Embryonic stem cells assume a primitive neural stem cell fate in the absence of extrinsic influences

Simon R. Smukler, Susan B. Runciman, Shunbin Xu, and Derek van der Kooy

Department of Medical Genetics and Microbiology, University of Toronto, Toronto, Ontario, M5S 1A8, Canada

The mechanisms governing the emergence of the earliest mammalian neural cells during development remain incompletely characterized. A default mechanism has been suggested to underlie neural fate acquisition; however, an instructive process has also been proposed. We used mouse embryonic stem (ES) cells to explore the fundamental issue of how an uncommitted, pluripotent mammalian cell will self-organize in the absence of extrinsic signals and what cellular fate will result. To assess this default state, ES cells were placed in conditions that minimize external influences. Individual ES cells were found to rapidly transition directly into neural cells, a process shown to be independent of suggested instructive factors (e.g., fibroblast growth factors). Further, we provide evidence that the default neural identity is that of a primitive neural stem cell (NSC). The exiguous conditions used to reveal the default state were found to present primitive NSCs with a survival challenge (limiting their persistence and proliferation), which could be mitigated by survival factors or genetic interference with apoptosis.

Introduction

The emergence of the earliest neural cells during mammalian development and the mechanisms that govern this process remain incompletely characterized. Such cells are likely to be neural precursors or stem cells, though the ontology of the neural stem cell (NSC), which can be isolated from embryonic and adult forebrain (Weiss et al., 1996; Gage, 2000), has not been fully elucidated. During development, neural cells arise from the ectodermal germ layer, which also produces epidermis. According to the classical model of this process, conceptualized largely from amphibian embryology studies, nascent embryonic ectoderm receives a positive signal from a specialized group of dorsal mesodermal cells, termed the organizer, which instructs the adjacent ectodermal cells to adopt a neural fate (Harland and Gerhart, 1997; Weinstein and Hemmati-Brivanlou, 1999; Spemann and Mangold, 2001). The structural equivalent of the organizer in amniotes is the node. It was thought that the ectodermal germ layer, which also produces epidermis.

Abbreviations used in this paper: aif, apoptosis-inducing factor; apaf1, apoptotic protease activating factor 1; BMP, bone morphogenic protein; cas9, caspase 9; cGMP, cyclic guanosine monophosphate; E, embryonic day; EB, embryoid body; ES, embryonic stem; ICM, inner cell mass; LiF, leukemia inhibitory factor; NAC, N-acetylcysteine; NFM, neurofilament-M; NS, neural stem cell; NSC, neural stem cell; pCPT, 8-(4-chlorophenylthio).

The online version of this article contains supplemental material.
These findings led to the development of the currently more widely accepted model, the default model, which states that each individual ectodermal cell has an intrinsic default program to become a neural cell (Munoz-Sanjuan and Brivanlou, 2002). In the context of the intact embryo, this default program is being actively suppressed by ubiquitously expressed BMPs. Thus, the organizer tissue does not provide a positive inductive signal but rather secretes factors that antagonize BMP signaling, thereby disinhibiting the default neural program in proximal ectodermal cells.

Several subsequent studies have challenged the default model of neural fate acquisition. For example, experiments in chicks have suggested that BMP inhibition may not be sufficient to induce neuralization (Streit et al., 2000; Linker and Stern, 2004). However, it is uncertain how complete the BMP inhibition was in these studies, and it is possible that the activity of individual BMPs was insufficiently suppressed to allow neuralization to occur (and/or that some BMP subtypes or other neural inhibitors escaped blockade). It has also been suggested that other factors, such as FGF and Wnt signaling, are involved with neural specification in several vertebrates (Baker et al., 1999; Streit et al., 2000; Wilson et al., 2000, 2001), though whether they are required for the initial neural fate change or for the later expansion of this neural population is currently unresolved. Further, their mechanism of action may be through modulation of BMP gene transcription (Bainter et al., 2001), consistent with a model of BMP inhibition–mediated neuralization.

There are currently few published studies examining the neural default model in mammalian cells, and there is controversy over whether such a default neural mechanism exists in mammals. In an effort to determine whether a default mechanism underlies neural fate specification from uncommitted mammalian precursors, we undertook studies using mouse embryonic stem (ES) cells, which are derived from the inner cell mass (ICM) of the blastocyst-stage embryo and represent a model of the earliest pluripotent mammalian cell (Evans and Kaufman, 1981; Martin, 1981). ES cells are capable of generating entire viable mice in vivo (Nagy et al., 1993) and are able to produce most, if not all, cell types in vitro (Beddington and Robertson, 1989; Rathjen and Rathjen, 2001; Wobus, 2001). Use of ES cells to investigate neural determination can potentially provide many insights into the developmental process. Importantly, though, their use in assessing a default fate model of neural fate acquisition is that single dissociated R1 ES cells were plated at low cell densities (≤10 cells/μl; 2,600 cells/cm²) in chemically defined serum- and growth factor–free media. As we reported previously (Tropepe et al., 2001), ES cells placed in these minimal conditions rapidly acquired a neural identity, with >90% of cells expressing nestin, an intermediate filament protein associated with neural precursors (Lendahl et al., 1990), within 4 h (Fig. 1 A). The neural precursor identity of these cells was supported further by their expression of Sox1, one of the earliest transcription factors expressed in cells committed to the neural fate (Pevny et al., 1998; Fig. 1 C). Undifferentiated ES cell colonies did not exhibit expression of these markers (Fig. 1, B and D). At 4 h, no significant cell mortality was observed and <1% of the plated cells had proliferated, indicating that single ES cells began a direct transition to neural cells, without requirement for cell division. After an additional 20 h, most (77.5 ± 1.3%) of these ES-derived neural cells did not survive, as these minimal conditions were not very supportive. However, 99.3 ± 0.2% of the remaining viable cells expressed nestin and Sox1 and maintained some expression of Oct4, a transcription factor expressed in ES cells (Nichols et al., 1998; Fig. 1, E and F). To verify that the observed up-regulation of nestin was not a nonspecific stress response, STO fibroblasts were placed in identical conditions; however,
RT-PCR and immunostaining did not show any nestin expression either before or after culture in minimal media (unpublished data).

Further, 24 h after cell plating, 90.4 ± 1.7% of the surviving cells expressed neurofilament-M (NFM) and 47.5 ± 3.6% expressed the early neuronal marker β3-tubulin (Fig. 1, G and H). RT-PCR analysis confirmed expression of nestin and Sox1 (Fig. 1 I). Additional confirmation of neural lineage commitment after 24 h was evidenced by the rapid down-regulation of brachyury and GATA-1 (mesodermal markers), as well as HNF3β, HNF4, and GATA-4 (endodermal markers), which were detectable in ES cells (Fig. 1 J). At 3 d, most surviving cells maintained a neural precursor identity, expressing nestin and Sox1 (Fig. 1, K and L). Cells with more elaborate neural morphologies were also evident, with the observation of neuronal and glial cells (Fig. 1, M and N). Cells with very advanced morphology, and positive for a neural cell subtype marker, typically displayed down-regulation of nestin expression (faint or absent immunostaining) and were Oct4− (unpublished data). By 7 d, viable cells were not observed, indicating that the exiguous nature of the media conditions was not supportive enough for the maintained survival of the neural cells. These data indicate that in the absence of extrinsic signals, ES cells rapidly begin to acquire a neural precursor identity, consistent with a default mechanism for an ES cell to transition directly into a neural cell.

Figure 1. ES cells rapidly transition into neural cells when placed in minimal conditions. (A and C) Immunocytochemical labeling showed that ES cells placed in minimal media conditions (for 4 h) initiated pronounced expression of the neural precursor markers nestin (A) and Sox1 (C). The nuclear stain DAPI was used to identify all cell nuclei within the field. (B and D) Undifferentiated ES cells (growing on a fibroblast feeder layer) exhibited Oct4 (B) but not nestin or Sox1 (D). The large Oct4+ nuclei are feeder cells. (E) By 24 h, almost all cells expressed nestin and maintained some nuclear Oct4 expression and Sox1 (F). (G and H) Most of the ES-derived neural cells express NFM (G), and many express the early neuronal marker β3-tubulin (H; at 24 h). (I) Expression of nestin and Sox1 was confirmed by RT-PCR analysis of ES-derived neural cells (at 24 h). (J) Brachyury and GATA-1 (mesodermal markers), as well as HNF3β, HNF4, and GATA-4 (endodermal markers), were detected by RT-PCR in undifferentiated ES cells, though they were rapidly down-regulated within 24 h in minimal conditions. (K and L) After 3 d, most surviving ES-derived neural cells retain a neural precursor identity, maintaining expression of nestin (K) and Sox1 (L). (M and N) Differentiating cells with more elaborate morphologies were also apparent as β3-tubulin+ neurons (M) and O4+ oligodendrocytes (N). Bars: (A–D and K–N) 25 μm; (E–H) 10 μm.
**ES cell neuralization occurs by default without requirement for any exogenous instructive factors**

Is this neural transition truly occurring by default? A protein component of the media formulation, transferrin, has been suggested to be required for ES cell neural fate initiation (Ying et al., 2003). We show that nestin and Sox1 expression was established after 4 h, even when ES cells were cultured in PBS alone, ruling out any requirement for media components in an instructive capacity (Fig. 2, A and B). After a 24-h culture period in PBS, very few cells remained viable, though they all expressed nestin and Sox1 (Fig. 2, C and D). The fragmented nuclei represent dead cells, which typically did not display immunoreactivity for the neural markers.

Though not exogenously required, autogenously produced FGFs have been proposed to be essential for ES cell neuralization, suggesting an instructive process (Ying et al., 2003). To the contrary, pharmacological inhibition of FGF signaling using the FGF receptor kinase antagonist SU5402 (5 μM) did not prevent the rapid acquisition of neural markers by ES cells placed in minimal conditions for 24 h, with expression of nestin (E), Sox1 (F), NFM (G), and β3-tubulin (H) observed. [E–H] Similarly, ES cells harboring deletion of the fgrf1 gene displayed the typically observed neural markers by 24 h, expressing nestin (I), Sox1 (J), NFM (K), and β3-tubulin (L). Bars, 25 μm.

**Primitive NSCs emerge from the default neural pathway**

What is the nature of the neural cells emerging from this default pathway? The early expression of nestin and Sox1 indicates that they are neural precursors. However, these markers are expressed not only in true NSCs but also in more restricted neural progenitor cells. Our previous work introduced a novel neural precursor arising under these conditions, the primitive NSC (Tropepe et al., 2001), identified with an assay analogous to the neurosphere (NS)-formation assay used to identify definitive forebrain-derived NSCs. When the cytokine LIF was included in our minimal condition assay, a very small percentage (0.18 ± 0.01%) of the initially plated ES cells became neural cells that proliferated over 7 d to form clonally derived floating sphere colonies, termed primitive NSs (Fig. 3 A). LIF was not required for early ES cell neuralization, as it could be added after 4 h without any decrement in primitive NS production (101 ± 4% of control). Rather, LIF appeared to prevent further differentiation down the neural lineage and thereby maintained these neural cells in an undifferentiated, proliferative state. This interpretation is supported by the findings that LIF addition after 2 d did not enable any primitive NS generation (unpublished data), indicating that there was an LIF-responsive temporal window, after which the neural cells had progressed to a non–LIF-responsive, unproliferative, and more differentiated state. These primitive NSs expressed both nestin and Sox1 (Fig. 3, B and C), similar to NSs generated by definitive NSCs, indicating that they are composed of neural precursors. The primitive NS cells also maintained some expression of Oct4 (by immunostaining and RT-PCR), as well as nanog (by RT-PCR), another transcription factor expressed by ES cells (Chambers et al., 2003; unpublished data). The neural precursor character of the primitive NSs was confirmed by expanded RT-PCR gene expression analysis. Various additional neural genes were expressed by primitive NSs, including Sox2, Sox3, Neurogenin1, NeuroD, Pax6, Nkx2.2, Mash1, Musashi-1, Otx2, and HoxB1, strongly supporting their neural nature (Fig. S1, available at http://www.jcb.org/cgi/content/full/jcb.200508085/DC1). Primitive NSs did not express CK-17 (an epidermal marker), nor did they express the mesodermal markers brachyury and GATA-1 or the endodermal markers HNF3β, HNF4, and Pax4. These multi–germ layer markers are typically found in EBs (Wassarman and Keller, 2003), and their absence further verifies that the primitive NS cells are specified to the neural lineage. Transcripts for most of these genes were found at detectable levels in undifferentiated ES cells, suggesting that there may be widespread, low-level, promiscuous gene expression in the uncommitted state and that...
differentiation may involve the down-regulation of certain genetic programs with the maintenance and up-regulation of others. The primitive NS cells could be maintained in an undifferentiated state by subcloning/passaging in serum-free conditions to produce secondary, tertiary, and successive spheres. Interestingly, secondary and subsequent sphere formation was dependent on the addition of exogenous FGF2, similar to the FGF2-dependent proliferation of forebrain-derived NSCs. When transferred to conditions promoting differentiation, individual primitive NSs differentiated to produce neurons, astrocytes, and oligodendrocytes (Fig. 3 D), the three neural lineage cell types. Cells negative for differentiation markers maintained a neural precursor identity, expressing nestin and Sox1 (Fig. 3 E). Differentiation of the passaged spheres was similar to that of the primary primitive NSs (unpublished data). Though the cells present after in vitro differentiation of primitive NSs appeared limited to differentiated and immature/precur sor neural cells, the maintained expression of some ES-like genes (Oct4 and nanog) in the undifferentiated primitive NS state suggested that cells constituting the primitive NS were not yet absolutely committed to the neural lineage. This is supported by our previous finding that primary primitive NSs could still manifest broader lineage potential when placed in the appropriate developmental environment, i.e., when used to generate morula-aggregation embryo chimeras (Tropepe et al., 2001). However, when the primitive NSs were passaged to produce secondary spheres (with FGF2 and without LIF), efficient chimera production was not observed (similar to our results obtained using NSs from forebrain-derived adult NSCs), suggesting that the secondary spheres were now fated exclusively to the neural lineage (Tropepe et al., 2001; unpublished data). Further, expression of Oct4 and nanog was not found by RT-PCR in the secondary spheres (unpublished data). Thus, a novel neural precursor, the primitive NSC, arose in these default conditions.

**Primitive NSC proliferation is dependent on autogenously produced FGF signaling**

Though not required for early default neural fate acquisition, autogenously produced FGFs were found to be necessary for primitive NS formation. Inclusion of 5 μM SU5402 in the primitive NS assay virtually eliminated primitive NS generation (reduced by 98 ± 1%; Fig. 4 A), and fgfr1+/− ES cells displayed dramatically diminished (by 91 ± 1%) primitive NS production (Fig. 4 B). This suggests that autogenous FGF signaling was important for the proliferation and/or survival of primitive NSCs. The proliferation interpretation is supported by results obtained from rescue experiments. When SU5402 was included in the primitive NS assay, minimal proliferation was observed by 3 d. If the drug was then removed and the assay was continued for 7 d, primitive NSC proliferation resumed and there was a large recovery of primitive NS formation (Fig. 4 C). Similarly, if the drug was maintained for a full 7 d and then removed, a substantial recovery of primitive NS formation was observed (Fig. 4 C). Further, if SU5402 was added on day 3 of the assay (at which point the cells have established their neural transition and primitive NSCs are proliferating), proliferation was impaired and there were consequently fewer primitive NSs observed after 7 d (Fig. 4 D). As FGF signaling was dispensable for the default neural fate switch in the short-term (4 and 24 h) experiments discussed previously, this suggests that primitive NSCs are still formed in the transition from ES cells, though their proliferation to form primitive NSs is dependent on signaling by autogenously produced FGFs. Sonic Hedgehog signaling, reported to be important in the regulation of some neural progenitors (Zhu et al., 1999), was found to be dispensable, as Sonic Hedgehog inhibition with cyclopamine had no inhibitory effect on primitive NS formation (unpublished data).

**Cell mortality limits the number of default pathway-derived primitive NSCs that form primitive NSs**

Given the low frequency of ES-derived neural precursors (~0.2%) that proliferate to form primitive NSs (which is our assay for the presence of a primitive NSC), the question remains as to whether the default fate pathway specifically gives rise to primitive NSCs or simply to a population of generalized and perhaps heterogeneous cells of the neural lineage. Though neural in nature, it is possible that this ES-derived population contains a very small subset of bona fide primitive NSCs, capable of long-term passaging and production of multilineage

![Figure 3. Primitive NSCs emerge from the ES cell default neural pathway. (A) When LIF was included in the minimal conditions, a small number of ES-derived neural precursor cells proliferated over 7 d to form clonally derived floating sphere colonies, termed primitive NSs. The primitive NS-initiating cell was termed a primitive NSC. (B) The primitive NS cells were found to coexpress nestin and Sox1 by immunocytochemical labeling, demonstrating their neural precursor identity. (C) RT-PCR analysis confirmed expression of nestin and Sox1 within primitive NSs (pNS). (D and E) When differentiated, primitive NSs yielded β-tubulin+ neurons, GFAP+ astrocytes, and O4+ oligodendrocytes (D), whereas most cells retained a neural precursor phenotype, maintaining expression of nestin and Sox1 (E). Bars: (A) 100 μm; (B–D) 50 μm.
Figure 4. Primitive NSC proliferation is dependent on autogenously produced FGF signaling. (A) Pharmacological inhibition of FGF signaling throughout the primitive NS assay using SU5402 (5 μM) virtually abolished primitive NS formation (pNS). *, P < 0.001 versus control. (B) ES cells with a deletion of the fgfr1 gene displayed drastically reduced primitive NS production. *, P < 0.001 versus wild type [wt]. (C) When SU5402 was included for 3 or 7 d of the primitive NS assay and then removed, the primitive NSCs resumed proliferation, and a substantial recovery of primitive NS formation was observed over an additional 7 d. *, P < 0.001 versus condition with maintained SU5402 (not depicted). (D) When SU5402 was added on day 3 of the primitive NS assay, proliferation of already proliferating primitive NSCs was impaired and fewer primitive NSs ensued. *, P < 0.001 versus control.

progeny, whereas the remainder may represent more restricted or committed neural cells. If the default pathway does specifically give rise to primitive NSCs, there must be some inhibitory influences preventing the exhibition of the primitive NS phenotype, i.e., primitive NS generation, in all of the individual neural derivatives of ES cells.

The early nestin+ neural cells present after 4 h in minimal conditions subsequently underwent a survival challenge because of the exiguous nature of the media, resulting in extensive cell mortality (77.5 ± 1.3% by 24 h). In the absence of LIF, mortality progressively increased over 7 d, until no viable cells remained. In the presence of LIF, a few of the neural cells retained a proliferative character, surviving and propagating to form primitive NSs, though the vast majority of the early nestin+ cells still died. Accordingly, primitive NSCs may have been produced but did not form primitive NSs because of the early survival challenge that they experienced. Thus, increasing cell viability should enhance primitive NS production. The survival factor N-acetyl-l-cysteine (NAC), which has multiple modes of action but is thought to act primarily through its antioxidant effects (Zafarullah et al., 2003), promotes survival of various neural cell types (Mayer and Noble, 1994; Ferrari et al., 1995; Yan et al., 1995). Inclusion of NAC dose-dependently increased primitive NS formation (up to ~35-fold), with toxic effects observed at very high concentrations (Fig. 5 A). We also explored the cAMP–protein kinase A pathway as a more physiological modulator of cell survival. cAMP signaling has been shown to enhance cell survival in several different neural cell systems (Rolletschek et al., 2001; Bok et al., 2003; Lara et al., 2003). Exogenous application of a membrane-permeable cAMP analogue, 8-(4-chlorophenylthio) (pCPT)-cAMP, was able to dose-dependently augment primitive NS formation (up to ~25-fold), with toxic effects observed at exceedingly supraphysiological concentrations (Fig. 5 A). Concentrations at which peak effects were observed were used in all subsequent experiments (1 mM NAC and 100 μM pCPT-cAMP). When added simultaneously, the effects of NAC and pCPT-cAMP were additive, if not synergistic, enhancing primitive NS formation (up to ~100-fold) such that ~20% of the initially plated ES cells became primitive NS–forming primitive NSCs (Fig. 5 B). This also suggests that the two compounds have distinct mechanisms of primitive NS survival promotion (leading to subsequent primitive NS formation). RT-PCR analysis of primitive NSs derived in each or both factors demonstrated the expression of the neural precursor markers without detection of the multi–germ layer markers (discussed previously). Addition of NAC and/or pCPT-cAMP after 4 h of the primitive NS assay (at which point the cells have already undertaken their neural transition) revealed no decrement in their ability to augment primitive NS formation (NAC addition at 4 h was 98 ± 3% of NAC added from start, and pCPT-cAMP addition at 4 h was 101 ± 3% of pCPT-cAMP added from start), suggesting that they were not effecting the NS increase through an instructive role in early differentiation. Additionally, this demonstrated that the survival-promoting effects of NAC and cAMP were not directly on ES cells but rather on their nestin+ clonal neural derivatives.

Viability assessment showed that NAC and pCPT-cAMP did indeed increase cell survival under minimal conditions. At 24 h, NAC increased survival of the initial 4-h nestin+ cells from 22.5 ± 1.3 to 44.8 ± 2.3% (P < 0.01), whereas pCPT-cAMP increased survival to 43.6 ± 2.4% (P < 0.01). Though the survival factor–induced increases in viability relative to the control media were observed over the course of the complete 7 d, progressive mortality was still observed. Even with both factors present in the primitive NS assay, there was still 34.4 ± 2.6% mortality for the initial 4-h nestin+ cells after 24 h and 54.3 ± 2.9% mortality by 3 d. There were very few cells that survived past the 7-d assay that were not in a primitive NS, suggesting that only cells that were in the context of a proliferating sphere were effective at long-term survival in these conditions. It is worth noting that although the survival factors greatly increased cell viability at 1 and 3 d in minimal media, the rapid acquisition of neural identity and the frequencies of marker-positive cells (discussed previously and in Fig. 1) were not altered (not depicted). Thus, these results indicate that the primitive NS is the primary identity of cells derived from the default pathway but that cell mortality limits the number that
can survive to form clonal primitive NSs and thereby be detected by our assay.

The ability of NAC and pCPT-cAMP to facilitate primitive NS formation decreased dramatically and progressively when the compounds were added during the derivation of secondary and tertiary spheres (Fig. 5 C). This suggests that the extensive survival challenge that limited primitive NS generation occurred only with the primary ES-derived primitive NSCs and not with the subsequently passaged NSCs, which are similar to forebrain-derived definitive NSCs (which also did not exhibit increases in NS formation with the survival factors; unpublished data). This effect highlights a fundamental difference between primitive NSCs and the passaged, more mature definitive NSCs. When individual primary spheres derived in the presence of pCPT-cAMP were passaged back into control media for secondary sphere formation, they generated 2.3 ± 0.2 times (P < 0.01) more secondary spheres compared with primary spheres that had been derived in control media. Spheres derived in NAC did not display any increased secondary sphere formation when passaged back into control media. This suggests that pCPT-cAMP may have had an additional effect, increasing the number of symmetric divisions that the primitive NS underwent, thereby expanding the stem cell numbers within the primary sphere. Alternately, there may have simply been a prolonged survival enhancement in the pCPT-cAMP-derived primary sphere cells, even when they were placed back into media without pCPT-cAMP. Primitive NSs derived in NAC and/or pCPT-cAMP did not display any notable alterations in differentiation (unpublished data). Pharmacological modulation of the endogenous pathways showed that the cAMP pathway positively regulated and the cyclic guanosine monophosphate (cGMP) pathway negatively regulated primitive NSC survival and primitive NS formation (see online supplemental material).

**Figure 5.** A survival challenge attributable to the minimal culture conditions limits the number of ES-derived neural precursors that survive to form primitive NSs. Extensive cell mortality was observed upon culture in the minimal conditions used for the default state assessment and primitive NS (pNS) assay. (A) Improving cell viability by inclusion of the survival factors NAC and the membrane-permeable cAMP analogue pCPT-cAMP (cAMP) dose-dependently enhanced primitive NS formation. *, P < 0.01 versus control. (B) Concurrent inclusion of 1 mM NAC and 100 μM pCPT-cAMP (a concentration at which peak effects were observed in A) in the primitive NS assay showed that they produced an additive, if not synergistic, effect. *, P < 0.001 versus control; †, P < 0.001 versus control and versus each factor alone. (C) The effectiveness of NAC (1 mM) and pCPT-cAMP (100 μM) in promoting NS formation decreased dramatically with passaging of NSs. *, P < 0.001 versus respective survival factor primary spheres.

**Genetic deletion of apoptotic pathway components enhances primitive NS formation**

To further substantiate that cell mortality limited the number of primitive NSCs that survived to proliferate and form clonal primitive NSs, we used ES cell lines with a survival advantage conferred by mutations in apoptotic signaling pathway components. *Apoptotic protease activating factor 1 (apaf1)* and *caspase 9 (cas9)* are components of the more common apoptotic pathway (Cai et al., 1998), whereas *apoptosis-inducing factor (aif)* is a mediator of an alternate, independent apoptotic pathway (Cande et al., 2002). The *apaf1−/−* (Yoshida et al., 1998), *cas9−/−* (Hakem et al., 1998), and *aif−/Y* (Joza et al., 2001) ES cells displayed substantially increased primitive NS formation relative to wild-type ES cells, with a more pronounced effect observed for the *apaf1* and *cas9* mutants (Fig. 6 A). Heterozygotes for *apaf1* and *cas9* displayed an intermediate effect, indicative of a gene dosage effect (Fig. 6 A). Differentiation of primitive NSs generated by these ES cell lines was similar to wild type (unpublished data). This demonstrates that, similar to the effect of survival factors, survival promotion through interference with apoptosis enhanced primitive NS formation. This also suggested that the primary apoptotic pathway responsible for mediating cell death in these cultures was the *apaf1−cas9* apoptotic pathway, although there was still some contribution from the *aif* pathway.

If the primitive NS–promoting effect of the survival factors NAC and pCPT-cAMP was achieved through a reduction in primitive NSC death mediated by these apoptotic pathways,
then the survival factor effect should be reduced in these mutant ES cell lines. Typical robust stimulation was observed for the wild-type line, which was much reduced, though still considerable, in the aif mutant (Fig. 6 B). Strikingly, the effects of NAC and pCPT-cAMP were drastically reduced in the apaf1−/− and cas9−/− ES cells to the extent that there was no longer any significant effect of NAC and only an approximately twofold stimulation by pCPT-cAMP (Fig. 6 B). This attenuation or occlusion of the survival factor effects in these mutant ES cell lines verifies that NAC and pCPT-cAMP promote primitive NS formation through a reduction in apoptotic cell death, primarily mediated by the more dominant apaf1–cas9 pathway.

**TGFβ inhibition independently cooperates with the survival factors to promote primitive NS production**

We have previously demonstrated that TGFβ-related signaling negatively regulates ES cell neuralization and the basal number of primitive NSs that form in our assay (Tropepe et al., 2001). To determine how survival promotion interacted with interference of the TGFβ pathway, the effects of NAC and pCPT-cAMP were assessed in an ES cell line that contains a mutation in smad4, a key downstream signaling molecule in multiple TGFβ-related pathways (Sirard et al., 1998). The smad4−/− ES cells displayed increased primitive NS generation relative to wild-type ES cells under both control media conditions as well as in the presence of the survival factors such that in the presence of both NAC and pCPT-cAMP ~35% of the initially plated ES cells became primitive NSCs that were able to survive and proliferate to form primitive NSs (Fig. 6 C). These results provide further support for the notion that the default pathway does indeed give rise specifically to primitive NSCs (though TGFβ signaling hinders this default to primitive NSCs), which then experience a separate survival challenge limiting the number of primitive NSCs that form primitive NSs.

**Discussion**

The present study demonstrates that in the absence of extrinsic influencing signals, ES cells will rapidly undergo a direct neural conversion, transitioning into neural precursor cells. This neuralization does not depend on any instructive factors but rather occurs by a default mechanism. The default neural pathway specifically gives rise to primitive NSCs (though TGFβ signaling hinders this default to primitive NSCs), which then experience a separate survival challenge limiting the number of primitive NSCs that form primitive NSs.
mediate their effects largely through abrogation of the Apaf1–Cas9 pathway. In the presence of LIF, the primitive NSC remains in an undifferentiated, proliferative state and generates a primitive NS containing a subpopulation of more mature [FGF2 dependent] definitive NSCs. Upon dissociation of the primitive NS, the definitive NSCs will proliferate in an FGF2-dependent manner to generate an NS. In the presence of FGF2, this self-renewal passaging can be performed repeatedly. With the addition of serum, the NS can be differentiated to produce trophoblasts, astrocytes, and oligodendrocytes.

al., 2004), although clonal analyses were not performed. Other work examining the neural conversion of mouse ES cells without EB formation reported that most cells adopted a neural precursor phenotype in serum-free conditions; however, they found that widespread neuralization did not occur until at least 3 d (Ying et al., 2003). Further, Ying et al. (2003) suggest that the media component transferrin and autocrine FGFs were required positive inductive factors, in contrast to a pure default model. The present results demonstrate that neural specification commences over a matter of hours, even when ES cells were placed in PBS alone, ruling out any necessity for media components in an instructive capacity. Certainly, cells cannot be maintained long-term in PBS alone, as nutritive factors are required in a permissive role. Further, we provide evidence against any positive inductive role for autogenously produced FGFs by demonstrating that neither pharmacological FGF inhibition (using the same inhibitor as Ying et al., 2003) nor genetic deletion of the FGF1–FGF receptor-1 effected any reduction in neuralization. Rather, such FGF signaling functions to allow the proliferation of the default pathway–derived neural precursors. Experiments of Ying et al. (2003) used cell densities up to ~20 times higher than those used in the present studies and in some cases allowed low-density plated ES cells to grow as colonies overnight in normal ES media (increasing the effective cell density) before removal of LIF and serum. These conditions are likely to facilitate the density-dependent accumulation and signaling of secreted neural inhibitors (e.g., BMPs), as we demonstrated previously (Tropepe et al., 2001), potentially explaining why their neuralization was delayed. They also reported the gradual increase in expression of BMP antagonists (e.g., noggin and follistatin) over 1–5 d in their conditions, which would then provide the BMP antagonism necessary to allow the default program to manifest. Furthermore, the extensive viability and exponential cell expansion in their conditions is in stark contrast to our findings, indicating that their high cell densities and/or richer media formulation produced an environment less minimal than that used in the current study.

The severely minimal nature of our culture conditions was necessary to effectively minimize potential extrinsic influences and thereby assess a default state. However, this was clearly responsible for the extensive cell mortality observed. In attempts to evaluate the default state, one encounters a catch-22. At some point, progressive removal of extrinsic factors will dispense with constituents essential for default fated cell survival, and there will be no experimental output to assess. Such factors are required for neuralization not in an instructive capacity but rather in a permissive, supportive one. In interpretation of our results, one might suggest that we were merely providing conditions that selected for the survival of neural cells while noneural cells perished, and thus the default state was not assessed. Two observations argue against this interpretation. First, in the initial 4 h after ES cell plating, no significant mortality was observed, yet the vast majority of individual ES cells had already begun their neural transition. Second, with the use of both survival factors in the 24-h assay, viability was increased from 23 to 65%; however, a multiple marker–based analysis showed that there was no difference in the uniform cell neuralization and that there was no expression of other germ layer markers. If the conditions used were selecting against the survival of nonneural lineage cells, they should have been apparent when survival was so dramatically increased.

The primitive NSCs described here exhibited characteristics (i.e., proliferative sphere formation, gene expression, and differentiation potential) that are analogous to forebrain-derived NSCs. Although it is clear that the primitive NSCs commenced neural lineage specification, it was apparent that they retained vestiges of ES cell features, suggesting that the commitment was not yet absolute. This was evidenced by the broader differentiation potential observed in blastocyst chimera experiments (Tropepe et al., 2001). Passaging of the primitive NSs yielded more committed neural precursors, increasingly similar to NSs formed by forebrain-derived NSCs (e.g., they acquired dependence on exogenous FGF2, ceased expression of ES cell markers Oct4 and nanog, and lost blastocyst chimerism potential). Recent work from our lab has demonstrated that LIF-dependent sphere-forming cells can be isolated directly from the embryonic day (E) 5.5–7.5 mouse epiblast (Hitoshi et al., 2004). These embryo-derived spheres possessed similar characteristics to the ES-derived primitive NSs and were found to give rise to FGF2-dependent, NS-forming, definitive NSCs upon passaging in vitro (Hitoshi et al., 2004). Thus, in vivo there are two distinct sphere-forming NSC populations that are present at different stages of development from E5.5 to adult. In addition, corresponding sphere-forming NSC populations could be sequentially derived from ES cells. This suggests the in vivo relevance of an ES cell–based model describing the ontogeny of neural stem cell fate.
of NSCs (Fig. 7). Pluripotent ES cells derived from the E3.5 ICM will commence a direct default transition in the absence of inhibitory influences (e.g., TGF-β-related signaling) to yield LIF-dependent primitive NSCs, similar to those that can be isolated from the epiblast (Hitoshi et al., 2004). The primitive NSs produced can be passaged to give rise to definitive, FGF2-dependent NSCs, analogous to the FGF2-dependent NSCs that can be isolated from the E8.5 neural plate (Tropepe et al., 1999). The in vitro-derived definitive NSCs can be extensively passaged to demonstrate long-term self-renewal, consistent with the effective isolation of definitive NSCs from the adult remnant of the embryonic brain germinal zone throughout the lifetime of the animal.

Is the propensity of ES cells to default to a neural identity developmentally relevant? ES cells have properties similar to the E3.5 ICM cells from which they are derived. Our data suggest that the ES cells can directly become neural without a discernible ectodermal intermediate phase. The developmental dogma states that neural tissue arises from the ectodermal germ layer, which is not defined until gastrulation (E6.5–8.5). ES cell differentiation has been suggested to recapitulate these stages (Pelton et al., 2002). It is possible that the neural tendency of the E3.5 cells never has the opportunity to manifest in vivo because of extensive antineural signaling via BMPs until after gastrulation, when node regions are formed to disinhibit the neural fate in what is now specified ectoderm. However, recent work in both Xenopus laevis and chicks has suggested that there may be some neural-fate acquisition even before gastrulation (Streit et al., 2000; Gamse and Sive, 2001). This may be attributable to preectodermal ICM-like cells that begin to express their default neural tendency after ineffective neural inhibition. In the context of the intact embryo, even with its milieu of secreted BMPs, it is conceivable that transient interruption or inefficiency in BMP delivery or activity would occur, allowing a cell or cells to escape neural inhibition and rapidly initiate the default neural program. In addition, the present data suggest that mouse ES cells can transition directly into neuronal cells; however, neurons are not produced in vivo until approximately E10. It is possible that neural precursors present in vivo before this stage are competent to form neurons but do not do so because of the presence of factors like LIF, which have been shown to prevent neural precursors from further differentiation down the neural lineage and maintain the undifferentiated primitive NSC state.

In addition to providing insights into the mechanisms of neural development, default fate studies using ES cells allow exploration of what is perhaps a more basic and fundamental issue, i.e., how an uncommitted, pluripotent mammalian cell will self-organize in the absence of extrinsic instructive or inhibitory signals and what cellular configuration/fate will result. Our data suggest that such uncommitted cells tend to self-organize in the configuration of a NSC, i.e., the default fate. What is meant by the term default? Clearly, such analyses begin with the definition of a system, and the default state represents the configuration that the system will autonomously acquire in the absence of extrinsic inputs to the system. However, this means that there can be different levels of analysis depending on the definition of the initial system. In the context of ES cell default neural fate, a fairly gross level of analysis would simply state that no other tissues or cells are required for an ES cell to become a neural cell, whereas all other factors (such as environmental/media components and endogenous/autocrine signaling) are included within the system. The present data would clearly satisfy this level of analysis. A finer level of analysis would state that a single ES cell would acquire a neural identity with only permissive molecules present (in the media) and without autocrine inductive signals. The present data seem to partially satisfy this level of analysis as well, as we have ruled out requirement for exogenously supplied instructive molecules and autogenous FGF signaling. However, we cannot yet definitively demonstrate that no other potential instructive endogenously supplied (and perhaps autocrinely acting) factors are present. An even finer level of analysis would be at the genomic level, with the system defined as the entire genome. Though comprehensive and definitive analysis of the genomic system in isolation is currently impossible (as the genome only exists in an active form in the context of living cells), a default neural state at this level would predict that the highest genes within the neural genetic program (i.e., a potential “master neural genes”) would have only repressor elements within its regulatory regions. In this model, signals extrinsic to the genome would not be necessary to activate the neural genetic program; rather, they could serve only to inhibit it. Transcriptional repressors have been identified that are responsible for inhibiting the neural program, e.g., neuron-restrictive silencing factor/RE1-silencing transcription factor (NSRF/REST; Chen et al., 1998). In vivo inhibition of NSRF/REST leads to derepression of neural genes in both neural and nonneural tissues (Chen et al., 1998). Thus, it is possible that NSRF/REST, as well as other transcriptional repressors, is required to prevent the default neural genetic program from being activated in all cells.

**Materials and methods**

**ES cell culture**

ES cells were maintained on mitotically inactivated mouse embryonic fibroblast feeder layers in DME media containing 15% FCS and LIF (1,000 U/ml). The main ES cell line used was R1, as well as E14K, ad−/−/Y (Joza et al., 2001), cas9+/−/cas9+/− (Hakem et al., 1998), apaf1−/− (Yoshida et al., 1998), and smad4−/− (Sirard et al., 1998). ES cells (provided by J. Rossant, A. Nagy, T. Mak, and J. Penninger, University of Toronto, Toronto, Canada).

**Animals and NSC culture**

NSCs were isolated from the forebrain ventricular subependyma of adult CD1 mice [Charles River Laboratories] and cultured/maintained using the NS assay as previously described (Seaberg and van der Kooy, 2002).

**ES cell minimal condition assays**

ES cells were dissociated into a single-cell suspension and plated at ≤10 cells/μl on laminin/polyornithine–coated culture plates (Nuncclone) in chemically defined, serum-free media (Tropepe et al., 1999, 2001), which consisted of DME/F-12 [1:1; Invitrogen] with 0.6% d-glucose, 5 mM Heps, 3 mM NaHCO 3, 2 mM glutamine, 25 μg/ml insulin, 100 μg/ml transferrin, 20 nM progesterone, 60 μM putrescine, and 30 nM sodium selenite. PBS used for cell incubation in the short-term neural conversion experiments included 20 mM glucose. Uncoated dishes were also tested, and though cells exhibited greatly decreased adhesion, no differences in the neural conversion assays were observed. Cells were fixed for immunocytochemical analysis or cells were collected for RT-PCR analysis.
regulated and the cGMP pathway negatively regulated primitive NSC survival and primitive NS formation. The Materials and methods section contains additional details regarding the primitive NS assay and the counting and normalization of data. Online supplemental material is available at http://www.jcb.org/cgi/content/full/jcb.200508085/DC1.

The authors wish to thank members of the van der Kooy laboratory for thoughtful discussion and critical reading of the manuscript; R. McKay and R. Goldman for generously providing nestin antibodies; and J. Penninger, T. Mak, A. Nagy, and J. Rossant for generously providing ES cell lines.

This work was supported by the Canadian Stem Cell Network and the Canadian Institutes of Health Research (grant 79750).

Submitted: 11 August 2005
Accepted: 28 November 2005

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