Rest-Mediated Regulation of Extracellular Matrix Is Crucial for Neural Development

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Abstract

Neural development from blastocysts is strictly controlled by intricate transcriptional programmes that initiate the down-regulation of pluripotent genes, Oct4, Nanog and Rex1 in blastocysts followed by up-regulation of lineage-specific genes as neural development proceeds. Here, we demonstrate that the expression pattern of the transcription factor Rest mirrors those of pluripotent genes during neural development from embryonic stem (ES) cells and an early abrogation of Rest in ES cells using a combination of gene targeting and RNAi approaches causes defects in this process. Specifically, Rest ablation does not alter ES cell pluripotency, but impedes the production of Nestin+ neural stem cells, neural progenitor cells and neurons, and results in defective adhesion, decrease in cell proliferation, increase in cell death and neuronal phenotypic defects typified by a reduction in migration and neurite elaboration. We also show that these Rest-null phenotypes are due to the dysregulation of its direct or indirect target genes, Lama1, Lamb1, Lame1 and Lama2 and that these aberrant phenotypes can be rescued by laminins.

Introduction

During mouse embryo development, the blastocyst differentiates into pluripotent primitive ectoderm and gives rise to a structure known as the epiblast [1]. The epiblast responds to extrinsic signals and generates three primary germ layers (ectoderm, mesoderm and endoderm) [2]. During neurulation, the ectoderm gives rise to the neuroectoderm in the form of a neural plate, which subsequently folds to generate the neural tube, composed of a single layer of neuroepithelial cells or neural stem cells (NSCs), where a series of ring-like constrictions mark the boundaries between the primordia of the major brain regions [3–4]. This process of neural development is orchestrated and accompanied by wholesale changes in transcriptional programmes and patterns of gene expression. However, due to the difficulties in accessing and manipulating early embryos, the transcriptional network that regulates neural development is poorly understood, especially in mammals. Embryonic stem (ES) cells derived from blastocysts retain the ability to recapitulate neural development in vitro, and offer an invaluable model to study early events in embryogenesis. The RE1 Silencing Transcription Factor / Neuron Restrictive Silencer Factor (Rest/Nrsf) is a zinc finger transcription repressor that has been postulated to act as a master regulator of neuronal gene expression in both the developing and mature nervous systems [5–6]. We and others have shown that Rest is highly expressed in blastocysts and ES cells, but that expression decreases as neural development proceeds [7–8]. In fact, down-regulation of Rest has been proposed to be obligatory for differentiation of neural progenitors [8] and more recently, it has been proposed that Rest haploinsufficiency results in loss of pluripotency markers and a reciprocal gain in differentiation markers [9]. Taken together, these observations suggest that Rest may play a crucial role at several stages of neural development. Here, we determine the function of Rest during neural development from ES cells through NSCs and neural progenitor cells (NPCs) to mature neurons using an in vitro ES cell-derived neural differentiation model.

Rest exerts its function by binding to both canonical and non-canonical RE1-sites identified at over 2000 loci in the mammalian genome [10–11] and is implicated in the regulation of both coding and non-coding genes [10,12], many of which represent neuron-specific transcriptional units. The observation that many of these target genes are expressed by differentiated neurons, including ion channels, neurotransmitter receptors, neurotrophins, synaptic vesicle associated proteins, cell adhesion molecules, growth-associated and cytoskeletal proteins, gave rise to the initial perception that Rest acted as a silencer of neuron-specific genes in NPCs and non-neural cells to prevent precocious expression of neuronal characteristics. However, recent studies emerge that Rest has more versatile roles and can regulate its target genes either by activation, repression or silencing, depending upon the developmental stage and cell type [7,13]. Rest recruits multiple cofactors, histone modifying and chromatin remodelling activities, all of which underline the complexity of Rest activity [14–16]. The diverse roles of Rest have been shown in both neural and non-neural pathologies including Huntington’s disease, cardiac hypertrophy, medulloblastoma, malignant rhabdoid tumor, small cell lung cancer, ovarian cancer, and ischemia (see review for references [13]).
Despite the wealth of knowledge in identifying target genes [10–12] and in delineating the mechanistic actions of Rest [14–16], the biological function of Rest during neural development remains unclear. Rest−/− mice die around embryonic day (E)11.5, with embryo degeneration, neural tube malformations and widespread apoptosis evident from E9.5 [6]. Constitutive expression of Rest in chick spinal cord does not cause defects in neurogenesis but does result in axon pathfinding errors [17]. However, in Xenopus, disruption of Rest function disturbs ectoderm patterning and expands the neural plate [18], suggesting that Rest is indeed required for normal neural plate formation and neurogenesis. Collectively, these studies paint a somewhat ambiguous picture of the role of Rest in the development of NSCs and neurons. We have sought to address this issue by using a combination of gene targeting and RNAi to create ES lines expressing a range of Rest concentrations, which we have used to investigate the effect of Rest deficiency during ES cell-derived neural development. Importantly, in contrast to a recent study [9], we find that deletion of a single Rest allele does not result in any change in neural differentiation. Instead, we find that Rest levels have to be decreased by more than 92% to precipitate any phenotype. Rest ablation impairs the extracellular matrix (ECM) components and impedes the production of Nestin+ NSCs, NPCs and neurons.

Furthermore, neurons derived from REST-null ES cells are devoid of elaborate processes, have defects in migration and undergo increased cell death. Importantly, all of these phenotypic effects of Rest ablation were rescued by treatment with laminins, a key component of the ECM that has been implicated in neuronal migration and more latterly in development of the neural plate. We propose a novel mechanism by which Rest regulates development of both NSCs and mature neurons by controlling expression of key components of the ECM.

Results

Rest expression during NSC and Neuron Development

We investigated the role of Rest in neural development using an in-vitro ES cell-derived neural differentiation model, which recapitulates events during neural development in vivo. ES cells firstly differentiate into neuroepithelial cells (early NSCs), which peak around 4–6 days of differentiation and express Sox1 and Nestin (about 80% of population), and then differentiate further into more restricted NSCs that peak around 10 days of differentiation and express either Ngn1 or Mash1 (about 80% of population) (Fig. 1C–D and Fig. S2B–C). In this paper, we refer to early NSCs as NSCs and late more restricted NSCs as NPCs. To
establish the time course during which NSCs, NPCs and neurons are formed from HM1 and 46C ES cells, we examined gene expression patterns for NSC markers (Nestin and Pax6), NPC markers (Mash1 and Ngn1) and neuronal markers (Tubb3, Syn1 and L1cam) during ES cell-derived neural differentiation (Fig. 1A). We found that Rest expression mirrored that of the pluripotent ES cell marker Oct4, and was expressed at highest level in ES cells with its expression level declining as differentiation proceeded; reaching its lowest level 4 days after differentiation just before NSC production reached its peak (Fig. 1B and Fig. S2A). Rest levels were maintained at low levels throughout neuron formation. Conversely, the expression patterns of Nestin and Pax6 reciprocated those of Rest and Oct4 indicating that NSC production started 1- or 2-days after differentiation, reached a peak around 6 days, and thereafter declined (Fig. 1C and Fig. S2B). Mash1 and Ngn1 expression indicated that NPCs started to be produced after 6-days of differentiation and peaked between 6 and 10 days (Fig. 1D and Fig. S2C). The early neuronal marker Tubb3 was observed around the same time but did not peak until 12 days of differentiation whereas the peak of mature neuron marker expression (Syn1 and L1cam) occurred at 14–16 days of differentiation (Fig. 1E and Fig. S2D). This time course closely recapitulates the sequential generation of NSCs, NPCs and neurons observed in vivo (Fig. 1A).

Rest ablation inhibits development of NSC and NPC

Next we examined the effects of Rest ablation on the development of NSCs and NPCs. The control REST-100, REST/KD-50 and REST-null ES cells, which express 100%, 50% and 0% wild-type Rest levels respectively (Fig. S1B and S1D), were differentiated into NSCs and then into NPCs identified using an array of NSC and NPC markers (Fig. 1A). We assessed the effects of Rest ablation on gene and protein expression of these markers using quantitative Real-time PCR, FACS and immunocytochemical analysis. In REST-100 ES cells, the expression profiles of Pax6, Msi1 and Nestin were similar to one another, with expression peaking around 4 days and thereafter gradually declining (Fig. 2A–2C). REST/KD-50 ES cells showed no significant difference in the expression level and pattern of these genes as compared to the control REST-100 cells (Fig. 2A–2C). Similarly, no change in the number of Sox1+/Nestin+ NSCs was seen (Table 1 and Fig. S3B–3C). However, in REST-null ES cells, the peak expression of Pax6 and Msi1 was significantly (P<0.01) reduced to 40% of control levels whilst expression of Nestin was reduced to 60% of control levels (Fig. 2A–2C). This mutant generated significantly (P<0.01) fewer Sox1+/Nestin+ NSCs (52%) as compared to the REST-100 (77%) and REST/KD-50 ES cells (76%), but generated double the number of Sox1+/Nestin− NPCs compared with the control cells (P<0.01) (Table 1 and Fig. S3D). Moreover, REST-null ES cells produced lower levels of Mash1 and Ngn1 expression (P<0.01) and this was reflected in a parallel reduction in the number of Mash1+ NPCs (~50%) as compared to the control (87%) (P<0.05) (Fig. 2D–2F; Table 1; Fig. S4B and 4D). Our results suggest that a 50% depletion of Rest shows no discernible effect on the production of NSCs and NPCs from ES cells. In fact, the production of early Sox1+/Nestin− NSCs from REST-null ES cells remained unaffected but formation of late Sox1+/Nestin+ NSCs and subsequent production of NPCs was inhibited.

Rest ablation impedes neuronal differentiation

Since Rest is known to be a repressor of neuronal gene expression and has been implicated in neurogenesis, we proceeded to examine the role of Rest in the generation of neurons from NSCs by assessing the expression of early and late neuronal markers (Fig. 1A). Similar to our studies on the generation of...
NSCs and NPCs, we noticed no significant difference in either the levels of gene expression, or in the number of cells expressing these neuronal markers in cells derived from REST/KD-50 ES cells compared with the control ES cells (Fig. 2F–2H; Table 1; Figs. S4B–S4C and S3B–S3C). However, REST-null ES cells showed a significant impairment in neuronal differentiation, as evidenced by a reduction in Tubb3 expressing early neurons (P<0.01) (Fig. 2F), and a reduced population of Tau+ neurons (25%) compared to the control cells (42%) (P<0.05) (Table 1; Fig. S4B and S4D). Furthermore, we did not detect Tuj1+ (Tubb3) immunoreactive cells in differentiated REST-null cells at any earlier stage (data not shown), suggesting that Rest abrogation did not result in premature neuronal differentiation. As differentiation proceeded, the number of Map2+ mature neurons was significantly (P<0.01) reduced in REST-null cells (64%) compared with control cells (83%) (Table 1; Fig. S4B and S4D). Expression of Sox1 and L1cam (Fig. 2G–2H) both showed parallel changes. Intriguingly, even though Tubb3, Sox2a and L1cam are all Rest target genes, their expression were not de-repressed in REST-null ES cells during early ES cell differentiation (Fig. 2F–2H), a view consistent with the absence of any effect of acute Rest abrogation on expression of these genes in ES cells (unpublished observations). In conclusion, reduction of Rest levels by 50% has no effect on neurogenesis in terms of the time course, sequence or production of neurons. In contrast, the absence of Rest impedes neurogenesis.

Constitutive expression of Rest rescues REST-null phenotypes

The REST-null ES cells were generated by a combination of gene targeting and RNAi knockdown. To confirm that their phenotype was not due to an off-target event, we constitutively expressed Rest using pMT-NRSF to create REST-null+REST, which raised REST levels to 8% of wild-type levels. Transfection of empty pMT vector was used to generate REST-null+MTV that served as a control (Fig. S1C–S1D). We then assessed the capacity of REST-null+REST ES cells to differentiate into NSCs, NPCs and neurons. REST-null+REST ES cells generated NSCs expressing Pax6 and Msi1 at a similar level to that of REST-100 ES cells while Nestin was rescued to 80% of REST-100 levels. As predicted, REST-null+MTV cells showed a similar phenotype to the REST-null mutant (Fig. S6A–S6C). In support of these findings, the number of Nestin+ NS cells derived from REST-null+REST ES cells significantly increased to 63% (Table 1 and Fig. S6E–S6F). Moreover, constitutively expressing Rest rescued the capacity of REST-null mutant ES cells to produce NPCs, and neurons (Table 1, Figs. S4E–S4F and S5E–S5F). Our findings suggested that the phenotypic effects of REST-null ES cells are not due to off-target effects. Furthermore, our results indicate that very low levels of Rest are both sufficient and necessary for normal generation and maturation of NSCs.

Rest ablation results in defects in adhesion, cell proliferation and survival

During investigation of the effects of Rest ablation on NSC and neuronal development, we observed that REST-null ES cells exhibited more severe phenotypes when plated on a glass surface than when plated on a plastic surface, whereas REST-100 and REST/KD-50 showed no discernible difference on either surface. On glass, REST-null cells showed defective adhesion and produced very few or no Nestin+ NSCs but did produce normal levels of Sox1+ NSCs as compared to those derived from REST-100 and REST/KD-50 ES cells (Fig. 3A–3C). This phenotype was much less pronounced when REST-null ES cells were plated on a plastic surface where a greater number of Nestin+ NSCs were seen (50%; Table 1). Intriguingly, NeuN+ and/or Map2+ neurons showed marked phenotypic defects on the glass surface, characterised by an absence of migration, and poor elaboration of processes and fasciculation among neuronal colonies as compared to REST-100 and REST/KD-50 ES cells (Fig. 3F–3H and 3K–3M). All phenotypes on the glass surface were rescued, at least in part, by raising Rest levels to 8% in REST-null+REST ES cells (Fig. 3D–3E; 3I–3J and 3N–3O). Interestingly, the phenotypic

| Table 1. The summary of FACS analysis of neural populations derived from control and Rest mutant ES cells with or without laminin treatment. |
|-----------------|---------------|---------------|---------------|---------------|
| REST mutant    | Cell type     | NSC            | NPC            | Neuron         |
|                 |               | Sox1+/Nestin   | Sox1+/Nestin   | Mash1+/Tau1+  |
|                 |               | 77.4±1.3       | 47.2±2.7       | 41.7±3.6      |
|                 |               | 77.4±1.3       | 44.5±2.6       | 36.4±1.8      |
|                 |               | 62.9±1.4**     | 29.8±2.1*      | 22.7±2.4      |
|                 |               | 62.9±1.4**     | 39.1±1.6*      | 31.5±2.1*     |
|                 |               | 78.8±3.1       | 38.3±2.1       | 31.8±2.4      |
|                 |               | 77.1±2.1       | 40.8±2.3       | 28.5±2.6      |
|                 |               | 74.9±2.5       | 37.8±1.9       | 31.9±2.2      |
|                 |               | 75.2±2.3       | 37.1±2.2       | 29.5±2.1      |
|                 |               | 78.4±1.8       | 35.9±2.4       | 31.4±1.8      |

Data are represented as mean±SEM.

*P<0.05 and **P<0.01, significantly different from REST-100 and REST/KD-50.

P 0.05 and ¥P 0.01, significantly different from REST-null and REST-null+MTV.

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null NSCs derived from the control, REST/KD-50 and REST-null ES cells. However, it was difficult to evaluate the proliferation of Sox1
+ cells now expressing Nestin (E). (F–O) Neurons derived from the control and 4 Rest mutant ES cells were identified using NeuN (red) and Map2 (green) after 14 days of differentiation. Neurons generated from REST-100 and REST-null+REST ES cells showed much higher levels of Nestin immunostaining in Sox1
+ cells (A–B). Constitutive expression of Rest (REST-null+REST) rescues, at least partially, this phenotype with many Sox1
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+ cells now expressing Nestin (E). (F–O) Neurons derived from the control and 4 Rest mutant ES cells were identified using NeuN (red) and Map2 (green) after 14 days of differentiation. Neurons generated from REST-100 and REST/KD-50 ES cells showed elaborate neurite outgrowth, fasciculation between aggregates/colonies (F, G and in higher power K, L) and migration of NeuN+ cells (see arrows in L). Conversely, neurons derived from REST-null and REST-null+MTV exhibited fragmented neuronal colonies that lacked elaborate processes and migration (H, I and in higher power M, N). Constitutively expressing Rest (REST-null+REST) attenuated the phenotypic effects of Rest ablation: neurite outgrowth, neurite fasciculation (J, O) and neuronal migration was observed (arrows in O). Scale bars: 20 μm (A to E and K to O) and 100 μm (F to J).

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Defects on the glass surface were also seen in NSCs and neurons derived from Rest–/– ES cells [6] (Fig. S7A–S7K), although in the latter case the Rest mutant retains alternatively spliced isoforms that may not be functional (Fig. S7A–S7B).

Either on plastic or glass surfaces, we found that REST-null ES cells generated fewer cells during neural differentiation than those from REST-100 and REST/KD-50 ES cells. Accordingly, we examined proliferation and cell death during the course of neurogenesis. Using BrdU incorporation, no differences were observed in the proliferation of either Sox1
+/Nestin– early NSCs derived from control and all Rest mutants or in Sox1
+/Nestin– late NSCs derived from the control, REST/KD-50 and REST-null+REST ES cells. However, it was difficult to evaluate the Sox1
+/Nestin– NSCs from REST-null and REST-null+MTV ES mutants, because of their low number and the ease with which they were lost from coverslips during the staining process. We then used TUNEL staining to assess the degree of apoptosis at 4 and 14-days of differentiation, i.e. at the peaks of NSC and neuron generation respectively. The ES monolayer culture system used in this study does not employ mitogens to induce neural differentiation [19], and accordingly, significant cell death occurs during NSC formation, especially during days 3 through 6 of differentiation, but thereafter this becomes much less marked as neuronal differentiation proceeds. At the NSC stage, cell death was equally prevalent in all samples derived from either control cells or Rest mutants (Fig. 4A–4E). However, at the neuronal stage, there was markedly more cell death in REST-null cells than those in the control and REST/KD-50 cells (Fig. 4F–4H), which is correlated with the ailing look of neurons derived from REST-null ES cells (Fig. 3H and 3I). Our results indicate that Rest ablation did not impair the proliferation of Sox1
+/Nestin– NSCs; however it is difficult to assess the proliferation of Sox1
+/Nestin– NSCs, due to the paucity of Nestin+ NSCs derived from REST-null ES cells. Moreover, Rest levels had to be reduced by more than 92% before an increase in cell death at the neuronal stage was observed.

Laminins rescue the phenotype of the REST-null mutant

REST-null defects in cell adhesion may be the cause of its phenotypic effects on Nestin+ NSC production and neuronal differentiation, because cell adhesion defects caused aberrant NSC and neuron development [20–21]. Thus, we further examined the causes of defective adhesion seen in the REST-null mutant. We and others have previously reported that several Rest target genes encode cell adhesion molecules or components of the extracellular matrix (ECM), particularly laminin subunits [7,11]. Accordingly, we considered the notion that dysregulation of the ECM by Rest ablation might be responsible for this aspect of the phenotype. To test this hypothesis, we examined whether we could rescue the adhesion defect and any other phenotypic effects caused by Rest ablation by pre-treatment with ECM components. We plated ES cells from all groups on to either plastic or glass pre-treated with EHS laminins (which contain predominantly Laminin 1 (α1β1γ1)) [22] and subsequently subjected them to neural differentiation. On the glass surface pre-treated with laminins, but not those pre-treated with gelatin, the Rest mutant ES cells behaved like control ES cells, both of which adhered firmly and proliferated well. After 4-days of differentiation (NSC stage), cells from all groups plated onto glass surfaces pre-treated with laminins showed greater survival and significantly less apoptosis than those on untreated surfaces (Fig. 4K–4O). There was no significant difference in cell growth among the treated groups (the control and mutants). The surviving cells were highly proliferative as adjudged by BrdU incorporation (data not shown). Laminin pre-treatment had an even more profound effect after 14-days of differentiation.
neuronal stage). No, or very few, apoptotic cells were detected in either the laminin-treated or control cells without laminin-treatment (Figs. 4F and 4P–4T), whereas significant cell death was found in the Rest mutants in the absence of laminin (Fig. 4H–4J). These data suggested that laminins prevented cell death in the REST-null cells. Intriguingly, the laminin rescue also extended to the specification of NSCs. We reported above that the REST-null ES cells were able to generate early Sox1+/Nestin− NSCs but were unable to produce normal numbers of late Sox1+/Nestin+ NSCs when they were differentiated on a gelatinised glass surface in the absence of laminins (Fig. 3C–3D). This deficiency was rescued by laminin-treatment. Under these conditions, REST-null mutants, like control ES cells, generated equivalent numbers of both early NSCs (Sox1+/Nestin− ) and late NSCs (Sox1+/Nestin+ ) (Fig. 5A–5E; Table 1). Furthermore, laminins rescued the production of NPCs and neurons from the REST-null mutant (Table 1). Neurons derived from REST-null ES cells showed widespread aggregation with extensive fasciculation (Fig. 5H–5I and 5M–5N) compared with those produced on glass in the absence of laminins (Fig. 3H–3I and 3M–3N).

To ensure that the rescue described above was not simply due to the enhanced adhesive properties of laminin, we investigated the ability of fibronectin (a different ECM component with similar adhesive properties to laminin) or poly-D-lysine (a commonly used non-biological substrate for neural cells) to mimic this response. Fibronectin was far less effective in rescuing the generation of Sox1+/Nestin+ cells (late NSCs) from REST-null ES cells than laminin. Some Sox1+/Nestin+ NSCs did develop when grown on fibronectin, however, the number observed was dramatically fewer when compared to that observed when REST-null cells were differentiated on laminins or when REST-100 cells were differentiated (Compare Fig. S8A with Fig. 5A and 5C). Some neurons were observed when REST-null cells were differentiated on fibronectin but again in far fewer numbers than seen after rescue with laminin or after differentiation of control cells (Fig. S8B). Phenotypically, neurons that developed on fibronectin had elongated process, suggesting that fibronectin was able to rescue the neuronal morphology of REST-null derived neurons, at least in part (Fig. S8B). The phenotype observed when REST-null ES cells were differentiated on poly-D-lysine was similar to that observed with gelatine; few Sox1+/Nestin+ NSCs developed and all neurons observed looked ailing, being rounded and devoid of processes (compare Fig. S8A with Fig. 3C and Fig. S8B with Fig. 3H and 3M). Collectively, our results indicate that laminins rescued the adhesion defects seen during differentiation of REST-null ES cells and concomitantly rescued their ability to differentiate into Nestin+ NSCs, NPCs and neurons. This rescue cannot solely be attributed to the adhesive properties of laminin.
Furthermore, a remarkable increase in Nestin (red) and Sox1 (green) (A–E). The results show that laminins restored the ability to generate Nestin+ cells in Rest mutants (C–E, compared with Fig. 3C–3E). Sister cultures were allowed to differentiate for 14 days to analyse neuronal differentiation using NeuN (red) and Map2 (green) (F–J and in higher power K–O). Laminins rescued the neuronal phenotypes derived from REST-null ES cells, which included elaboration processes, neurite outgrowth and migration (see arrows in M and N) similar to that seen in neurons derived from control ES cells. Nuclei were stained with DAPI (blue) in all. Scale bars: 20 μm (A to E and K to O) and 100 μm (F to J).

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Figure 5. Laminins rescue the phenotypic effects of Rest ablation on the neural differentiation of ES cells. Control (REST-100) and 4 Rest mutants ES cells were grown under neural-differentiating conditions on laminin coated glass coverslips. After 4 days of differentiation some cultures were fixed and labelled with the NSC markers Nestin (red) and Sox1 (green) (A–E). The results show that laminins restored the ability to generate Nestin+ cells in Rest mutants (C–E, compared with Fig. 3C–3E). Sister cultures were allowed to differentiate for 14 days to analyse neuronal differentiation using NeuN (red) and Map2 (green) (F–J and in higher power K–O). Laminins rescued the neuronal phenotypes derived from REST-null ES cells, which included elaboration processes, neurite outgrowth and migration (see arrows in M and N) similar to that seen in neurons derived from control ES cells. Nuclei were stained with DAPI (blue) in all. Scale bars: 20 μm (A to E and K to O) and 100 μm (F to J).

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but must involve laminin-directed signalling since fibronectin has similar adhesive properties yet was dramatically less effective at rescuing both the NSC and neuronal differentiation.

Impairment in laminins is caused by Rest ablation

The laminin rescue of the REST-null phenotype indicated that laminins are downstream effectors of Rest. To test this idea, we determined whether the expression of laminins during neurogenesis was impaired by Rest ablation. Laminins have over 15 isoforms, each consisting of a combination of α, β, and γ subunits. In this experiment, we only focused on the effect of Rest deficiency on the expression of the genes encoding α1 (Lama1), β1 (Lamb1), γ1 (Lama1) and α2 (Lama2) subunits since the EHS laminins used in the rescue experiments are composed mainly of laminin 1 (α1β1γ1). Furthermore α2 is a known Rest target gene [11] and a major component of the basement membrane in the embryonic CNS that is known to be involved in NSC development [23]. Each of these 4 laminin subunits exhibited distinct expression patterns during neural differentiation of ES cells (Fig. 6). In REST-100 and REST/KD-50 cells, the expression pattern of Lama1 closely followed the time course of NSC and neuron development. Expression of Lamb1 and Lama1 both showed a very similar pattern with high expression levels in ES cells followed by an initial decline during NSC differentiation and a subsequent gradual increase as neuronal differentiation proceeded. These expression patterns of Lama1, b1 and c1 are similar to those reported in an earlier in vitro study [24]. Intriguingly, the expression pattern of Lama2 correlated closely with the time course of NSC formation, peaking at 4-days of differentiation, at a similar developmental stage as expression of α2 in vivo [23]. In contrast, REST-null cells exhibited significantly (P<0.01) decreased expression levels of Lama1 throughout neurogenesis (Fig. 6A). Remarkably, raising the Rest level to 8% (REST-null+REST) restored Lama1 levels to 40–50% of those of the control, although the levels remained significantly (P<0.01) lower than those seen in control cells. A similar change was also seen in the expression patterns of Lamb1, c1 and a2. Rest ablation reduced the levels of Lamb1, c1 and a2 to below 40% of the level seen in control cells throughout neurogenesis, while raising the Rest level to 8% restored, at least partially, the expression levels of Lamb1 and a2 (Fig. 6B–6D). Taken together, our results suggest that Rest ablation impairs expression of laminins 1 and 2 (α2β1γ1) during neurogenesis, which leads to defects in cell adhesion, expansion and fasciculation.

Discussion

Development of the vertebrate nervous system is orchestrated by transcriptional programs executed by both transcriptional activators and repressors. Studies suggested that the transcription factor Rest acts as a master regulator to suppress premature differentiation of neuronal progenitors and secure orderly neuronal maturation. Despite a wealth of information on the mechanism of Rest action and on identification of over 2000 target genes, we know very little about the biological function of Rest in the developing and mature nervous system. In our efforts to delineate the function of Rest during neural development, we generated several Rest deficient ES cell mutants to elucidate the role of Rest in the transition from pluripotent ES cells to multipotent NSCs and subsequently to mature neurons. Here, we show that deletion of a single Rest allele has no discernible effect on either NSC formation or neurogenesis but severe depletion of Rest to levels less than 8% impedes NSC and neuron development, and further that these impairments are mediated by attenuation of laminin levels.

Rest is dispensable for ES cell pluripotency

Recently, a study showed that knocking down Rest by 50% altered ES cell pluripotency and promoted ES cell differentiation into endoderm, mesoderm and ectoderm in ES medium [9].
However, our findings show that depletion of Rest levels by 50% to 100% does not change ES cell pluripotency as assessed by expression of pluripotent markers and by lineage competence. In this study, we have generated Rest mutant ES cells, which expressed Rest at 50%, 8% and 0% of the wild-type level. We found that these mutant ES cells did not significantly decrease the expression of Oct4 and Nanog, a finding that is congruent with observations of Rest+/− and Rest−/− ES cells derived from Rest knockout mice in which gastrulation proceeds normally [6]. Moreover, we also found that two independently derived ES lines with only one allele of Rest, REST/KD-50 and Rest−/+−, both behave like their wild-type counterparts, and are fully capable of normal in vitro neural differentiation. These findings are consistent with the findings that, in vivo, Rest−/+− ES cells are pluripotent and are able to generate germ-line mice in which gastrulation proceeds normally [6]. Moreover, we also found that two independently derived ES lines with only one allele of Rest, REST/KD-50 and Rest−/−, both behave like their wild-type counterparts, and are fully capable of normal in vitro neural differentiation. These findings are consistent with the findings that, in vivo, Rest−/+− ES cells are pluripotent and are able to generate germ-line mice in which gastrulation proceeds normally [6]. Moreover, we also found that two independently derived ES lines with only one allele of Rest, REST/KD-50 and Rest−/−, both behave like their wild-type counterparts, and are fully capable of normal in vitro neural differentiation. These findings are consistent with the findings that, in vivo, Rest−/+− ES cells are pluripotent and are able to generate germ-line mice in which gastrulation proceeds normally [6].

Rest ablation impedes NSC and neuron development

There are over 2000 RE1 sites in the human and murine genomes [12] and many RE1-bearing genes encode ion channels, neurotransmitters, growth factors and hormones, and factors involved in axonal guidance and vesicle trafficking, and molecules involved in maintenance of the cytoskeleton [5,25]. The observation that many of these Rest target genes are expressed by postmitotic neurons, helped to foster the initial belief that the role of Rest was to prevent premature expression of neuron-specific genes in NPCs [26–27,6]. Further, it has been proposed that down-regulation of Rest in NPCs is required for their subsequent differentiation [8]. However, we found that Rest ablation in ES cells and NSCs results neither in increased NSC production nor in precocious neuronal differentiation. Rather, Rest ablation significantly decreases the production of Nestin+, Pax6- or Msi1-expressing NSCs without affecting that of Sox1+ NSCs, suggesting that Nestin+ NSCs are derived from Sox1+ NSCs. This view is confirmed by previous studies showing that the expression of Nestin is regulated by co-operative action between Sox1-3 and class III POU transcription factors [28]. Our results indicate that the function of Rest is not restricted to regulating neuronal differentiation as previously conceived, but that Rest also takes part in upstream events regulating the generation of Nestin+ NSCs from Sox1+ NSCs. This regulation of early NSC development is echoed in a previous study that showed that disruption of Rest function in Xenopus disturbs ectoderm patterning and expands the neural plate [18].

Additionally, Rest ablation also reduces the production of Mash1+ (and Mash1+) or Ngn1+ NPCs. This may be secondary to the diminished NSC population since there are no detectable differences in proliferation or cell death during derivation of NSCs from either the control or REST-null ES cells. Our data show that the role of Rest in NSCs and NPCs is not simply to repress expression of neuronal target genes and to prevent precocious neuronal differentiation, but rather that Rest plays manifold roles at the level of both NSC and NPC generation.

We also find that the absence of Rest in ES cells and NSCs does not cause precocious neuronal differentiation. In fact, even direct Rest target genes, such as the neuronal markers Tubb3, L1cam and Scn2A, are not precociously expressed. This finding resonates strongly with the initial experiments on Rest−/− mice, which displayed embryonic lethality around E11, but importantly, showed no widespread precocious expression of Rest target genes.

Figure 6. The effects of Rest ablation on the gene expression of laminin subunits during neural differentiation. The expression patterns of Lama1 (A), Lamb1 (B), Lamc1 (C) and Lama2 (D) were analysed throughout ES cell-derived neural differentiation by real time-PCR. Data are represented as mean ± SEM. *P < 0.05 and **P < 0.01, significantly different from REST-100 and REST/KD-50. #P < 0.05 and ¥P < 0.01, significantly different from REST-null and REST-null+MTV.

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Rest in Neural Development

However, our findings show that depletion of Rest levels by 50% to 100% does not change ES cell pluripotency as assessed by expression of pluripotent markers and by lineage competence. In this study, we have generated Rest mutant ES cells, which expressed Rest at 50%, 8% and 0% of the wild-type level. We found that these mutant ES cells did not significantly decrease the expression of Oct4 and Nanog, a finding that is congruent with observations of Rest+/− and Rest−/− ES cells derived from Rest knockout mice in which gastrulation proceeds normally [6]. Moreover, we also found that two independently derived ES lines with only one allele of Rest, REST/KD-50 and Rest−/+−, both behave like their wild-type counterparts, and are fully capable of normal in vitro neural differentiation. These findings are consistent with the findings that, in vivo, Rest−/+− ES cells are pluripotent and are able to generate germ-line mice in which gastrulation proceeds normally [6].

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Only Tubb3 was seen to be de-repressed in non-neuronal tissues but not in the developing CNS. Furthermore, constitutive expression of Rest in chick spinal cord causes axon pathfinding errors, but only two Rest target genes encoding Tubb3 and L1cam were repressed [17]. Taken together, our findings suggest that the role of Rest in neuronal differentiation is not to simply regulate its neuronal target genes by a simple “on-off” switch. The regulation of Rest in these target genes may be compounded by other transcriptional repressors or co-repressors, such as having been reported for SMCX, a transcriptional repressor that co-regulates Rest neuronal target genes in X-linked mental retardation [29] and NFkappaB that synergistically interacts with Rest in the suppression of TAC1 in non-neuronal cells [30].

Strikingly, there is also no evidence that the absence of Rest promotes or increases neuron number [31–32]. On the contrary, we find that Rest knockout decreases neuron production and causes severe defects in neuronal interaction (judged by malformation of neuronal processes and fascicles and neuronal migration). The decline in neuron production, in part, is attributed to cell death, because we found that Rest ablation increases neuronal death. These phenomena however can be rescued by increasing levels of Rest expression, indicating that Rest is a causal effect. Taken together, the role of Rest during neuronal development is not restricted simply to repress neuron-specific genes in NPCs prior to terminal neuronal differentiation.

Laminin deficiency causes impairment in NSC and neuron development in REST-null ES cells

In this study, the most striking phenotype of Rest mutant ES cells is their defective adhesion, especially on a glass surface. The other phenotypes include impairment in the production of Nestin+ NSCs (albeit the mutants can generate a comparable amount of Sox1+ NSCs), and defects in neuronal migration and process formation. Additionally, Rest ablation results in greater cell death and a consequent reduction in the number of neurons. Interestingly, we also observed these phenotypes in ES cells derived from Rest−/− mice [6]. Our finding suggests these are bona fide phenotypes of Rest-null mutants and not due to the artefact of a single ES cell line used in this study. In previous studies, we and others identified several Rest target genes that are involved in cell-matrix adhesion and cell-cell interaction, such as Lama1, Lama2, Arc, Cyp3g, Unc3d, Adam23, Catnd2, Cldn4, L1-cam and Mmp24 [7,11,12]. This latter observation prompted us to consider the possibility that dysregulation of cell-matrix adhesion molecules may be responsible for Rest-knockout phenotypes.

Cell-matrix adhesion molecules mediate direct interactions of the cell with its extracellular environment by binding of cell surface molecules with components of ECM, and are crucial for cell migration, differentiation, organisation and embryogenesis [33–35]. In the developing central nervous system, the ECM is present in the basement membrane of the ventricular zone and is essential for ordered differentiation of neuronal subtypes in the cerebral cortex [36]. In vitro, the ECM has been shown to play an important role in lineage decision and cell type selection [21,37]. The ECM mainly consists of laminins, fibronectin, type IV collagen, nidogen and heparan sulfate proteoglycans [38–39]. Laminins are a family of extracellular glycoproteins expressed throughout developing neural tissues and each laminin is composed of α, β, and γ polypeptide chains [40]. In this study, EHS laminins rescue the phenotypes of Rest mutant, by restoring their adhesion ability and their differentiation into Nestin+ NSCs, NPCs and mature neurons. Not only do laminins promote neuron migration, neurite outgrowth and elaboration, but they also prevent neuronal cell death caused by Rest ablation. These findings resonate with the known functions of laminins in the NS cell niche where they are required for proliferation, neuronal differentiation and survival, and neurite outgrowth [20,23,41]. Additionally, we found that although fibronectin exhibited similar adhesiveness to that of laminins, it nevertheless was markedly less effective than laminins in restoring the REST-null phenotype, indicating that specific ECM signalling rather than adhesion alone is the key player in REST-null phenotypes.

How does Rest impair the expression of laminins, which subsequently generate phenotypic effects in Rest mutants? We know that the genes encoding laminin subunits α1 and α2 are direct Rest target genes [11] and γ1 is an indirect target since it is regulated by miR-124, which is a direct Rest target gene [42]. During neural differentiation of ES cells, the expression pattern of Lama1 showed an inverse relationship with that of Rest, suggesting Rest may act as a repressor in Lama1 expression. However, Rest ablation in ES cells caused impairment in stage-dependent Lama1 expression, indicating that a minimal amount of Rest is required to maintain Lama1 expression. This is the case that can be seen from the rescue of Lama1 expression by raising Rest levels to 8%. This phenomenon also applies to the Lama2 expression. In fact, the expression of laminin subunits correlates with severity of phenotypic effects of Rest deficiency. In conclusion, our studies show that one of the roles of Rest in neural development is to regulate ECM components, which in turn are required for the transition from ES cell to neural lineage. Furthermore, we show that Rest is required at multiple stages during neural development from production of NSCs through to terminal neuronal differentiation. This contrasts with the contemporary idea that limits Rest to regulation of neuronal development by simply repressing neuron-specific target genes prior to loss of Rest expression during the final stages of terminal neuronal differentiation.

Materials and Methods

Generation of Rest mutant ES cells

To generate Rest mutant ES cells, firstly, we have constructed a conditional knockout targeting vector, REST/ck vector (Fig. S1A) to replace one allele of the Rest gene in HM1 ES cells (a gift from Dr. J. McWhir, the Roslin Institute) using homologous recombination. In general, HM1 ES cells were transfected with the REST/ck vector digested with AlII/BglII using electroporation (800 V, Time constant 0.2 msec) and in G418 selection from the second day after transfection. Colonies were screened for targeted clones using RT-PCR with two primer pairs, primers 1/2 and primers 3/4 (Fig. S1A). Of 500 colonies, one clone was identified as a REST/ck-targeted clone, which was used to create the control ES cells (called REST-100) and Rest+/− ES cells (called REST/KD-50). REST-100 ES cells were generated by stably transfecting with the empty pSuper vector, whereas REST/KD-50 ES cells were created by transiently expressing Cre recombinase (a gift from Dr. Jeremy Brown) to delete one allele of Rest. The REST/KD-50 ES cells were then used to create REST-null ES cells by stably transfecting with the empty pSuper vector, whereas REST/ KD-50 ES cells were created by transiently expressing Cre recombinase (a gift from Dr. Jeremy Brown) to delete one allele of Rest. The REST/KD-50 ES cells were then used to create REST-null ES cells by stably expressing a Rest shRNA (5'-GTGTAATCTACAATACCAT-3') in the presence of puromycin (2 μg/ml) using a pSuper- Rest-shRNA vector [43].

The Rest levels expressed from REST-100, REST/KD-50 and REST-null ES cells were 100%, 50% and 1%, respectively as adjudged by quantitative Real-time PCR (Fig. S1B), and confirmed by Western blot analysis (Fig. S1D). In our Western blot analysis, the level of Rest in REST-100 (lane 1) is approximately double that seen in REST/KD-50 (lane 2) when normalised to levels of the house-keeping gene, Gapdh (lower panel) (Fig. S1D). Although REST-null ES cells produced 1% of
wild type Rest mRNA levels, as detected by PCR analysis, Rest protein level was undetectable by Western blot analysis using a Rest antibody (Upstate; 07-579) at a 1:1000 dilution. Therefore, we consider this to be a Rest-null mutant. For a rescue experiment, we constitutively expressed Rest in the REST-null ES cells with the pMT-NR1SF vector (a gift from David Anderson, Caltech [27]) to produce REST-null+REST, which expresses 0% of the control Rest levels. Control cells were produced by transfection of REST-null ES cells with empty pMT vector REST-null+MTV (Fig. S1C–S1D). Although Rest in REST-null+REST ES cells was not detectable by Western blot analysis, it was detected by immuno-
cytochemical analysis (data not shown). We also showed that Rest is effectively silenced throughout the neuronal differentiation process (Fig. S1B–S1C). Furthermore, in contrast to recent observations [9], we also found that Rest deficiency (50% to 100%) did not affect the pluripotency of ES cells as judged by the expression levels of Oct4 and Nanog (Fig. S1E).

Quantitative Real Time-PCR

Primer design and experimental details were carried out as described previously [7] and in Methods S1. Primers used in this study are shown in Table S1. All expression levels were normalised to cyclophilin levels and then as a percentage of the highest level of expression of the REST-100 clone. All data were performed in duplicate and repeated three times.

Statistical Analysis

Statistical significance was determined using a two-tailed Student test. P values of <0.05 and <0.01 were considered statistically significant. All results are presented as mean±standard error of the mean (SEM) from experiments that have been repeated three times.

Neuronal differentiation

Rest mutant and control ES cells were differentiated into NSCs, NPCs and then neurons using a monolayer culture in N2B27 medium [19]. For studying gene expression patterns and for cell type analysis, control and Rest mutant ES cells (600,000 cells/plate) were plated onto 10-cm Petri dishes coated with 0.1% gelatin in ES medium. The next day, the ES cell medium was replaced with N2B27 medium. Medium was changed every 2 days and differentiation continued for 18 days. For studying gene expression patterns, samples were collected at day 0 (ES cell stage, before differentiation) and 2, 4, 6, 8, 10, 12, 14, 16 and 18 days after differentiation. For cell type analysis using FACS, samples were collected separately from 4, 10 and 14 days of differentiation, which were plated out either on gelatin or laminin-treated surface. In immunoocytochemical analysis, ES clones were seeded onto gelatinised, polyornithine/fibronectin (10 μg/ml, Sigma), polyornithine/laminin (10 μg/ml, Sigma) or poly-D-lysine (60 μg/ml, Sigma) coated glass coverslips (VWR, 631-0150) in 24-well plates at a density of 1×10⁴/cm² in ES cell medium. The next day, cells were subjected to the differentiation process in N2B27 medium. All experiments were repeated three times.

Immunocytochemistry

ES cells, NSCs and neurons were fixed in 3% paraformalde-
hyde for 20 min at RT. Antibodies, Nestin (1:500, MAB353, Chemicon) and Sox1 (1:300, AB5934, Chemicon) were used to identify neural stem cells, whereas NeuN (1:500, MAB377, Chemicon) and MAP2 (1:400, ab10588-50, Abcam) were used for staining neurons. All primary antibody staining was carried out at 4°C, overnight. Samples were then stained with appropriate fluorescence-conjugated secondary antibodies at RT for 1 hr and examined under an Axiovision fluorescent microscope (Zeiss).

Flow cytometry

Cells were collected 4, 10 and 14 days after differentiation for FACS analysis of NSCs, NPCs/early neurons and mature neurons, respectively. Cells were trypsinised to dissociate into single cells, fixed in 3% paraformaldehyde for 15 min, permea-
bised with 0.1% saponin (Invitrogen) for 30 min and then stained with Nestin/Sox1 (both 1:250) or Mash1/Chemicon/Tau (Che-
mic) (both 1:200) or Map2 (1:300) in the presence of 0.1% saponin for 1 hr. All procedures were carried out at room temperature. Cells were subsequently stained with their corresponding secondary antibodies: Cy3-anti-mouse/FTTC-anti-chick-
en (both 1:300) for Nestin/Sox1, Cy3-anti-goat/FTTC-anti-rabbit (both 1:300) for Mash1/Tau and FTTC-anti-chicken (1:300) for Map2. The corresponding controls were stained only with secondary antibodies. Cells were acquired with a FACS LSRII with CellQuest software (BD biosciences). Flow cytometry data were analysed using Summit v4.3 (Dako Colorado, Inc). All FACS analysis experiments were repeated three times.

Cell proliferation and apoptosis

5-bromo-2-deoxyuridine (BrdU; 10 μM; Sigma) was added to ES cells grown on polyornithine/laminin (10 μg/ml) coated 13 mm glass coverslips. 16 hrs later, cultures were fixed with 2% paraformaldehyde for 15 min, followed by 95% methanol for 30 min at room temperature. After rinsing with PBS, coverslips were incubated with a biotin conjugated sheep anti-BrdU antibody (1:250; Abcam) overnight at 4°C in PBS containing 0.1% Triton X-100 and 10% normal goat serum (Sigma). BrdU labelled cells were visualised using streptavidin conjugated to Alexa Fluor 488 (1:500; Invitrogen). After three washes with PBS, coverslips were mounted on slides using Fluoromount-G (Invitrogen) and analysed using an Axiovision fluorescent microscope (Zeiss). Apoptotic cell death was detected by terminal deoxynucleotidyl transferase-mediated dUTP-digoxigenin nick end labelling (TUNEL) according to manufacturer’s instructions (Promega).

Supporting Information

Figure S1 The generation of Rest mutants. (A) The REST conditional knockout targeting vector (REST/ck) (bottom) was designed according to Rest gene (top). The REST/ck vector was constructed by inserting a FRT-ik-neor-FRT-LoxP cassette at the Aat II site in intron 2 and a LoxP site at the Asc I site in the exon 1 of Rest gene. This design is to abrogate Rest production by snapping off exons 1 and 2, which encode the transcriptional start sites, the N terminal repression domain and 4 zinc fingers of Rest protein, in the presence of Cre recombinase. Primers shown were used to identify targeted clones. (B–C) Rest expression levels during ES cell-derived neural development from the control (REST-100), REST/KD-50, REST-null, REST-null+MTV and REST-null+REST. (Data are represented as mean±SEM.) (D) Western blot analysis in the control and Rest mutants. Lane 1: REST-100; 2: REST/KD-50; 3: REST-null; 4: REST-null+MTV and 5: REST-null+REST. Gapdh as an internal control. (E) The pluriptoty of embryonic stem (ES) cells was not altered in Rest mutants judged by the expression of ES cell markers, Oct4 and Nanog. (Data are represented as mean±SEM.)

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Figure S2 Gene expression patterns of stage-specific markers during 46C ES-derived neural differentiation. ES cell differentiation into neurons recapitulates neurogenesis in vivo down-
regulation of ES cell pluripotent marker Oct4 and Rest (A) to the
sequential development, firstly neural stem cells (Nestin and Pax6) (B), then neural progenitor cells ( Mash1 and Ngn1) (C) and to neurons (Syn1, Llicam and Tubb5) (D). This experiment corroborated the findings in H1M ES cells.

Found at: doi:10.1371/journal.pone.0003656.s002 (0.07 MB TIF)

**Figure S3** FACS analysis of Sox1+ and/or Nestin+ NSCs derived from control and Rest mutant ES cells. (A) Background control showing control (REST-100) cells co-stained with the secondary antibodies (Cyc3-anti-mouse/FTTC-anti-chicken) used to visualise the anti-Nestin and anti-Sox1 antibodies respectively. (B-F) Nestin and Sox1 expression on NSCs generated from control and Rest mutant ES cells. In the R5, R2 and R3 areas are classified as Sox1+, Nestin+ and Sox1+/Nestin+ respectively. These NSC populations are summarised in Table 1.

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**Figure S4** FACS analysis of Mash1+ and/or Tau+ NPCs/early neurons from control and Rest mutants. (A) Background control showing control (REST-100) cells co-stained with the secondary antibodies (Cyc3-anti-mouse/FTTC-anti-rabbit) used to visualise the anti-Mash1 and anti-Tau antibodies respectively. (B-F) Mash1 and Tau expression on NPCs/early neurons generated from control and Rest mutant ES cells. Cells in the R5, R2 and R3 areas are classified as Tau+, Mash1+ and Tau+/Mash1+ respectively. These populations of NPCs/early neurons are summarised in Table 1.

Found at: doi:10.1371/journal.pone.0003656.s004 (0.22 MB TIF)

**Figure S5** FACS analysis of Map2+ neurons from control and Rest mutant ES cells. (A) Background control showing control (REST-100) cells stained with the secondary antibody (FTTC-anti-chicken) used to visualise the anti-Mash1 antibody. (B-F) Map2 expression on neurons generated from control and Rest mutant ES cells. Cells in the R5 area are classified as Map2+ mature neurons. This population of mature neurons is summarised in Table I.

Found at: doi:10.1371/journal.pone.0003656.s005 (0.21 MB TIF)

**Figure S6** Constitutively expressing Rest in REST-null ES cells rescues their differentiation defects. By constitutively expressing Rest in REST-null cells (REST-null+REST) using pMT-NRSF, we raised Rest levels to 5% of wild-type levels. We compared the ability of REST-null+REST ES cells to generate NSCs, NPCs and neurons with that of the REST-null ES cells transacted with empty vector (REST-null+MTV) by real-time PCR analysing the expression of stage-specific differentiation markers: Pax6, Mash1 and Nestin to detect neural stem cells (NSCs) (A-C); Ngn1 and Mash1 to detect neural progenitor cells (NPCs) (D, E); Tubb5 to detect young neurons (F) and Snai2a and Llicam to detect mature neurons (G-H). Raising Rest expression level to 8% restored the gene expression levels of stage-specific markers to those observed in the control ES cells. Data are represented as mean±SEM. *P<0.05 and **P<0.01, significantly different from REST-null+REST.

Found at: doi:10.1371/journal.pone.0003656.s006 (0.09 MB TIF)

**Figure S7** Confirmation of REST-null phenotypes in the NSCs derived from Rest knockout mice (Chen et al., 1998). (A) Rest protein expression analysed by Western blot. An alternatively spliced Rest isoform, which has a deletion in the entire exon 2 encoding the N-termius of Rest, can be seen in ES cells derived from Rest+/− and Rest−/− mice. Gapdh was used as an internal control. (B) The expression levels of Rest in wild-type, Rest+/− and Rest−/− ES cells quantified by real-time PCR using primer sets designed against the N-terminus and the C-terminus of Rest gene. Rest is expressed in Rest−/− ES at the wild-type level when primers against the C-terminus of Rest were used, indicating the existence of an alternatively spliced isoform which corroborated the result analysed by Western blot analysis (A). (C-E) The neural stem cells (NSCs) derived from wild-type, Rest+/− and Rest−/− ES cells were identified by Sox1+ (green) and Nestin (red) from a 4-day differentiation. Nuclei were stained with DAPI (blue). The NSCs from Rest−/− ES cells show less Nestin expression in Sox1+ cells as compared to those from wild-type and Rest+/− ES cells, which is resemble to the finding in REST-null mutants (Fig. 3C and 3D). (F-H) Low power and (I-K) high power images of neurons derived from the wild-type, Rest+/− and Rest−/− ES cells. Neurons were identified by NeuN (red) and Map2+ (green) after 14 days of differentiation. Nuclei were counterstained with DAPI (blue). The neurons generated from wild-type and Rest+/− ES cells show similar phenotypes to those derived from REST-100 and REST/KD-50 ES cells (Fig. 3F-G and K-L), whereas the neurons from Rest−/− to those from REST-null ES cells, which are devoid elaborated processes and migration (Fig. 3H-I and M-N).

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**Figure S8** The effect of fibronectin and ploy-D-lysine on REST-null phenotypes. Control (REST-100) and REST-null ES cells were plated onto coverslips coated either with poly-D-lysine or fibronectin. The next day, cells were driven along a neural differentiation pathway in B2N27 medium. After 4 days of differentiation cells were fixed and analysed for the presence of neural stem cells by staining with Sox1 (green) and Nestin (red) (A), NSCs derived from wild-type, REST-100 and REST/KD-50 ES cells show similar phenotypes to those derived from REST-100 and REST/KD-50 ES cells (Fig. 3F-G and K-L), whereas the neurons from Rest−/− to those from REST-null ES cells, which are devoid elaborated processes and migration (Fig. 3H-I and M-N).

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**Methods S1** Supplemental data

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**Author Contributions**

Conceived and designed the experiments: YMS ZFC BPW. Performed the experiments: YMS ZFC SF HHL. Analyzed the data: YMS. Contributed reagents/materials/analysis tools: YMS ZFC. Wrote the paper: YMS. Edited the manuscript: BPW NJB. Financial support: NJB.

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