Lineage Reprogramming of Astroglial Cells from Different Origins into Distinct Neuronal Subtypes

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http://dx.doi.org/10.1016/j.stemcr.2017.05.009

SUMMARY

Astroglial cells isolated from the rodent postnatal cerebral cortex are particularly susceptible to lineage reprogramming into neurons. However, it remains unknown whether other astroglial populations retain the same potential. Likewise, little is known about the fate of induced neurons (iNs) in vivo. In this study we addressed these questions using two different astroglial populations isolated from the postnatal brain reprogrammed either with Neurogenin-2 (Neurog2) or Achaete scute homolog-1 (Ascl1). We show that cerebellum (CerebAstro) and cerebral cortex astroglia (CtxAstro) generates iNs with distinctive neurochemical and morphological properties. Both astroglial populations contribute iNs to the olfactory bulb following transplantation in the postnatal and adult mouse subventricular zone. However, only CtxAstro transfected with Neurog2 differentiate into pyramidal-like iNs after transplantation in the postnatal cerebral cortex. Altogether, our data indicate that the origin of the astroglial population and transcription factors used for reprogramming, as well as the region of integration, affect the fate of iNs.

INTRODUCTION

Direct lineage reprogramming of somatic cells into induced neurons (iNs) is a promising strategy to study the molecular mechanisms of neuronal specification, identify potential therapeutic targets for neurological diseases, and eventually repair the CNS after acute or neurodegenerative injury (Arlotta and Berninger, 2014). Brain resident cells such as forebrain astrocytes present some great advantages when we consider the easiness and efficiency of conversion into iNs both in vitro (Berninger et al., 2007) and in vivo (Guo et al., 2014; Hu et al., 2015; Niu et al., 2013, 2015) compared with other cell types (Vierbuchen et al., 2010; Pang et al., 2011; Victor et al., 2014).

However, still little is known about the role of the environment in the specification of iNs fates during astrogial lineage reprogramming in vivo (Heinrich et al., 2015). Also, which are the subtypes of iNs generated by lineage reprogramming and how could be controlled remain poorly understood. Following retroviral-mediated expression of ACHAETE SCUTE HOMOLOG-1 (ASCL1) or NEUROGENIN-2 (NEUROG2) in vitro, forebrain astrocytes are reprogrammed mostly into GABAergic or glutamatergic neurons, respectively (Heinrich et al., 2010). These observations led to the suggestion that ASCL1 and NEUROG2 would be instructing neuronal phenotypes, reminiscent of their reported role in cortical development (Schuurmans and Guillemot, 2002). However, the same transcription factors (TFs) are also expressed by non-overlapping progenitor populations in the developing cerebellum, contributing to the generation of GABAergic neurons in cerebellar nuclei, Purkinje cells, and inhibitory interneurons of the cerebellar cortex (Zordan et al., 2008). Thus, it is possible that astroglial cells isolated from cerebral cortex or cerebellum and lineage reprogrammed with NEUROG2 or ASCL1 could retain a “molecular memory” of their origin and generate iNs with distinctive phenotypes.

To test this possibility, we first investigated the potential to reprogram astroglial cells isolated from the postnatal cerebellum (CerebAstro) into iNs. Next, we compared the phenotypes of iNs derived from lineage-reprogrammed CerebAstro and cerebral cortex astroglia (CtxAstro), both in vitro and in vivo, following transplantation in the mouse cerebral cortex or subventricular zone (SVZ). Our results show that either ASCL1 or NEUROG2 is sufficient to convert CerebAstro into iNs adopting mostly a GABAergic phenotype. Following transplantation in the postnatal subventricular zone, both types of astroglial cells generate iNs that migrate to the olfactory bulb and integrate as granular or periglomerular neurons, albeit at different ratios depending on the reprogrammed astrogial population. However, after transplantation in the postnatal cerebral cortex, only iNs derived from CtxAstro reprogrammed with NEUROG2 adopted fates reminiscent of cortical pyramidal neurons. Collectively, our results suggest that both the origin of the astrogial population used for reprogramming and the region of grafting in the brain affect the phenotype of iNs.
RESULTS

Expression of ASCL1 or NEUROG2 Efficiently Reprograms Cerebellum Astroglia in iNs

Direct lineage reprogramming of astroglial cells into iNs following viral or chemical delivery of neurogenic TFs is well established for astrocytes isolated from the cerebral cortex of postnatal mice (Berninger et al., 2007; Gascón et al., 2015; Heinrich et al., 2010; Masserdotti et al., 2015). However, it remains unknown whether astroglial cells from different regions of the CNS hold the same potential. To address this possibility, we set out to investigate whether the expression of single proneural TFs could lineage reprogram CerebAstro into iNs. Toward this aim, we generated cultures enriched for cerebellum astroglia and nucleofected the cells with plasmids carrying the genes encoding for NEUROG2 (Neurog2-DsRed), ASCL1 (Ascl1-DsRed), or only the reporter protein dsRed (dsRed). To determine the cellular composition of cultures prior to nucleofection, we immunostained cells for the astrocytic marker glial fibrillary acidic protein (GFAP), the pan-neuronal marker class III β-tubulin (BIII-TUBULIN), the neural stem cell transcription factor SOX2, and the oligodendrocyte marker O4. We observed that the vast majority of cells were positive for GFAP, whereas only a very small percentage of cells were positive for BIII-TUBULIN (GFAP: 93±1%; BIII-TUBULIN: 3%±1%; n = 2,254 cells) (Figures S1A–S1E). We did not detect expression of SOX2 and O4 in the cultures (data not shown). One day post nucleofection (dpn), we observed that virtually all transduced cells expressed GFAP (Figures S1H–S1J). However, at 7–8 dpn most CerebAstro transduced with ASCL1 or NEUROG2 adopted a neuronal-like morphology and expressed BIII-TUBULIN (Figures 1A–1K; ASCL1: 73% ± 2% BIII-TUBULIN+ cells, n = 1,320 dsRed+ cells; NEUROG2: 54% ± 2% BIII-TUBULIN+ cells, n = 1,234 dsRed+ cells). In sharp contrast, cells nucleofected with dsRed kept astrocyte morphology and did not express BIII-TUBULIN (Figures S1F–S1G). Similar rates of iNs were observed with cerebral cortex astroglia nucleofected with Neurog2-DsRed, Ascl1-DsRed, or control plasmids (Figures S1K–S1Q), indicating that CerebAstro and CtxAstro are equally prone to lineage reprogramming into iNs after expression of NEUROG2 or ASCL1. According to previous data in the literature (Law-yell et al., 2000; Masserdotti et al., 2015), we also observed that the capacity of CerebAstro and CtxAstro to generate multipotent neurospheres and to lineage reprogramming decreased after successive passages in vitro (Figures S1R–S1U).

Next, we studied the process of CerebAstro lineage reprogramming into neuron using time-lapse video-microscopy (Figure S2 and Movies S1, S2, and S3). We observed that most transfected cells in control or experimental conditions displayed astroglial morphology. However, only cells transduced with either Neurog2-DsRed or Ascl1-DsRed underwent a thorough process of morphological changes ending with the acquisition of a neuronal-like morphology and the expression of neuron-specific markers (Figures S2A–S2L and Movies S1, S2, and S3). Yet expression of ASCL1 and NEUROG2 induced cell death in a high percentage of cells (Figure S2M), suggesting that, similar to cerebral cortex astroglia, the reprogramming process can be hampered by metabolic constraints (Gascón et al., 2015). Notably, we also observed that few transfected cells underwent cell division (Figure S2N), suggesting that mostly postmitotic astroglial cells are converted into iNs. Accordingly, most iNs were generated without cell division, as indicated by the low proportion of iNs incorporating bromodeoxyuridine (BrdU) after expression of NEUROG2 or ASCL1 (Figure 1Y). To further confirm the astroglial nature of lineage-converted cells, we genetically labeled astrocytes using the double-transgenic GLAST-CreERT/CAG-CAT-GFP mouse (Heinrich et al., 2010). Animals received tamoxifen through the mother milk from postnatal day 5 (P5) to P7, leading to the Cre-mediated recombination in cerebellum astroglia in vivo (Figure S3A). We observed that 95% of iNs also expressed GFP, confirming the astroglial origin of the original cells.

Figure 1. ASCL1 and NEUROG2 Convert Cerebellum Astroglia into Induced Neurons

(A–J) Representative pictures taken from CerebAstro cultures 7 days post nucleofection (dpn) with Ascl1-DsRed (A) or Neurog2-DsRed plasmid (F). Many nucleofected RFP+ cells (inside dotted boxes) adopt neuronal-like morphologies and express the immature neuronal marker BIII-TUBULIN (BIII-TUB) after transduction with (A) ASCL1, magnification of boxes 1 and 2 in (B) to (E) or (B) NEUROG2, magnification of boxes 1 and 2 in (G) to (J).

(K) Quantification of BIII-TUB expression in CerebAstro culture 7 dpn with Ascl1-DsRed (black bar), Neurog2-DsRed (gray bar), or control plasmid dsRed (white bar) (see Figure S1 for control condition). ***p < 0.001. N.F, not found.

(L–W) Expression of mature neuronal markers in lineage-reprogrammed CerebAstro iNs 15 dpn. Example of RFP+ ASCL1-iNs (L–N) and NEUROG2-iNs (O–Q), stained for MAP2. Example of RFP+ ASCL1-iNs (R–T) and NEUROG2-iNs (U–W) stained for NEUN.

(X) Quantification of MAP2+ and NEUN+ among RFP+ cells 15 dpn with ASCL1 (black bar) or NEUROG2 (white bar).

(Y) Quantification of BrdU incorporation and BIII-TUB expression in nucleofected cells 8 dpn with ASCL1 or NEUROG2. Nuclei are stained with DAPI (blue).

Data are derived from six independent experiments (mean ± SEM). Scale bars, 25 μm. See Figures S1–S3.
**Functional Properties of Cerebellum Astroglia-Derived iNs**

After, we investigated the phenotypic maturation of lineage-reprogrammed CerebAstro iNs. For this, nucleofected cultures were maintained for 2 weeks in differentiation medium and assessed for the expression of the neuronal microtubule-associated protein 2 (MAP2) and the neuronal nuclei protein Foxb3 (NEUN). We observed that CerebAstro transfected with either ASCL1 or NEUROG2 reprogrammed into iNs expressing MAP2 (ASCL1: 83% ± 7%, n = 321 cells; NEUROG2: 91% ± 9%, n = 236 cells) and NEUN (ASCL1: 90% ± 9%, n = 365 cells; NEUROG2: 92% ± 5%, n = 210 cells) (Figures 1L–1X). To further stimulate the synaptic maturation of iNs, we grew transfected CerebAstro in the presence of co-cultured neurons (see Supplemental Experimental Procedures). Thirty days after nucleofection we could observe expression of SYNAPSIN 1, a synaptic vesicle protein involved in the control of neurotransmitter release (Hvalby et al., 2006), in juxtaposition to dsRed' processes (Figures 2A–2H), suggesting that iNs could be establishing synaptic contacts with co-cultured neurons.

To study the electrical properties of iNs, we performed patch-clamp recordings of fourteen ASCL1-iNs and nine NEUROG2-iNs grown in the absence of co-cultured primary neurons (Figures 2I–2L). Of the 14 ASCL1-iNs, ten responded with action potentials and had a mean input resistance of 337 ± 76 MΩ, a mean resting membrane potential of −52 ± 2 mV, and a mean action potential amplitude of 50 ± 7 mV following minimal depolarizing current injections (50 pA, 500 ms) (Figure 2I). NEUROG2-iNs (n = 9 cells) had a mean input resistance of 352 ± 78 MΩ, a mean resting membrane potential of −56 ± 2 mV, and a mean action potential amplitude of 42 ± 4 mV following the same current injection (50 pA, 500 ms) (Figure 2L). Hyperpolarizing currents (−100 pA, 500 ms) generated rebound action potentials in 50% (5/10) of the ASCL1-cells with a mean amplitude of 63 ± 4 mV and 22% (2/9) of the NEUROG2-cells with a mean amplitude of 39 ± 2 mV (Figure 2I). Notably, some iNs showed spontaneous electrical activity, further suggesting that iNs established synaptic contacts in the culture (Figure 2I). Recorded iNs could be classified into cells with spikelets (40% of ASCL1-iNs; 66% of NEUROG2-iNs), cells with a few spikes (40% of ASCL1-iNs; 22% of NEUROG2-iNs), and regular spiking cells (30% of ASCL1-iNs; 11% of NEUROG2-iNs) (Figures 2I and 2J). Moreover, current-voltage relationships of ASCL1-iNs and NEUROG2-iNs revealed a significant difference between instantaneous values in response to hyperpolarizing current injections (Figure 2K). Collectively, these observations suggest that NEUROG2 or ASCL1 can efficiently reprogram cerebellum astroglia into functional iNs.

**ASCL1 and NEUROG2 Induce Different Neuronal Subtypes in Astroglia of Different Origin**

Another important aspect of neuronal differentiation is the establishment of axial polarity and process growth (Barnes and Polleux, 2009; Whitford et al., 2002; Takano et al., 2015). Interestingly, we noted that iNs displayed different morphologies 15 days after expression of ASCL1 or NEUROG2 in CerebAstro or CtxAstro (Figure S4). To quantify these differences, we analyzed morphological features of iNs using the Sholl analysis. We found that iNs derived from CerebAstro expressing ASCL1 displayed significantly longer processes, with an increased amount of secondary and tertiary branches compared with iNs expressing NEUROG2 (Figure S4F). Notably, we also found that iNs derived from CerebAstro expressing ASCL1 were more complex than those derived from CtxAstro expressing the same TF, whereas the opposite was observed with NEUROG2 (Figures S4H–S4I).

Next, we set out to measure the axial distribution of cell processes (Figures S4G and S4K). We observed that expression of ASCL1 in CerebAstro generate higher polarized iNs than NEUROG2 (Figure S4G). The opposite result was observed when we compared iNs derived from CtxAstro (Figure S4K). Collectively, these morphological parameters indicate that both the TF and the type of reprogrammed astroglial cell may interfere with iNs phenotypes.

To further investigate possible phenotypic distinctiveness of iNs, we analyzed the expression of the neurotransmitter GABA and glutamate, as well as TBR1, a TF associated with glutamatergic neurons (Hevner et al., 2006), in iNs derived from CerebAstro or CtxAstro (Figures 3 and S5). We observed that the percentage of GABA was higher than GLUTAMATE-expressing iNs following expression of both ASCL1 and NEUROG2 in CerebAstro (Figures 3A–3S; CerebAstro GABA+ iNs ASCL1: 70% ± 6%, n = 68 cells and NEUROG2: 59% ± 7%, n = 58 cells; CerebAstro Glut1+ iNs ASCL1: 38% ± 6%, n = 52 cells and NEUROG2: 48% ± 2%, n = 63). The GABAergic phenotype of astroglia-derived iNs was also confirmed using astrocytes isolated from GAD67-GFP mice (Figures 3T–3Z). In contrast, the same TFs induced different fates in CtxAstro. While iNs derived from CtxAstro reprogrammed with NEUROG2 expressed mostly GLUTAMATE, ASCL1 induced more GABA expression (Figures 3S and S5G–S5L; CtxAstro GABA+ iNs ASCL1: 64% ± 15%, n = 60 cells and NEUROG2: 19% ± 7%, n = 50 cells; CtxAstro Glut1+ iNs ASCL1: 50% ± 9%, n = 67 cells and NEUROG2: 81% ± 4%, n = 54 cells). We also noted that NEUROG2 induced the expression of TBR1 only in CtxAstro (Figures S5A–S5F; 79% ± 2% of TBR1+, n = 30 iNs).

Next, to evaluate whether iNs derived from CtxAstro or CerebAstro adopt the phenotype of distinct GABAergic classes, we compared the expression of the calcium binding
Figure 2. Electrical and Synaptic Development of CerebAstro-Derived iNs

(A–H) Lineage-reprogrammed CerebAstro iNs express SYNAPSIN 1 (green) 30 dpn with Ascl-DsRed or Neurog2-DsRed plasmids. Example of RFP+/SYNAPSIN+ iNs 30 dpn with ASCL1 (A–D) or NEUROG2 (E–H). (D–H) Magnifications of dashed boxes in (C) and (G), respectively.

(I–L) Electrophysiological properties of ASCL1- and NEUROG2-iNs. Current clamp traces from ASCL1-iNs showing spikelet (left), few spikes (middle), and regular spiking (right) in response to depolarizing current injections (50 pA, 500 ms) (I, top). Arrows highlight spontaneous excitatory postsynaptic potentials in recorded cells (I). Fluorescence images of the recorded ASCL1-iNs (I, bottom).

(J) Example of current clamp traces from a NEUROG2-iNs responding with a few spikes to depolarizing current injections (50 pA, 500 ms) (J, top). Note the deeper sag due to hyperpolarizing current injections (−100 pA, 500 ms) compared with ASCL1-iNs. Fluorescence image of the recorded NEUROG2-iN (J, bottom).

(K) Current-voltage relationships for ASCL1-iNs (circles) and NEUROG2-iNs (triangles) measured at the beginning of the onset of the current steps.

(L) Plot of mean input resistance, resting membrane voltage, and mean action potential amplitude following the current injection (50 pA, 500 ms) of ASCL1-iNs (black plot) and NEUROG2-iNs (white plot).

Data are derived from three independent experiments (mean ± SEM). Scale bars, 50 μm.
Figure 3. Lineage-Reprogrammed Cortical and Cerebellar iNs Adopt Different Neurotransmitter Phenotypes

(A–J) Examples of CerebAstro-derived RFP+ iNs expressing GABA 20 dpn with either ASCL1 (A) or NEUROG2 (D). Single confocal z stacks are shown in higher magnification to confirm the colocalization of GABA and RFP in iNs (B, C, and E–J). Observe the similar pattern of GABA expression between co-cultured hippocampal neurons (RFP+, yellow arrowhead in D) and iNs (RFP+, white arrowheads in D).

(K–R) Example of CerebAstro-derived RFP+ iNs expressing GLUT 20 dpn with either ASCL1 (K–O) or NEUROG2 (P–R). Single confocal z stacks are shown in higher magnification to confirm the colocalization of glutamate and RFP in iNs (B, C, and E–J). Observe the presence of GLUT+/C0 (K–M, yellow arrowhead) and GLUT+ (K, N, and O, white arrowhead) RFP+ iNs.

(S) Quantification of GABA+ and GLUT+ cells among RFP+ iNs derived from CerebAstro nucleofected with ASCL1 (orange) or NEUROG2 (yellow) and CtxAstro nucleofected with ASCL1 (dark green) or NEUROG2 (light green). *p < 0.05, **p < 0.01, ***p < 0.001.

(T–Z) Expression of GFP in CerebAstro and CtxAstro derived from GAD67-GFP mice 12 dpn with ASCL1. Example of RFP+/GFP+/BIII-TUB+ iN derived from CerebAstro (T, dashed box magnified and showing single channels in U–W) or CtxAstro (X, dashed box magnified and showing single channels in Y–Z). GLUT, GLUTAMATE; BIII-TUB, BIII-TUBULIN. Statistical test: two-way ANOVA followed by Bonferroni’s post hoc test (mean ± SEM).

Data are derived from 3 independent experiments. Scale bars, 25 µm. See also Figure S5.
proteins CALBINDIN (CALB) and PARVALBUMIN (PARV) in iNs after expression of NEUROG2 or ASCL1 (Figure 4). We observed that ASCL1 expression in CerebAstro induced a higher ratio of CALB+ iNs, whereas NEUROG2 induced a higher ratio of PARV+ iNs (Figure 4; ASCL1: 72% ± 11%, and NEUROG2: 40% ± 10% CALB+ iNs; n = 170).

Figure 4. Expression of Calcium Binding Proteins Indicates that iNs Derived from Cortical and Cerebellar Astroglia Adopt Distinct GABAergic Phenotypes

(A–N) Examples of CerebAstro-derived iNs expressing PARV or CALB 30 dpn with either ASCL1 (A and G) or NEUROG2 (D and L). Single confocal z stacks of the two iNs (white and yellow arrowheads) are shown in higher magnification to confirm the colocalization of RFP with PARV (B, C, E, and F) or CALB (H–K, M, and N).

(O) Quantification of CALB+ and PARV+ cells among RFP+ iNs derived from both CerebAstro nucleofected with either ASCL1 (orange) or NEUROG2 (yellow) and CtxAstro nucleofected with ASCL1 (green). *p < 0.05, ***p < 0.001.

(P) Example of NEUROG2-iN reprogrammed from CerebAstro isolated from PARV-Cre/CAG-CAT-GFP mice.

(Q–S) Single confocal z stacks are shown in higher magnification to confirm the colocalization of RFP, GFP, and BIII-TUB.

(T) Quantification of ASCL1-iNs and NEUROG2-iNs expressing GFP after lineage reprogramming of PARV-Cre/CAG-CAT-GFP CerebAstro (two-way ANOVA followed by Bonferroni’s post hoc test, mean ± SEM). CALB, CALBINDIN; PARV, PARVALBUMIN; TUB, TUBULIN. **p < 0.01, ***p < 0.001.

Data are derived from three independent experiments. Scale bars, 20 μm. See also Figure S6.
118 cells, respectively; ASCL1: 30% ± 4% and NEUROG2 65% ± 7% PARV+ iNs; n = 120 and 143 cells, respectively. Interestingly, we also observed that more than half of CerebAstro-derived CALB+ iNs expressed CTIP2 (Figure S6) a transcription factor present in Purkinje neurons (Leid et al., 2004). Additionally, the parvalbuminergic phenotype of iNs was assessed using astrocytes isolated from PARV-Cre/CAG-CAT-GFP mice (Figures 3P–3T). We observed that 10 dpn with ASCL1 a similar proportion of iNs expressed GFP compared with iNs expressing PARV at 20 dpn (Figures 4O and 4T). However, the proportion of GFP+ iNs at 10 dpn after NEUROG2 expression was lower than that expressing PARV at 20 dpn (Figures 4O and 4T), suggesting that the expression of PARV in NEUROG2-iNs is regulated at some point between 10 and 20 dpn.

In contrast, ASCL1 expression in CtxAstro induced similar rates of CALB+ and PARV+ iNs (Figure 4O; CALB: 53% ± 17% iNs; PARV: 55% ± 6% iNs; n = 138 and 127 cells, respectively). Yet ASCL1 induced a significantly higher fraction of PARV+ iNs in CtxAstro compared with CerebAstro (Figure 4O). Altogether, these results indicate that ASCL1 and NEUROG2 instruct different subtypes of GABAergic iNs depending on the origin of the reprogrammed astroglial cell.

**CtxAstro, but Not CerebAstro, Nucleofected with NEUROG2 Differentiate into Pyramidal-like Neurons after Transplantation in the Postnatal Mouse Cerebral Cortex**

The distinctive iNs phenotypes observed in vitro raised the question as to whether these cells would keep such hallmarks after integration in a pre-existing circuitry. To investigate this possibility, we transplanted cortical and cerebellum astroglia following nucleofection with NEUROG2, ASCL1, or control plasmid in the postnatal mouse cerebral cortex and studied the phenotypes of grafted cells. To facilitate the identification of grafted cells, we transplanted astroglial cells isolated from GFP mice (Okabe et al., 1997) into wild-type animals. Twenty days post transplantation (dpt), animals were perfused and grafted cells were analyzed for their morphology and chemical markers (Figure 5). We observed that virtually all CtxAstro population nucleofected with control plasmid kept astroglial morphologies (Figures 5A and 5C; n = 1,145 GFP+ cells, n = 4 host animals). Interestingly, astroglial cells adopted morphologies similar to those of endogenous astrocytes in the gray and white matter (Figures S7A–S7C; Emsley and Macklis, 2006). In sharp contrast, about one-fifth of CtxAstro nucleofected with NEUROG2 and transplanted into the postnatal cerebral cortex adopted neuronal morphologies in all animals analyzed (n = 2,338 GFP+ cells in five host animals, 18% ± 7%) (Figures S5B–S5O). Intriguingly, we noted that about half of iNs settled in the layers II/III of the cerebral cortex (Figure 5D). Induced neurons adopted a pyramidal neuron-like morphology with apical dendrites toward the molecular layer and basal dendrites projecting radially (Figures 5E and 5F). We also observed basal axonal processes directed toward the white matter (Figure 5E). Interestingly, some GFP+ processes were observed in the contralateral corpus callosum (Figures S7K–S7O), suggesting that such processes originated from transplanted iNs. Some iNs showed very complex morphologies with several secondary and tertiary dendrites (Figure 5G), and the majority of these cells also showed dendritic spines (Figure 5I, 66% ± 8% of pyramidal-like iNs, n = 29 cells). Most cells with neuronal morphology also expressed NEUN+ (Figures S5J–S5L) and about 60% of iNs in layer II/III expressed the TF CUX1 (Figures 5M–5O, n = 15 cells). These data suggest that cortical astroglia reprogrammed with NEUROG2 can differentiate into pyramidal-like iNs in vivo.

In contrast, however, we found only a very small number of GFP+ cells with neuronal morphology following transplantation of CerebAstro nucleofected with NEUROG2, as well as CtxAstro or CerebAstro nucleofected with ASCL1 (Figures S7D–S7I; >500 cells counted/animal). Still, different from iNs derived from CtxAstro reprogrammed with NEUROG2, those iNs displayed morphologies reminiscent of non-spiny GABAergic interneurons (Figures S7J–S7F) and some expressed CALBINDIN (Figure S7G), suggesting a GABAergic phenotype. These observations suggest that reprogramming and survival of iNs following transplantation are affected by both the origin of the reprogrammed cell and TF used.

**CerebAstro and CtxAstro iNs Integrate as Olfactory Bulb Interneurons upon Transplantation in the SVZ**

Next, we set out to test whether integration of iNs could be facilitated by transplantation into a neurogenic milieu. Toward this aim, we transplanted CerebAstro and CtxAstro in the SVZ of postnatal animals following their nucleofection with either ASCL1 or dsRed. Thirty days after transplantation, we observed a substantial number of GFP+ cells in the rostral migratory stream (RMS) and olfactory bulb (OB) of host animals (Figures 6A–6F). In the control group, virtually all GFP+ cells nucleofected with control plasmid retained astrocytic morphologies in the SVZ, RMS, and OB (Figures S7P–S7Q, n > 300 cells). In contrast, a significant fraction of CtxAstro and CerebAstro nucleofected with ASCL1 adopted morphologies typical of bona fide OB neurons. Notably, iNs adopted typical morphologies of neurons of the granule cell layer (GCL) or periglomerular (PGL) (Figures 6H–6U) and expressed the mature neuronal marker NEUN (Figures 6Q and 6R). However, the ratio of iNs in the GCL and PGL varied depending on the reprogrammed astroglia (Figure 6G). While iNs derived from
CtxAstro mostly adopted the fate of GCL neurons (types I and III) (CtxAstro+ ASCL1: 68% ± 14%, n = 117 cells; CerebAstro+ ASCL1: 28% ± 10%, n = 87 cells), iNs derived from CerebAstro preferentially integrated as PGL neurons (CerebAstro+ ASCL1: 71% ± 10%; CtxAstro + ASCL1: 31% ± 14%). We also observed a small number of iNs with granular- and periglomerular-like morphologies after transplantation of CtxAstro nucleofected with NEUROG2 in the SVZ (Figures S7 R–S7S). Notably, the expression of calcium binding proteins in OB iNs was different from that observed in vitro (Figure 4). We could not observe PARV+ iNs or CALB+ iNs in the OB after transplantation of both CerebAstro and CtxAstro in the postnatal SVZ (Figures 6S–6U and data not shown).

Finally, we transplanted ASCL1 lineage-reprogrammed CtxAstro in the adult SVZ. We observed that, similar to our experiments in the postnatal brain, GFP cells also migrated throughout the RMS and reached the OB (Figure 7). However, many cells, albeit with neuronal morphology, remained in the RMS (Figures 7 C, 7D, and 7H). Among the cells that differentiate within the OB, we observed exclusively iNs in the GCL with morphologies of granular neurons (Figures 7 A, 7E, and 7H, 58.9% ± 8%, n = 98 counted cells, n = 2 animals).

Figure 5. Cortical Astroglia Nucleofected with NEUROG2 Integrate as Pyramidal Cell-like iNs In Vivo
(A and B) Examples of GFP+ CtxAstro 20 days post transplantation (dpt) in the postnatal mouse cerebral cortex. Observe the astrocytic morphology of CtxAstro nucleofected with control plasmid (Ctrl) in the cortical layer II (A), and the presence of GFP+ cells with neuronal morphology in animals transplanted with CtxAstro nucleofected with NEUROG2 (B, arrowhead).
(C) Quantification of cells showing astroglial or neuronal morphology 20 days after transplantation. ***p < 0.001.
(D–H) Examples of GFP+ pyramidal-like iNs observed in the cerebral cortex following transplantation of CtxAstro nucleofected with NEUROG2. Note the typical pyramidal cell morphology of iNs in layers II and III of the host cerebral cortex (D, dashed boxes). (E and F) Magnification of dashed boxes shown in (D) revealing the apical dendrite (white arrowheads), basal dendrites (white arrows), and axonal process of iNs (yellow arrowheads). (G) Example of iN in layer II of the cerebral cortex showing spiny dendrites (dashed box magnified in H), suggestive of a glutamatergic identity.
(I) Quantification of spiny and non-spiny iNs in the cerebral cortex of host animals. **p < 0.01.
(J–L) GFP+ IN 20 dpt expressing the mature neuronal marker NEUN (J and K, white arrows).
(M–O) GFP+ IN 20 dpt expressing the transcription factor CUX1 (M–N, white arrowheads). Nuclei are stained with either DAPI (blue) or TO-PRO3 (red).
Statistical test in (C) and (I): Student’s t test (mean ± SEM). n = 5 animals/condition. WM, white matter. Scale bars, 50 μm. See also Figure S7.
Some of these cells expressed GABA (Figures 7F and 7G). Still different from transplantation in the postnatal SVZ, we did not detect GFP+ cells with astroglial morphology in the OB following transplantation of CtxAstro in the adult SVZ.

Altogether, these data suggest that the neurogenic environment in the SVZ plays an instructive role in the phenotypic specification of iNs. However, the origin of astroglial cells and the TF used for reprogramming interfere with the final fate of iNs.
DISCUSSION

Here we show that, similar to their CtxAstro counterparts, postnatal CerebAstro can be efficiently lineage reprogrammed into functional iNs using a single TF. More importantly, we reveal that astroglial cells isolated from different regions and reprogrammed by overexpression of NEUROG2 or ASCL1 generate iNs with different morphological and neurochemical phenotypes. Finally, we demonstrate that integration of iNs after transplantation depends on several factors such as the origin of astroglia population, the TF used, the region of transplant, and the age of the transplanted animal.

Studies focused on astrogial cells reprogramming into neurons exclusively used CtxAstro as starting cell population (Berninger et al., 2007; Heinrich et al., 2010). However, it remained unclear whether astroglial cells isolated from other regions of the CNS could be reprogrammed into iNs using the same TFs. Here, we show that postnatal cerebellum astroglia can be reprogrammed into neurons following overexpression of NEUROG2 or ASCL1 at rates similar to those observed in postnatal cortical astroglia. This study therefore extended neurogenic potential to cerebellum astroglia, suggesting that neurogenic potential may be a hallmark of every astroglia population of the CNS. We chose to use cerebellum because NEUROG2 and ASCL1 lineages contribute different neuronal phenotypes compared with the cerebral cortex.

In fact, ASCL1 and NEUROG2 are TFs belonging to the basic helix-loop-helix family and expressed in different regions of the developing CNS. In the telencephalon, ASCL1 is mostly expressed by progenitors in the ganglionic eminences and contributes to the generation of cortical GABAergic neurons, whereas NEUROG2 is mostly expressed by dorsal progenitors that generate glutamatergic neurons (Fode et al., 2000; Parras et al., 2002; Schuurmans...
These developmental roles of TFs have been suggested to explain the phenotype of lineage-reprogrammed CtxAstro iNs (Heinrich et al., 2010). However, the very same TFs are expressed by progenitors contributing inhibitory interneurons and Purkinje cells of the cerebellar cortex (Zordan et al., 2008; Kim et al., 2008).

According to these roles in the developing cerebellum, we show that ASCL1 and NEUROG2 reprogram CerebAstro mostly into GABAergic iNs. Moreover, expression of calcium binding proteins reveals that ASCL1 and NEUROG2 lineage-reprogrammed CerebAstro iNs are distinct: while ASCL1 induces mostly CALBINDIN expression, NEUROG2 induces PARVALBUMIN. These observations are in line with previous suggestions of these TFs instructing distinct neuronal phenotypes in the developing cerebellum (Zordan et al., 2008). Moreover, we observed that most iNs displaying complex morphologies after ASCL1 also expressed CALBINDIN and CTIP2, which are typical hallmarks of Purkinje cells.

Electrophysiological recordings of CerebAstro-derived iNs indicate similarities and possible differences to CtxAstro-derived iNs. In fact, resting membrane potential, action potential amplitudes, and input resistance of CerebAstro-derived iNs are similar to values previously reported to CtxAstro-derived iNs (Berninger et al., 2007). However, ASCL1 expression in CerebAstro induces the generation of 30% regular spiking iNs, whereas no regular spiking iNs could be observed in CtxAstro reprogrammed with ASCL1 and Dlx2 (Heinrich et al., 2010). Future experiments should systematically compare the electrophysiological properties of iNs generated from distinct astroglia, reprogrammed with different TFs.

Considering that ASCL1 and NEUROG2 reprogram CtxAstro into iNs adopting mostly a GABAergic or glutamatergic phenotype, respectively (Heinrich et al., 2010 and our own data), which is reminiscent of the roles of these TFs in the developing telencephalon, a parsimonious explanation for these data is that astroglial cells retain a molecular memory of the region from where they were isolated. In fact, corroborating this idea, many recent data indicate that reprogrammed somatic cells retain residual DNA methylation signatures characteristic of their somatic tissue of origin. These are called “memory” of origin and indeed favor their differentiation toward lineages related to the donor cells (Hu et al., 2010; Kim et al., 2010; Polo et al., 2010; Tian et al., 2011). Astroglial cells from separate regions of the CNS may present different chromatin modifications in genes targeted by neurogenic TFs. These modifications are likely to occur in early progenitor cells, under influence of distinct morphogenetic signals at different domains of the developing CNS (Kiecker and Lumsden, 2005; Lupo et al., 2006), before generation of neurons and glial cells. This patterning contributes to generate neuronal diversity but would also be inherited by astroglial cells within the same lineage (Costa et al., 2009; Gao et al., 2014). Alternatively, astroglial cells obtained from different regions could express different sets of microRNAs or long non-coding RNAs involved in the specification of neuronal fates (Flynn and Chang, 2014; Jönsson et al., 2015). Future experiments should help to elucidate the exact molecular machinery controlling the acquisition of neuronal phenotypes during lineage reprogramming. It will also be interesting to test whether astroglial cell types isolated from other CNS regions, such as the spinal cord and retina, generate iNs phenotypically similar to neurons of these regions.

In accordance with our observations in vitro, iNs derived from NEUROG2 lineage-reprogrammed CtxAstro and transplanted in the postnatal cerebral cortex mostly adopted a phenotype of pyramidal spiny neurons, which are glutamatergic in this region (Shepherd, 2003). Similarly, previous data in the literature have shown that NEUROD1, a downstream target of NEUROG2, converts cerebral cortex reactive astrocytes into TBR1+ iNs in situ (Guo et al., 2014). However, iNs morphologies described here are much more elaborate, showing typical apical and basal dendrites as well as long-distance axonal projections. One possible explanation for this thorough differentiation of iNs could be that NEUROG2 targets genes important for morphological maturation of cortical pyramidal cells that are not regulated by NEUROD1. In fact, it has been shown that phosphorylation of a single tyrosine residue (T241) of NEUROG2 is necessary and sufficient to control radial migration, neuronal polarity and dendritic morphology of pyramidal neurons (Hand et al., 2005). We assume that NEUROG2 phosphorylation happens in grafted iNs within the host-developing cortex and therefore permits the development of a mature pyramidal morphology. In contrast, however, CerebAstro expressing NEUROG2 and transplanted in the postnatal cerebral cortex differentiate into a very small number of iNs with GABAergic interneuron phenotypes. Thus, NEUROG2 expression alone is not sufficient to reprogram all astroglial populations into pyramidal-like iNs in vivo. Finally, we found very few cells with GABAergic interneuron-like morphologies in the cerebral cortex of animals transplanted with cortical or cerebellum or cerebral cortex astroglia nucleofected with ASCL1, further supporting the notion that both the origin of the astroglial cell and the TF used for reprogramming are important in determining the final fate of iNs in vivo.

The environment in the postnatal cerebral cortical may not be permissive for all lineage-reprogrammed astroglial cells to differentiate into iNs. In fact, after transplantation in the neurogenic subventricular zone, we observed that both cerebellar and cortical astroglia nucleofected with ASCL1 could migrate throughout the RMS and differentiate in the OB as GCL- and PGL-like interneurons. This suggests that the milieu in the postnatal SVZ is not only more permissive to lineage-reprogrammed astroglia iNs, but also plays instructive roles in...
Regardless of the plasmid used for transfection (both cortical and cerebellar astroglia at postnatal stages, Sohn et al., 2015), there is evidence that astrocytes in the RMS and OB of all animals transplanted with Neurog2 contribute to the generation of a very small proportion of juxtaglomerular neurons (Wippenny et al., 2011).

Of note, we observed GFP+ cells with astrocytic morphologies in the RMS and OB of all animals transplanted with both cortical and cerebellar astroglia at postnatal stages, regardless of the plasmid used for transfection (control, Neurog2-, or Ascl1-DsRed). Most of these cells did not express dsRed (non-transfected cell) and some did (transfected but not reprogrammed). This observation suggests that transplanted astrocytic cells can also respond to migration cues in the SVZ-RMS-OB system and integrate in the OB. Accordingly, it has been recently shown that astrocytes are constantly added to the OB after generation in the SVZ (Sohn et al., 2015).

Finally, we also observed that integration in the adult brain is more limited than in the postnatal brain. In fact, we could not observe integration of iNs after transplantation in the adult cerebral cortex (data not shown). In the adult OB we could detect some iNs following transplantation of cortical astroglia transfected with ASCL1. Interestingly, an important fraction of iNs in the adult OB seems to have a problem in reaching the GCL.

Altogether, our results indicate that lineage reprogramming of astroglial cells into neurons by neurogenic TFs is more complex than previously thought. In fact, they show that a same TF can induce the generation of glutamatergic or GABAergic iNs, which would have a completely different role in a neuronal circuitry. Moreover, they demonstrate that both the origin and the region of integration play an important role in the phenotypic specification of iNs. These results have critical relevance for future cell-based therapies, using either transplantation of exogenous lineage-reprogrammed astroglial cells or direct in situ lineage reprogramming of resident astroglia.

**EXPERIMENTAL PROCEDURES**

**Astroglia Culture and Nucleofection**

Postnatal CerebAstro and CtxAstro were isolated from mice on P5–P7. Cerebral cortex gray matter and entire cerebellum were removed and mechanically dissociated. Tissues from both regions were plated separately in culture flasks containing Astromedium (see **Supplemental Experimental Procedures**). After 3–4 days, medium was replaced with fresh Astromedium. After confluence, astroglial cells were nucleofected with pCAG-Neurog2-IRES-DsRed, pCAG-Ascl1-IRES-DsRed, or the control plasmid pCAG-IRES-DsRed using 4D nucleofector (LONZA) (see **Supplemental Experimental Procedures**). Next, cells were plated at densities from 7 × 10^4 to 1 × 10^5 cells/well on poly-D-lysine-coated 24-well tissue plates containing serum-free differentiation medium. For some experiments, primary cells isolated from the neonatal brain were co-cultured at 5 days post nucleofection (see **Supplemental Experimental Procedures**).

**Cell Transplantation**

CerebAstro and CtxAstro were isolated from postnatal GFP animals and cultured as described above. After nucleofection, cells were counted, suspended in serum-free DMEM-F12 (Gibco) at 3–5 × 10^5 cells/µL, and maintained on ice until transplantation procedure. One microliter of cell suspension was gently injected using a pulled glass capillary coupled to a manual injector in the cerebral cortex or SVZ of P0–P2 C57BL/6 mice anesthetized by hypothermia. Transplantation in young adults (P30-P60) C57BL/6 mice were performed under isoflurane anesthesia. Cells were injected using a nanoinjector (Nanoliter 2010, WPI) coupled to a glass capillary using the following stereotactic coordinates (in mm): SVZ (anteroposterior [AP], 0.6; mediolateral [MV], 1.2; dorsoventral [DV], 1.8) and cortex (AP, 1.58; ML, 3.44; DV, 1.40).

**Quantifications and Statistical Analysis**

Quantification of neuronal reprogramming and iNs phenotype in vitro was performed in at least three independent batches of cell culture. For the transplantation studies in the postnatal brain, we analyzed three to five animals for each condition (type of astroglial cell, TF used, and region of grafting). Total number of cells analyzed is described throughout the text (see also **Supplemental Experimental Procedures**). Statistical tests were performed using GraphPad Prism version 5.00 for Windows (www.graphpad.com). Confidence interval is 95%. Statistical significance is indicated in the figures as follows: *p < 0.05; **p < 0.01; ***p < 0.001.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes Supplemental Experimental Procedures, seven figures, one table, and four movies and can be found with this article online at http://dx.doi.org/10.1016/j.stemcr.2017.05.009.

**AUTHOR CONTRIBUTIONS**

M.C. and M.R.C. designed the work, collected and analyzed data, and wrote the article. A.R.M.F. and D.M.S.M. performed video-microscopy time-lapse and fate-mapping experiments. M.M.H. and R.N.L. performed electrophysiological recordings. T.S. provided the software for acquisition and analysis of time-lapse video microscopy experiments. All authors read and approved the final version of the article.
ACKNOWLEDGMENTS
This work was supported by CNPq (MRC 473254/2013-1 and 466959/2014-1) and CAPES fellowships to M.C. and D.M. We thank Dr. Benedikt Berninger for providing the plasmids encoding for pronuclear factors and for helpful discussion of the manuscript. We also thank Dr. Cecilia Hedin Pereira, Dr. Cláudio Queiroz, Dra. Katarina Leão, and Dr. Ricardo Reis for valuable discussion of the work, and Rebecca Diniz, Ana Cristina, and Giovanna Andrade for technical assistance.

Received: April 19, 2016
Revised: May 8, 2017
Accepted: May 9, 2017
Published: June 8, 2017

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