presence of IGF1 did not show any change in reversal potential for GABA$_A$R-mediated responses ($E_{\text{GABA}}$), perforated voltage-clamp recordings were made from hECNs at various time points after aNPC differentiation. Figure 3D illustrates a series of GABA-activated current traces obtained from a week 5 hECNs and recorded at holding potentials that allowed $E_{\text{GABA}}$ to be determined. Figure 3E summarizes the extent to which $E_{\text{GABA}}$ changes with time in culture. At week 1, the mean $E_{\text{GABA}}$ is $-43.0 \pm 2.0$ mV ($n = 12$, $N = 3$) and becomes progressively more negative such that, at week 5, it is $-59.2 \pm 1.5$ mV ($n = 11$, $N = 2$, $p < 0.001$). To examine the possibility that $E_{\text{GABA}}$ could become more negative, we extended the culture period of hECNs to 7 weeks, which necessitated the inclusion of IGF1 in the culture media to improve cell viability (see Materials and Methods), and observed a further decrease in $E_{\text{GABA}}$ to $-70.3 \pm 0.7$ mV ($n = 13$, $N = 3$, $p < 0.001$). IGF1 was unlikely to have directly affected the change in $E_{\text{GABA}}$ because hECNs at week 5 cultured in the presence of IGF1 did not show any change in $E_{\text{GABA}}$ ($-62.9 \pm 2.3$ mV; $n = 10$, $N = 2$). In these experiments, BDNF and GDNF were omitted because these neurotrophic factors have been reported to cause alterations in Cl$^-$ transporter function and expression (Blaesse et al., 2009). Indeed, although $E_{\text{GABA}}$ becomes more negative in the presence of BDNF and GDNF, it remains $\sim 10$ mV more positive than is observed for hECNs cultured in the absence of neurotrophic factors (Fig. 3E).

**Discussion**

We have demonstrated the ability of hECNs to functionally express mature GluA2(R)-containing AMPARs and exhibit a reduction in [Cl$^-$]$_{\text{INT}}$ to adult-like levels. Importantly for the interpretation of our results, differentiating aNPCs are likely to be the predominant cellular phenotype at early culture time points (weeks 1 and 2) and, by later time points ($\geq$week 4), the cultures are predominantly of hECN identity (Fig. 1A; Bilican et al., 2014).

We demonstrate that AMPAR single-channel conductance at weeks 2 and 5 significantly reduces from 10.9 to 4.5 pS in accordance with increased functional expression of the GluA2(R) subunit that confers a lower conductance to the AMPAR complex. Indeed, the conductance values at weeks 2 and 5 are directly comparable to conductance measurements of AMPAR-mediated events from rodent excitatory cortical neurons during a postnatal developmental period involving the upregulation of the GluA2(R) subunit (Brill and Huguenard, 2008). Furthermore, week 5 data correspond well to those of recombinantly expressed heteromeric GluA2(R)-containing AMPARs (Swanson et al., 1997). A GluA2(R)-lacking to GluA2(R)-containing switch in AMPAR subunits is confirmed by the observed high sensitivity of
AMPAR-mediated currents at week 2, but not at week 5, to GluA2(R)-lacking the AMPAR blocker NASPM.

The principal AMPAR composition in native cortical preparations has been described to develop from an immature GluA1/GluA2(R)-lacking the AMPAR blocker NASPM. GluA4 subunit. Nevertheless, these data do not indicate the level of functional protein. At week 2, the cultures contain a mixture of neurons and aNPCs, so it is interesting that native primary human embryonic cortical NPCs express unedited forms of the GluA2 subunit [GluA2(Q)] and, upon neuronal differentiation, express the edited GluA2(R) subunit over an equivalent culture period (Whitney et al., 2008).

The underlying mechanism of this AMPAR subunit shift has not yet been described in native excitatory cortical neurons. Here, we demonstrate that the reduction of AMPAR conductance is not prevented by antagonists of neurotransmitter ligand-gated ion channels and voltage-gated ion channels, indicating that the developmental upregulation of GluA2(R)-containing AMPARs is activity-independent. Indeed, it has been reported that NPC differentiation results in a large upregulation of the ADAR2 enzyme responsible for the editing of the GluA2 subunit Q/R site (Monyer et al., 1991). Our data are consistent with an upregulation of the ADAR2 enzyme at week 2, but not at week 5, to GluA2(R) coincided with a switch in the relative mRNA expression levels of the Cl⁻ transporters NKCC1 and KCC2 (Blaesse et al., 2009). This implies that the high [Cl⁻]ᵢNT observed at week 1 is due to dominant NKCC1 activity and the low [Cl⁻]ᵢNT observed at week 7 is a result of overriding KCC2-mediated Cl⁻ extrusion.

To obtain a reduction in [Cl⁻]ᵢNT, hECNs had to be cultured for extended periods in the absence of BDNF and GDNF but with IGF supplementation. IGF1 was shown not to have an effect upon E_{GABA} in week 5 hECNs, but has previously been associated with an upregulation of KCC2 expression in rodent hippocampal neurons (Kelsch et al., 2001). The inclusion of BDNF and GDNF, however, appears to slow the developmental [Cl⁻]ᵢNT reduction. Neurotrophic supplements are commonly used in hPSC-derived neuronal cultures with little regard for their potential impact on neuronal Cl⁻ transporter function (Blaesse et al., 2009) and receptor desensitization (Frank et al., 1996). We thus advocate caution when using these agents, particularly for functional studies.

The mechanistic control of neuronal [Cl⁻]ᵢNT development has received considerable attention and specific reports of mechanisms of NKCC1 and KCC2 regulation have been controversial (Ben-Ari et al., 2012). The inability to prevent the reduction of [Cl⁻]ᵢNT by chronic blockade of ion channel activity are in agreement with an activity-independent mechanism in hECNs that has been observed in native neuronal preparations (Ben-Ari et al., 2007).

In summary, we show the ability of hECNs to functionally express mature GluA2(R)-containing AMPARs. This enables investigation into disease mechanisms that may affect the editing at the Q/R site. Furthermore, the ability of hECNs to exhibit a reduction in [Cl⁻]ᵢNT to adult levels enables homeostatic and disease studies that potentially affect Cl⁻ transporter function.

References


