Canonical MicroRNA Activity Facilitates but May Be Dispensable for Transcription Factor-Mediated Reprogramming

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INTRODUCTION

MicroRNAs (miRNAs) are short, endogenous, non-coding RNAs that repress gene expression post-transcriptionally by destabilizing and/or repressing translation of target mRNAs. In the canonical biogenesis pathway, primary microRNA transcripts (pri-miRNAs) are processed in the nucleus by the microprocessor complex, which consists of the RNase III enzyme DROSHA and the double-stranded RNA-binding protein DGCR8, to generate ~70-nt precursor miRNAs (pre-miRNAs). The pre-miRNAs are then exported to the cytoplasm by EXPORTIN-5 and further processed by another RNase III enzyme, DICER, to generate ~22-nt mature miRNAs (Figure S1) (Kim et al., 2009). More than 400 miRNAs have been identified in the human (Landgraf et al., 2007), and up to 60% of all human genes may be regulated by miRNAs (Friedman et al., 2009).

Given the potentially vast regulatory influence of miRNAs on gene expression and the critical roles of these molecules in embryo development (Bartel, 2009; Sun and Lai, 2013), it is not surprising that miRNAs have emerged as important regulators in reprogramming somatic cells into induced pluripotent stem cells (iPSCs). Together with the Yamanaka factors (OCT4, SOX2, KLF4, and c-MYC) (Takahashi and Yamanaka, 2006), co-expression of the miRNA cluster 302/367 or 106a/363; members of the miR-302, miR-294, or miR-181 family; or miR-93 and miR-106b greatly enhance iPSC derivation efficiency (Judson et al., 2013; Li et al., 2011; Liao et al., 2011; Lin et al., 2011; Subramaniam et al., 2011). Furthermore, expression of the miR-302/367 cluster or miR-200c, miR-302, and miR-369 without the Yamanaka factors is sufficient to reprogram human and mouse fibroblasts (Anokye-Danso et al., 2011; Miyoshi et al., 2011). How these miRNAs promote reprogramming is only partially understood. Several mechanisms have been proposed, such as acceleration of mesenchymal to epithelial transition and antagonism of the activities of let-7 family members, MBD2, NR2F2, and/or other reprogramming suppressors (Hu et al., 2013; Judson et al., 2013; Lee et al., 2013; Liao et al., 2011; Melton et al., 2010).

In addition to the miRNAs that promote reprogramming, several miRNAs that inhibit reprogramming, such as the let-7 family members, have been reported (Melton et al., 2010; Unternaehrer et al., 2014). Therefore, it remains unclear whether miRNA activity as a whole facilitates reprogramming and whether miRNAs are required to convert somatic cells into iPSCs. Previous attempts to reprogram Dicer null mouse embryonic fibroblasts (MEFs) were unsuccessful (Kim et al., 2012); however, this observation cannot rule out a requirement of miRNAs in reprogramming because DICER is also critical for the biogenesis of several other small RNAs, such as endogenous small hairpin RNAs (shRNAs), mirtrons, and endogenous small interfering RNAs (siRNAs) (Figure S1) (Babiarz et al., 2008). In this study, we addressed the question of whether miRNAs are required for generating iPSC by reprogramming mouse cells that lack Dgcr8, a factor required specifically for the biogenesis of canonical miRNAs (Figure S1), including all miRNAs implicated in reprogramming (Babiarz et al., 2008; Judson et al., 2013; Wang et al., 2007). We report that Dgcr8-deficient fibroblasts and NSCs can be reprogrammed by the Yamanaka factors, albeit at decreased
efficiencies. These results demonstrate that while canonical miRNAs as a whole facilitate reprogramming, they may be dispensable for the derivation of iPSCs.

RESULTS

Reprogramming of Dgcr8Δ/Δ MEFs and Tail Tip Fibroblasts

To assess the requirement of miRNAs in iPSC derivation, we first tested whether Dgcr8-deficient MEFs and tail tip fibroblasts (TTFs) could be reprogrammed by Yamanaka factors. Because Dgcr8 null embryos become grossly malformed by embryonic day (E) 6.5 and absorbed by E10 (Wang et al., 2007), isolation of MEFs or TTFs from Dgcr8 null mice was not possible. Instead, we obtained Dgcr8Δ/Δ fibroblasts by Cre-mediated disruption of Dgcr8 in Dgcr8flx/flx MEFs or TTFs (Figure 1A) (Suh et al., 2010; Wang et al., 2007). To monitor Cre activity and enable purification of Dgcr8Δ/Δ fibroblasts, we isolated MEFs or TTFs from Dgcr8flx/flx mice carrying a ROSA26-LoxP-STOP-LoxP-YFP (R26-LSL-YFP) reporter (Srinivas et al., 2001). A previous report demonstrated that mature miRNAs are eliminated in DicerΔ/Δ MEFs by 6 days after transduction of Cre-expressing lentivirus (Kim et al., 2012). To measure the levels of mature miRNAs after Dgcr8 disruption, we performed qPCR analyses on Dgcr8flx/flx and Dgcr8Δ/Δ TTFs 7 and 10 days after Cre expression. Among the miRNAs examined, we found that let-7b, miR-20a, and miR-181a were reliably expressed in Dgcr8flx/flx TTFs, but expression of all three miRNAs was reduced to negligible levels in the Dgcr8Δ/Δ TTFs (Figure 1B), which is consistent with the previous report (Kim et al., 2012). To ensure that only Dgcr8Δ/Δ fibroblasts were used for reprogramming and to exclude those cells that may disrupt Dgcr8 during reprogramming, we sorted YFP+ cells 48 hr after mock (left) or Cre adenovirus (right) transduction. PI, propidium iodide.

(F) PCR genotyping of wild-type, Dgcr8flx/flx TTFs, and Dgcr8Δ/Δ TTF-derived iPSC clones derived from a representative reprogramming experiment. Although most iPSC clones have Dgcr8 disrupted completely, approximately 15% of YFP+ clones, such as iPSC-5, retain one functional Dgcr8 allele. Diamond, Dgcr8+; arrow, Dgcr8flx; arrowhead, Dgcr8Δ.
confirmed that the majority of the resulting iPSCs had both Dgcr8 alleles disrupted; however, approximately 15% of YFP+ iPSCs retained one functional allele of Dgcr8, suggesting that the R26-LSL-YFP reporter is imperfect in monitoring disruption of endogenous genes and that those fibroblasts expressing a single Dgcr8 allele would have a reprogramming advantage (Figure 1F; Table S1).

Reprogramming of Dgcr8<sup>D</sup>/<sup>D</sup> Mouse Neural Stem Cells

Though the miRNAs in Dgcr8<sup>D</sup>/<sup>D</sup> fibroblasts were under the qPCR detection limit (Figure 1B), we could not exclude the possibility that a residual amount of miRNAs remains in a small percentage of fibroblasts 7–10 days after Cre transduction and is required for reprogramming. Dgcr8<sup>D</sup>/<sup>D</sup> fibroblasts quickly deteriorate in culture (data not shown), which precludes long-term passaging to eliminate any residual miRNAs through cell division-mediated dilution and miRNA degradation. In contrast, neural stem cells (NSCs) can be cultured long term in vitro (Andersson et al., 2010; Kawase-Koga et al., 2010), so we used Dgcr8<sup>lox/lox</sup> NSCs to further examine the requirement of miRNAs in reprogramming (Figure 2A). We isolated NSCs from brains of E13.5 Dgcr8<sup>lox/lox</sup>, R26-LSL-YFP mice and disrupted Dgcr8 by transduction of Cre adenovirus (Figure 2A). YFP+ NSCs underwent fluorescence-activated cell sorting (FACS) 48 hr after Cre transduction to exclude cells that had not yet activated Cre. We continuously cultured the sorted Dgcr8<sup>D</sup>/<sup>D</sup> NSCs for 45–60 days (9–12 passages) (Figure 2B) to ensure exhaustion of any residual miRNAs by cell division-mediated dilution and degradation. PCR-based genotyping analysis detected no contamination of cells with incomplete Dgcr8 disruption in the prolonged culture of Dgcr8<sup>D</sup>/<sup>D</sup> NSCs (Figure 2C). The qPCR analysis confirmed that Dgcr8<sup>D</sup>/<sup>D</sup> NSCs did not express mature miRNAs such as miR-20a, miR-181a, let-7b, and miR-9/9*, which are abundantly expressed in Dgcr8<sup>lox/lox</sup> NSCs (Figure 2D). The resulting Dgcr8<sup>D</sup>/<sup>D</sup> NSCs were then transduced with STEMCCA lentivirus to generate iPSCs. The control Dgcr8<sup>lox/lox</sup> NSCs were reprogrammed at an efficiency of 0.5%, which is comparable to published data (Kim et al., 2008). We detected YFP+ iPSC colonies 4 weeks after STEMCCA transduction of Dgcr8<sup>D</sup>/<sup>D</sup> NSCs at efficiencies of 0.01%–0.05% (Figures 2E and 2F). Genotyping of the resulting iPSCs confirmed that Dgcr8 was disrupted in all examined clones (Figure 2C; Figure S2).

Characterization of Dgcr8<sup>D</sup>/<sup>D</sup> iPSCs

The Dgcr8<sup>D</sup>/<sup>D</sup> iPSCs derived from fibroblasts or NSCs expressed pluripotency-associated markers such as alkaline phosphatase (AP), SSEA-1, and NANOG (Figures 3A–3C; Figure S3A). The qPCR analysis confirmed the lack of Dgcr8 expression in Dgcr8<sup>D</sup>/<sup>D</sup> iPSCs (Figure 3D). Karyotyping analyses demonstrated that normal Dgcr8<sup>D</sup>/<sup>D</sup> iPSCs could be isolated (Figure 3E; Figures S3B and S3C). The qPCR analyses revealed that Yamanaka factors delivered
by the STEMCCA lentivirus were largely silenced in Dgcr8<sup>Δ/Δ</sup> iPSCs (Figure 3F). Furthermore, transgene-free Dgcr8<sup>Δ/Δ</sup> iPSCs could be isolated and stably maintained after removal of the STEMCCA lentivirus by Cre adenovirus transduction (Figure 3G; Figure S3D) (Somers et al., 2010).

Next, we evaluated the differentiation capacity of Dgcr8<sup>Δ/Δ</sup> iPSCs in embryoid bodies (EBs). EBs of Dgcr8<sup>Δ/Δ</sup> iPSCs failed to form cystic cavities over an 11-day period, suggesting a lack of differentiation (Figures 4A and 4B). The qPCR analyses revealed that pluripotency-associated markers Oct4 and Nanog were maintained but lineage-specific markers, such as Fgf5 and Krt18 (ectodermal), Brachyury (mesodermal), Afp and Hnf4a (endodermal), and Eomes (extraembryonic), were weakly expressed or absent in EBs of Dgcr8<sup>Δ/Δ</sup> iPSCs. The only gene modestly upregulated in EBs of Dgcr8<sup>Δ/Δ</sup> iPSCs was Sox1 (Figure 4C), which is expressed by neural progenitor cells (Ying et al., 2003). To test whether Dgcr8<sup>Δ/Δ</sup> iPSCs could differentiate into more mature neuronal cells, we extended the differentiation protocol under pro-neuronal conditions. Unlike wild-type embryonic stem cells (ESCs), mature TuJ1+ neurons were not differentiated from Dgcr8<sup>Δ/Δ</sup> iPSCs (Figures 4D and 4E).

These data are consistent with the previous finding that Dgcr8<sup>−/−</sup> ESCs poorly produce mature somatic cells (Wang et al., 2007).

Next, we restored DGCR8 expression to levels similar to wild-type ESCs using a human DGCR8 cDNA (Figure 5A). The DGCR8-rescued iPSCs exhibited an accelerated cell cycle with a shortened G1 phase compared to Dgcr8<sup>Δ/Δ</sup> iPSCs (Figure 5B), which underwent slower proliferation, similar to Dgcr8<sup>Δ/Δ</sup> ESCs (Wang et al., 2008). To test whether DGCR8 rescue restored the differentiation potential of the Dgcr8<sup>Δ/Δ</sup> iPSCs, we performed a colony-forming assay to examine the number of differentiation-resistant cells within the Dgcr8<sup>Δ/Δ</sup> and DGCR8-rescued iPSCs. Mutant and rescued iPSCs were first induced to differentiation by retinoic acid and then plated back to conditions supporting self-renewal of iPSCs to form colonies. We found that significantly more colonies were formed by Dgcr8<sup>Δ/Δ</sup> iPSCs than by wild-type control ESCs and DGCR8-rescued iPSCs (Figure 5C). We further evaluated the differentiation potential of rescued iPSCs in a teratoma assay. When injected into immunodeficient mice, the Dgcr8<sup>Δ/Δ</sup> iPSCs formed tumors containing predominantly undifferentiated cells.
In contrast, the teratoma formed by DGCR8-rescued iPSCs consisted of tissues from all three embryonic layers (Figures 5E–5E*).

Together, our data support that somatic cells lacking Dgcr8 and deficient in the biogenesis of canonical miRNAs can be reprogrammed into iPSCs by the Yamanaka factors alone, albeit at decreased reprogramming efficiencies; therefore, canonical miRNA activity facilitates but may be dispensable for iPSC derivation. Consistent with previous reports (Kanellopoulou et al., 2005; Wang et al., 2007), however, miRNAs do appear to be important for subsequent iPSC-derived tissue differentiation.

**DISCUSSION**

miRNAs may confer robustness to biological systems by integrating into transcriptional regulatory circuitry to reinforce genetic programs and buffer stochastic perturbations (Ebert and Sharp, 2012; Hornstein and Shomron, 2006). Mutant mice with deletions of individual miRNA clusters often exhibit only relatively subtle phenotypic defects (Park et al., 2012). More severe phenotypes are usually observed in mutants with compound deletions of functionally redundant miRNA clusters, suggesting that the subtle defects of individual mutations are at least partially due to functional compensation (Park et al., 2012). The Dgcr8 and Dicer mutants, which have complete miRNA loss, exhibit the most extreme phenotypic defects. The mutant ESCs can self-renew and express stem cell markers but are functionally defective in spontaneous differentiation (Kanellopoulou et al., 2005; Wang et al., 2007). These results suggest that the regulatory circuitry of pluripotent cells can be sustained solely by transcription factors, while miRNAs are required to initiate and/or sustain the differentiation. Our data support this notion. Because reprogramming is generally considered to be a de-differentiation process, our data suggest that miRNA activity may not be essential for de-differentiation but is essential for normal tissue differentiation.
The mechanisms involved in reprogramming somatic cells to iPSCs by the Yamanaka factors remain poorly understood. Because of the low efficiency and slow kinetics of most reprogramming systems, molecular events that direct somatic cells to pluripotency have been difficult to define. Recent work has demonstrated that miRNAs such as miR-294, miR-302, and miR-181 family members facilitate (Judson et al., 2013; Li et al., 2011; Liao et al., 2011; Lin et al., 2011; Melton et al., 2010; Subramanyam et al., 2011), but let-7 family members inhibit, reprogramming (Melton et al., 2010; Unternaehrer et al., 2014). Therefore, it remains unclear whether miRNA activity as a whole promotes reprogramming and whether miRNAs, in particular those miRNAs shown to promote reprogramming, are necessary for the derivation of iPSCs. Here, we present data demonstrating that while miRNA activity as a whole promotes reprogramming, the derivation of iPSCs may be achieved without any miRNA activities. Kim et al. (2012) reported that iPSCs could not be isolated from MEFs 6 days after disruption of Dicer, which is inconsistent with our data on reprogramming Dgcr8<sup>-/-</sup> fibroblasts (Figure 1). DICER is required for the biogenesis of Cre-expressing lentivirus (Kim et al., 2012). Nevertheless, to exclude the possibility that residual miRNAs may be present and essential for reprogramming, we reprogrammed Dgcr8<sup>-/-</sup> NSCs, which can be propagated for longer terms to ensure exhaustion of residual miRNAs before transduction of reprogramming factors (Figure 2A). The prolonged culture of Dgcr8<sup>-/-</sup> NSCs exhausts residual miRNAs by two mechanisms. First, the Dgcr8<sup>-/-</sup> NSCs are proliferative; therefore, residual miRNAs are diluted out with each cell division. We split Dgcr8<sup>-/-</sup> NSCs at a 1:5 ratio for each passage, resulting in the expansion of any single cell to 1.9<sup>±</sup>10<sup>6</sup>–2.4<sup>±</sup>10<sup>8</sup> (59–512) progeny cells and making it highly unlikely that any residual miRNAs could persist at a biological meaningful concentration by the end of 9–12 passages. Second, the sorted Dgcr8<sup>-/-</sup> NSCs were reprogrammed after a continuous culture for 45–60 days, which is a sufficient duration to achieve complete degradation of residual miRNAs. Therefore, our data conclusively demonstrate that reprogramming of NSCs may be achieved solely by transcriptional factors without any miRNA activities.

Kim et al. (2012) reported that iPSCs could not be isolated from MEFs 6 days after disruption of Dicer, which is inconsistent with our data on reprogramming Dgcr8<sup>-/-</sup> fibroblasts (Figure 1). DICER is required for the biogenesis of
not only canonical miRNAs but also other small RNA species, such as endogenous siRNAs, shRNAs, mirtrons, and short interspersed nuclear element-derived RNAs (Figure S1) (Babiarz et al., 2008). The discrepancy between the data on reprogramming of Dicer-deficient cells and those of Dgcr8-deficient cells probably reflects the activities of some DICER-dependent but DGCR8-independent small RNAs. Alternatively, the poorer proliferation capacity of Dicer<sup>−/−</sup>/A fibroblasts may contribute to the failure of iPSC derivation (Kim et al., 2012), which is known to be proliferation dependent (Smith et al., 2010). Recently, Zhang et al. (2013) reported that they were unable to isolate iPSCs from human foreskin fibroblasts that were null for the endogenous miR-302/367 cluster. These data suggested the miR-302/367 cluster is required for human somatic cell reprogramming. Although this result is not consistent with our findings, the discrepancy may be explained by the potential difference in somatic cell reprogramming and/or in the self-renewal of human and mouse pluripotent stem cells (Nichols and Smith, 2009). Alternatively, the discrepancy may be caused by the different miRNA deficiencies of the reprogrammed fibroblasts. In our study, the Dgcr8<sup>−/−</sup> fibroblasts lacked miRNAs both promoting reprogramming, such as the miR-290s and miR-302s, and inhibiting reprogramming, such as the let-7s; however, the fibroblasts used by Zhang et al. (2013) were only deficient in the reprogramming-promoting miR-302/367 cluster. The fine balance between pluripotency-promoting and differentiation-inducing miRNAs has been demonstrated to play critical roles in the maintenance of the ground state of pluripotency (Kumar et al., 2014), which could be similarly required in reprogramming. Nonetheless, this is an interesting observation that deserves further investigation.

**EXPERIMENTAL PROCEDURES**

**Mice and the Derivation of ESCs, MEFs, TTFs, and NSCs**  
All animal experiments were performed in accordance with guidelines from the University of Alabama at Birmingham (UAB) and NIH. Dgcr8<sup>flox/flox</sup>; LSL-YFP mice were generated by crossing Dgcr8<sup>flox/flox</sup> mice (Wang et al., 2007) and R26-LSL-YFP mice (Srinivas et al., 2001). ESCs were derived from E3.5 blastocysts as described (Kim et al., 2010; Liu et al., 2011). MEFs were isolated from E12.5 embryos, and TTFs were derived from 1-week-old mice. NSCs were isolated from brains of E13.5 embryos following from E12.5 embryos, and TTFs were derived from 1-week-old mice (vas et al., 2001). ESCs were derived from E3.5 blastocysts as described previously (Kim et al., 2010). For EB differentiation, trypsinized wild-type or mutant iPSCs were suspended in Costar ultra-low-attachment cell culture plates (Corning) at a density of 1 × 10<sup>5</sup> cells/ml in differentiation medium (ESC maintenance medium without ESGRO). EB samples were collected on the indicated days for total RNA extraction. For neuronal differentiation, EBs (day 4) were plated onto tissue culture plates and cultured in N2 medium (DMEM/F12 and N2 supplement [Gemini Bio]) for up to 25 days. All fibroblasts were cultured in D10 medium (DMEM and 10% FBS). NSCs were cultured in Mouse Neural Stem Cell Expansion medium (EMD Millipore) on tissue culture plates coated with polyornithine (Sigma-Aldrich) and laminin (EMD Millipore).

**Lentiviral Production, iPSC Derivation, and Rescue of Dgcr8 Deficiency**  
Lentivirus expressing STEMCCA (Somers et al., 2010) were prepared as described (Zhao et al., 2014). Dgcr8<sup>−/−</sup> fibroblasts or NSCs were obtained by Cre adenovirus (Vector Biolabs) transduction of Dgcr8<sup>flox/flox</sup>, LSL-YFP MEFs or TTFs at an MOI of 500 and FACS sorting of YFP<sup>+</sup> cells 48 hr after viral transduction. For fibroblast reprogramming, sorted MEFs or TTFs were continuously cultured for seven or ten days before transduction with STEMCCA lentivirus at an MOI of 2. For NSC reprogramming, sorted NSCs were continuously cultured for 45–60 days before transduction with STEMCCA lentivirus at an MOI of 2. The transduced fibroblasts or NSCs were plated directly onto irradiated MEF feeders in mouse ESC maintenance medium (DMEM, 15% FBS, and 1,000 U/ml ESGRO [Millipore]) for up to 4 or 6 weeks, respectively. A human Dgcr8 cDNA was subcloned from pFLAG/HA-DGCR8 (Addgene 10921) (Landthaler et al., 2004) into pSIN-EF2-DEST-Pur, a derivative of pSIN-EF2-Oct4-Pur (Addgene 16579) (Yu et al., 2007), to generate the lentiviral vector pSIN-EF2-DGCR8-Pur. Dgcr8-deficient iPSCs were transduced with lentivirus expressing Dgcr8 and selected for puromycin resistance.

**Immunostaining, Immunoblotting, and AP Staining**  
For immunostaining, iPSCs or EBs were fixed in 4% paraformaldehyde, blocked in Protein Block (Dako), and incubated with the appropriate primary antibodies overnight at 4 °C and secondary antibodies for 2 hr at room temperature. Images were acquired by a Nikon Ti-S microscope and processed by Photoshop CS6. For immunoblotting, whole cell extracts were prepared in RIPA buffer (50 mM Tris-HCl [pH 8.0], 150 mM NaCl, 1% NP-40, 0.5% sodium deoxycholate, and 0.1% SDS), separated on a 4%–20% SDS-polyacrylamide gel (Bio-Rad), and transferred to polyvinylidene fluoride membrane (Thermo Scientific). Antibodies used were Dgcr8 (10996-1-AP, ProteinTech), GAPDH (sc-25778, Santa Cruz), SSEA-1 (MC-480, Hybridoma Bank), NANOG (AF2729, R&D Systems), and Tuj1 (801202, BioLegend). For AP staining, cells were fixed in 4% paraformaldehyde and stained using the Leukocyte Alkaline Phosphatase Kit (Sigma-Aldrich).

**Genotyping, Karyotyping, and Teratoma Analysis**  
Genotyping was performed as described (Suh et al., 2010). All cell lines were submitted to Cell Line Genetics for G-band karyotyping. Non-obese diabetic severe combined immunodeficiency
gamma mice 4–8 weeks of age were injected subcutaneously with $1 \times 10^5$–$5 \times 10^5$ iPSCs. Tumors were harvested, fixed with 10% formalin, and processed by the Comparative Pathology Laboratory at UAB or by Histowiz.

**RNA Extraction and qPCR Analyses**

Total RNA was isolated with the DirectZol RNA Kit (Zymo Research), and cDNA were synthesized using the Verso cDNA Synthesis Kit (Thermo Scientific). The qPCR was performed using 2x Absolute Blue qPCR Master Mix (Thermo Scientific) on a Viia 7 real-time PCR system (Life Technologies). Primers are listed in Table S2. The miRNAs were reverse transcribed using the TaqMan MicroRNA Reverse Transcription Kit (Life Technologies). The qPCR was performed using the TaqMan Universal PCR Master Mix and TaqMan MicroRNA Assays for indicated miRNAs (Life Technologies) on the Viia 7 system.

**Colonies-Forming Assay**

The colony-forming assay was performed as previously described (Wang et al., 2007). In brief, undifferentiated wild-type, Dgcr8–/–, and rescued Dgcr8+/– iPSCs were cultured in differentiation medium supplemented with 2 μM retinoic acid (Sigma-Aldrich) for the indicated days, trypsinized to single cells, replated at a density of 100 cells/cm² onto gelatin-coated plates, and cultured in ESC maintenance medium for 7 days before AP staining. Experiments were repeated three times, and only AP-positive colonies were scored.

**Cell-Cycle Analysis**

Cell-cycle analysis was performed as described (Zhao et al., 2014). In brief, cells at 30%–50% confluency were trypsinized and fixed in cold 70% ethanol at $-20\,^\circ C$ overnight. Cells were washed twice in PBS, treated with 10 μg/ml DNase-free RNase A at 37°C for 30 min, and resuspended at a density of $5 \times 10^5$ cells/ml in PBS with 5 μg/ml propidium iodide. Cells were analyzed on a Becton Dickinson Fortessa flow cytometer, and data were analyzed by the FlowJo VX software.

**SUPPLEMENTAL INFORMATION**

Supplemental Information includes three figures and two tables and can be found with this article online at [http://dx.doi.org/10.1016/j.stemcr.2015.11.002](http://dx.doi.org/10.1016/j.stemcr.2015.11.002).

**ACKNOWLEDGMENTS**

We thank Drs. Robert Bléloch (UCSF) for the Dgcr8–/– ESCs, Jinyang Feng (Loyola Medical Center) for interpretation of histology results, Tim Townes (UAB) for comments on the manuscript, and the UAB Comprehensive Flow Cytometry Core (P30AR048311 and P30AI027767) for cell sorting. R.Z. is supported by UAB Developmental Fund, and K.K. is supported by the NIH (5R00HL093212-04 and R01AG043531), TriStem (2013-049), MSK Cancer Center Support Grant/Core Grant (P30CA008748), Louis V. Gerstner Jr. Young Investigators awards, Geoffrey Beebe Junior Chair Award, Alfred W. Bressler Scholars Endowment Fund, and Sidney Kimmel Scholars Award.

Received: March 6, 2014
Revised: November 3, 2015
Accepted: November 12, 2015
Published: December 8, 2015

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