Lineage tracing of neuromesodermal progenitors reveals novel Wnt-dependent roles in trunk progenitor cell maintenance and differentiation

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ABSTRACT

In the development of the vertebrate body plan, Wnt3a is thought to promote the formation of paraxial mesodermal progenitors (PMPs) of the trunk region while suppressing neural specification. Recent lineage-tracing experiments have demonstrated that these trunk neural progenitors and PMPs derive from a common multipotent progenitor called the neuromesodermal progenitor (NMP). NMPs are known to reside in the anterior primitive streak (PS) region; however, the extent to which NMPs populate the PS and contribute to the vertebrate body plan, and the precise role that Wnt3a plays in regulating NMP self-renewal and differentiation are unclear. To address this, we used cell-specific markers (Sox2 and T) and tamoxifen-induced Cre recombinase-based lineage tracing to locate putative NMPs in vivo. We provide functional evidence for NMP location primarily in the epithelial PS, and to a lesser degree in the ingressed PS. Lineage-tracing studies in Wnt3a/β-catenin signaling pathway mutants provide genetic evidence that trunk progenitors normally fated to enter the mesodermal germ layer can be redirected towards the neural lineage. These data, combined with previous PS lineage-tracing studies, support a model that epithelial anterior PS cells are Sox2/T+ multipotent NMPs and form the bulk of neural progenitors and PMPs of the posterior trunk region. Finally, we find that Wnt3a/β-catenin signaling directs trunk progenitors towards PMP fates; however, our data also suggest that Wnt3a positively supports a progenitor state for both mesodermal and neural progenitors.

KEY WORDS: Neural progenitor, Neuromesodermal progenitor, Wnt signaling, Brachyury, Paraxial mesoderm, Sox2

INTRODUCTION

How progenitor populations are directed to differentiate into different cell lineages is poorly understood in vivo, particularly for the embryonic progenitors of the neural and mesodermal lineages. Neuroectoderm and mesoderm progenitors were originally thought of as separate descendants of the pluripotent epiblast during early gastrulation (Bellairs, 1986; Garcia-Martinez and Schoenwolf, 1992; Tam and Behringer, 1997). However, in mammalian embryos, a common neuromesodermal progenitor (NMP) is thought to give rise to both neural and paraxial mesoderm progenitors (PMPs) (Cambray and Wilson, 2002, 2007; Tzouanacou et al., 2009; Wilson et al., 2009). Regional PS progenitors show the capacity to contribute to multiple cell types along the body axis and can be passaged through multiple successive host embryos, suggesting that the PS region contains stem cell populations (Cambray and Wilson, 2002; Wilson et al., 2009). However, recognizing axial stem cells in vivo by molecular criteria remains a challenge as the PS region is a site of dynamic and continuous cell movement with different, indistinguishable populations of cells entering, exiting and residing in the PS (Garcia-Martinez and Schoenwolf, 1992; Schoenwolf et al., 1992; Wilson and Beddington, 1996). NMPs are proposed to co-express Sox2, a pluripotency and neural progenitor marker, and the PS marker T, also known as brachyury (Martin and Kimelman, 2012; Wilson et al., 2009). Although Sox2/T co-expressing regions have been documented in the PS region in mammals (Tsakiridis et al., 2014), no study has demonstrated that Sox2/T co-expression functionally represents multipotent NMPs.

One way to address whether T-expressing cells include NMPs is to take a gene-specific transgenic lineage-tracking approach. The recently developed TCreERT2 transgenic line, in which the tamoxifen (TAM)-inducible CreERT2 recombinase is driven by the T promoter, has the potential to trace NMPs in vivo as it is expressed in the anterior epithelial PS region where NMPs are thought to exist (Anderson et al., 2013; Wilson et al., 2009). TCreERT2 also has the advantage of excluding pluripotent epiblast stem cells that are capable of giving rise to most embryonic cell types, including neural and mesodermal cells. Previous transgenic tracing experiments have successfully used TAM-inducible Cre-based transgenics to follow the population dynamics of diverse progenitor populations in vivo (Boyle et al., 2008; Göhert et al., 2005; Masahira et al., 2006; Schepers et al., 2012; Srinivasan et al., 2007).

Wnt3a has been proposed to be a crucial regulator of NMP maintenance and differentiation, and presumably does so through β-catenin/Tcf transcriptional complexes and the subsequent activation of downstream target genes (Cleverson and Nusse, 2012) such as T (Yamaguchi et al., 1999). In the absence of Wnt3a or T, the embryo fails to generate the posterior-most ~55 of 65 skeletal elements/vertebrae, resulting in a severe posterior axis truncation (Herrmann, 1992; Takada et al., 1994). The loss of
skeletal elements corresponds to a reduction in progenitor markers, indicating a collapse in the progenitor populations that build the embryonic trunk (Yamaguchi et al., 1999). Importantly, Wnt3a, Tcf1;Lef1 double mutants, and T mutant embryos show an expansion of neural tissue in the form of an ectopic neural tube, giving support to the hypothesis that the Wnt3a/β-catenin/Tcf1-Lef1/T axis is directly regulating the differentiation of NMPs into neural or mesodermal progenitors (Galceran et al., 1999; Herrmann, 1992; Yamaguchi et al., 1999; Yoshikawa et al., 1997). Single cell studies in zebrafish further this argument by showing that embryonic progenitors will selectively form striated skeletal muscle tissue when exposed to high Wnt signaling and, by contrast, form neural tissue when Wnt signaling is inhibited (Martin and Kimelman, 2012). From these, and other studies, a model of Wnt3a function has evolved to incorporate the concept of the NMP (Fig. 1A) (Galceran et al., 1999; Li and Storey, 2011; Martin and Kimelman, 2008, 2012; Takada et al., 1994; Yamaguchi et al., 1999; Yoshikawa et al., 1997). This model predicts that Wnt has a direct role in NMP maintenance, inducing PM cell fate and repressing neural cell fate. However, the model remains hypothetical and has not been directly tested in the native mammalian niche. The Wnt3a-null and conditional Ctnnb1 (β-catenin) mouse mutants provide excellent opportunities to study NMP behavior in vivo as the fate of these cells can be dramatically modulated through these single gene mutations.

RESULTS

Imbalance of neural progenitors and PMPs, and differentiated descendants in Wnt3a−/− mutants

To assess neural progenitors and PMPs in Wnt3a−/− mutants, we examined representative markers of each population in sections taken just posterior to the forelimb, corresponding to the level of the 13th-16th somite (s13-16) of wild-type E9.5 embryos. Phenotypic differences are clearly evident between control and Wnt3a−/− mutant embryos at this stage and axial level (Takada et al., 1994; Yoshikawa et al., 1997). E9.5 Wnt3a−/− mutant embryos showed enlarged and malformed neural tissue evident by detection of the neural progenitor marker Sox2, while fibronectin expression revealed a reduced PM (Fig. 1B). Quantification showed that Wnt3a−/− mutants had significantly more neural progenitors and fewer...
mesodermal cells than control embryos (Fig. 1C). At later stages of development, this phenotype manifests itself as a malformed neural tube and loss of musculoskeletal tissues, among others (supplementary material Fig. S1).

Although the observed progenitor imbalance can be explained by fate changes of NMPs, Wnt-dependent differential progenitor growth could also account for the phenotype. For example, Wnt3a could act to promote mesodermal growth and inhibit neural progenitor growth. To test this, we examined the cell proliferation rates of differentiating neural and mesodermal cells by detection of p-Histone H3 (Fig. 1D). The rate of cell proliferation was not significantly different in neural versus mesodermal populations, comparing littermate control embryos with Wnt3a−/− mutants, despite the apparent loss of PM tissue and an increase in neural tissue in Wnt3a−/− mutants at E9.5 (Fig. 1C; supplementary material Fig. S1). The only significant difference was the proliferation rate of the ectopic neural tissue in Wnt3a−/− mutants, which was lower than controls in or in Wnt3a−/− primary neural tube (Fig. 1E). We also observed no difference in neural versus mesodermal proliferation rates between control and Wnt3a−/− mutants in the E8.5 trunk region (supplementary material Fig. S1). These data suggest that differential neural and mesodermal growth is insufficient to explain the Wnt3a-dependent progenitor imbalance.

Detection of Sox2 and T co-expressing cells in the primitive streak

The PS is the source of progenitors for the trunk region during the extension of the body axis. In Wnt3a−/− mutants, changes in the fate of NMPs residing in the streak are presumably responsible for the observed imbalances in neural and mesodermal cell types in the trunk. To localize these putative NMPs by immunohistochemistry in wild-type embryos at E8.5, we used antibodies to Sox2 and T that are thought to mark the NMP. Sox2 was detected in the epithelial PS and, to a much lower degree, in ingressed cells of anterior, but not posterior, regions of the PS. T was strongly expressed in the epithelial and the ingressed regions of the PS along the anterior-posterior (A-P) extent of the PS. Co-detection of Sox2 and T was primarily observed in the epithelial PS, and to a lesser degree in ingressed cells (Fig. 1F), but was not observed near the notochord/node transition, where Sox2 and T marked distinct cell populations (supplementary material Fig. S1). Although our data indicate zones of putative NMPs, it is unclear whether these Sox2+ T+ populations retain the potential to give rise to both neural and mesodermal cells.

Tracing trunk progenitors in Wnt3a−/− mutants

To determine whether NMPs are the source of the cells that contribute to the ectopic neural tube in Wnt3a−/− mutants, T+PS cells were lineage traced. Wnt3a−/− PS cells labeled on E7.5 with TCreERT2 and examined at E8.5 showed more labeled cells in the neural midline region of the mutant trunk than in control embryos (supplementary material Fig. S3) (n=9 control, n=3 Wnt3a−/− embryos). Analysis of the trunk region (s13-16) at E9.5 showed an increased proportion of labeled cells in the ectopic neural tube of Wnt3a−/− embryos relative to the primary neural tube (Fig. 2B,C) (n=18 control, n=7 Wnt3a−/− embryos). Transverse sections of three TCreERT2-traced wild-type embryos showed that roughly 20% of neural tube cells were labeled within the s13-16 region, whereas ~80% of PM cells were traced (Fig. 2D). In three sectioned Wnt3a−/− mutants, the ectopic neural tissue showed more than 60% TCreERT2-labeled cells, a significantly higher number than observed in the primary neural tube and comparable with the PM (Fig. 2E). These data suggest that wild-type TCreERT2-labeled cells differentiate into neural progenitors and PM, among others, but, in the absence of Wnt3a, cells normally fated for PM change their fate and differentiate into neural tissue. However, as the TCreERT2 lineage also included other mesodermal and gut progenitors, a more-refined analysis was required to determine whether the ectopic neural tube in Wnt3a−/− mutants derived solely from the epithelial PS population.

To distinguish the Sox2+T+ epithelial PS lineage from the Sox2+T− mesoderm and hindgut lineages (all of which are labeled with TCreERT2), we used Cited1-CreERT2 (Boyle et al., 2008), which labels only mesodermal (and hindgut) progenitors and not the epithelial PS. Following the same protocol described above for the TCreERT2-based tracings, Cited1-CreERT2 mice were crossed to R26R-lacZ and cell-tracing studies were initiated with an injection of TAM at E7.5. βGal detection was performed on E8.5-E10.5 embryos to identify Cited1-CreERT2-labeled cells (Fig. 2; supplementary material Fig. S2). As expected, wild-type cells labeled with Cited1-CreERT2 were not found in the Sox2+ epithelial PS at E8.5 but were detectable in mesodermal cells that had already ingressed, including Tbx6+ PM (Fig. 2F; supplementary material Fig. S2). Importantly, the Cited1-CreERT2 did not trace neural tissue; however, labeled cells were maintained throughout the extending body axis, including the posterior-most tailbud progenitor zone, days after a single pulse of TAM (Fig. 2G; supplementary material Fig. S2). This result is consistent with Cited1-CreERT2 labeling a mesodermal progenitor population. Tracing Cited1-CreERT2-labeled cells in Wnt3a−/− mutant embryos showed a persistence of labeled cells in the trunk region at E9.5
Histological analysis of Wnt3a−/− embryos (n=3) showed that only a few Cited1-CreERT2-labeled cells were present in the ectopic neural tube of Wnt3a−/− embryos (Fig. 2G). Typically two or three cells were observed in the ectopic neural tube per section of the mutant embryo. Considering that the ectopic neural tube contains 137.2±16.2 (s.d.) cells per section this represents roughly 2% of the ectopic neural tube cells. These data show that for cells originally destined to become mesoderm, only very few of them can change their fate, enter the neural cell lineage and contribute to the Wnt3a−/− ectopic neural tube, after they have ingressed through the PS. The subtractive comparison between TCreERT2 and Cited1-CreERT2 tracings suggests that T+ cells (including NMPs) originating in the epithelial PS, and not the ingressed PS, primarily contribute to the ectopic neural tube.

To show that the observed change of fate from mesoderm to neural is unique to PS cells, we also examined the Shh-Cre traced gut and notochord lineages (Harfe et al., 2004). Examination of Shh-Cre traced cells in Wnt3a−/− mutant embryos (n=3) showed no labeled cells occurring in the ectopic neural tube, demonstrating that NMPs do not reside in Shh-expressing notochord or gut progenitors (supplementary material Fig. S2).

**Tracing TCreERT2-labeled cells in β-catenin loss-of-function mutants**

Although Wnt3a signals through β-catenin, previous conditional deletion of Ctnnb1 (β-catenin) in posterior progenitors with the non-inducible T-Cre transgenic mouse resulted in a severe loss of all trunk progenitors with no apparent ectopic neural tubes (Dunty et al., 2008). To bypass this early requirement for Ctnnb1 and test whether NMP fate decisions are β-catenin dependent, we used the TAM-inducible TCreERT2 transgene to conditionally delete Ctnnb1 (βcatLOF) (Fig. 3). TAM was administered at E6.5, not at E7.5 as in previous tracing experiments, to ensure that endogenous β-catenin protein was depleted after excision of the conditional βcatLOF allele. At E9.5 the conditional βcatLOF mutants showed a malformed posterior trunk region (n=13/13) reminiscent of a Wnt3a−/− mutant, with an absence of T-expressing trunk progenitors and defective somitogenesis (n=5/5) (Fig. 3A). The detection of small aggregates of somite cells can be explained by incomplete excision of β-catenin, as suggested by immunofluorescence (supplementary material Fig. S4). Histological analysis of TCreERT2;βcatLOF mutants (n=3) revealed enlarged and malformed neural tissue containing increased numbers of Sox2+ neural tissue descendants (TAM administered at E7.5) are absent from the E8.5 epithelial PS but overlap with Tbx6+ ingressed streak cells. (G) βGal-stained Cited1-CreERT2 descendants (TAM at E7.5) in E9.5 control embryos show no neural contribution, and only a few traced cells in the Wnt3a−/− ectopic neural tube (arrows), es, epithelial PS; is, ingressed PS; nt, neural tube; ect. nt, ectopic neural tube. Arrows in G indicate Cited1-CreERT2 descendants in ectopic neural tube.

**Fig. 2. Tracing trunk progenitors in Wnt3a−/− mutants using CreERT2;R26R-lacZ reporters.**

(A) E8.5 embryo showing TCreERT2-traced βGal+ cells and Sox2+ cells (presumptive NMPs) in the epithelial PS (es), and with Tbx6+ cells in the ingressed PS (is). (B) βGal-stained TCreERT2 descendants (TAM administered at E7.5) in E9.5 control and Wnt3a−/− embryos showing many βGal+ cells in the ectopic neural tube of Wnt3a−/− mutants (sections through the s13-16 region). (C) Detection of βGal and DAPI to determine the number of TCreERT2-traced cells in neural and mesodermal tissues at E9.5 used in D,E. (D) Quantification of TCreERT2-labeled cells in neural versus PM tissues. Cell counts from sampled areas of three embryos. (E) Quantification of TCreERT2-labeled cells in the primary and ectopic neural tube of Wnt3a−/− mutants (n=3 control, n=3 Wnt3a−/−). Data are mean±s.e.m. Student’s t-test. (F) βGal+ Cited1-CreERT2 mutants showing many Wnt3a−/−/− mutants containing an enlarged and malformed neural tube (βGal+ Cited1-CreERT2;R26R-lacZ).
β-Catenin gain-of-function is compatible with both neural and mesodermal progenitor fates

Previously it has been shown in zebrafish that activation of Wnt3a/β-catenin signaling drives a cell to a mesodermal fate and inhibits the adoption of neural fate in a non-native niche (Martin and Kimelman, 2012), as modeled in Fig. 1A. To directly test whether Wnt signaling inhibits neural cell fate and promotes mesodermal cell fate in mammals, we increased Wnt signaling in NMPs in the native niche, using TCreERT2 to conditionally express a dominant stabilized β-catenin (Ctnnb1lox(ex3)βcatGOF, hereafter referred to as βcatGOF) (Harada et al., 1999), and simultaneously trace these cells with the R26R- βGal reporter. TAM was administered at E6.5 for uniformity with the βcatGOF studies. The resulting TCreERT2; βcatGOF mutant embryos showed similar elongation of the trunk region compared with control embryos but had an irregular segmental pattern (n=31 controls, n=22 mutants). Surprisingly, section analysis of TCreERT2; βcatGOF embryos (n=3) showed a similar distribution of TCreERT2; βcatGOF-labeled cells in neural and mesodermal tissues at E10.5 to controls (n=3) (Fig. 4A). Examination of TCreERT2-traced cells for Sox2 expression showed that βcatGOF NMPs were able to enter the neural cell fate and contribute to the neural tube (Fig. 4A). Additionally, a Wnt/β-catenin reporter transgene [BATlacZ (Nakaya et al., 2005)] showed ectopic expression in the ventral neural tube of sectioned βcatGOF embryos (n=3) that was not observed in control embryos (n=3), suggesting that TCreERT2-labeled cells can differentiate into the neural lineage in the presence of elevated Wnt signaling (Fig. 4B, arrows). Only a small proportion of Wnt-reporting cells were distinctively stained in the ventral neural tube, likely due to the lower Cre-excision/tracing of TCreERT2 cells in the neural tube (supplementary material Fig. S2). We also observed spheroid-shaped aggregates of Wnt reporter-responding cells throughout the trunk region of mutants (n=7) that were not observed in controls (n=8) (Fig. 4B,C). The observation that Wnt does not block entry into the neural lineage contrasts with previous reports (Martin and Kimelman, 2012; Yamaguchi et al., 1999; Yoshikawa et al., 1997); however, these data are consistent with the sustained detection of Wnt signaling in the neuroepithelium of BATlacZ embryos, including the epithelial PS, neuroepithelial folds and neural tube (supplementary material Fig. S5).

TCreERT2; βcatGOF embryos elongate surprisingly normally up to E10.5 with no obvious truncation of the embryonic axis (Fig. 4B). Analysis of T and Sox2 in sectioned PS regions at E8.5 showed both proteins were easily detected in the PS of βcatGOF embryos (n=3) (Fig. 4D). However, the morphology of the βcatGOF PS was abnormal with a broader and thinner epithelium (Fig. 4D). βcatGOF embryos also showed increased expression of T, consistent with the known role of T as a transcriptional target of Wnt/β-catenin signaling (Yamaguchi et al., 1999). Combined, these data are consistent with Wnt signals maintaining multipotent NMPs in vivo, as recently demonstrated in vitro (Tsakiridis et al., 2014).

Notably, the most prominent phenotype of constitutive Wnt/β-catenin signaling in the trunk region of the TCreERT2; βcatGOF embryos was the formation of aggregates of undifferentiated progenitors along the length of the embryo (Fig. 4C). These aggregates were similar histologically to PS progenitors and, like PS tissue, highly responsive to Wnt signaling (Fig. 4C). To determine which progenitors were being promoted by Wnt/β-catenin signaling, markers of NMPs (Sox2+/T+), PMPs (Tbx6+/T+) and presomitic mesoderm (Tbx6−T+) were used in immunofluorescent studies of sectioned βcatGOF embryos (n=3) (Fig. 5A). This analysis suggests that Wnt signaling promotes both mesodermal progenitors...
and presomitic mesoderm types. We did not observe Sox2+T− co-expressing cells in aggregates, indicating that Wnt signaling, or at least prolonged Wnt signaling, maintained TCreERT2-labeled progenitors in a mesodermal state (Fig. 5A).

To distinguish whether Wnt-induced mesodermal progenitor maintenance occurred specifically in NMPs or in mesodermal progenitors, we took advantage of the Cited1-CreERT2 transgene (Fig. 2F). Cited1-CreERT2;βcatGOF embryos (TAM administered at E6.5) elongated similarly to controls but had abnormal trunk morphology at E10.5 (n=11 control, n=9 mutants). Sections of Cited1-CreERT2;βcatGOF embryos (n=3) showed that the constitutive activation of β-catenin in mesoderm led to the formation of irregularly shaped aggregates of PMPs (Tbx6+T−) and presomitic mesoderm (Tbx6+T−), but not Sox2+T− NMPs (Fig. 5B). This phenotype is very similar to that observed when β-catenin is expressed in T+ progenitors (via TCreERT2) (Fig. 5A), suggesting that the maintenance of mesodermal progenitor fate by Wnt signaling occurs in already-specified mesodermal progenitors.

Implicit in the βcatGOF studies above is that Wnt functions to maintain progenitors. To test this hypothesis, we traced the progeny of permanently labeled TCreERT2;βcatGOF cells (TAM administered at E6.5) using R26R-lacZ to determine whether they differentiated into neurons or PM derivatives such as myocytes (Fig. 6). This analysis showed that traced βcatGOF cells were disproportionately associated with Sox2+ neuronal progenitors in the ventricular zone in the medial neural tube, whereas control traced cells were widely distributed throughout the neural tube (n=4 control, n=4 mutants) (Fig. 6A). Quantification showed that traced βcatGOF cells were twice as likely as control cells to be associated with Sox2+ progenitor populations (Fig. 6B). During development, Sox2-expressing neural progenitors do not express neural differentiation markers (Graham et al., 2003). Analysis of the expression of Sox2 and the neuronal differentiation marker neurofilament indeed showed that these βcatGOF-expressing Sox2+ progenitors were undifferentiated (Fig. 6C). Similarly, we used skeletal muscle myosin (MF20), a terminal myocyte marker, to determine whether TCreERT2;βcatGOF cells become myocytes (Fig. 6D). This was assessed in E9.5 embryos (injected with TAM at E7.5) by examining whether βGal+/βcatGOF cells entered the MF20 lineage. In control embryos, traced TCreERT2 cells were frequently observed entering a MF20+ myocyte fate; however, βcatGOF cells were much less likely to do so (Fig. 6E). These data suggest that persistent Wnt signaling in TCreERT2-traced progenitors inhibits their differentiation.

Loss of NMPs in Wnt3a−/− embryos
To determine the fate of NMPs in Wnt3a−/− mutants, we examined E8.5 embryos for the presence of Sox2- and T-expressing populations. In Wnt3a−/− embryos (n=3), Sox2 and T colocalization was not detected in the anterior PS: Sox2 was readily detected; however, T was not (Fig. 7A). For embryo-wide perspective of Sox2 and T co-expression, we employed whole-mount in situ hybridization detection. Two-color whole-mount in situ hybridization showed that Sox2 and T transcripts were co-expressed in control anterior PS regions at E8.5 (n=8). In Wnt3a−/− embryos, the Sox2-expressing domain was expanded posteriorly while the T expressing domain was dramatically reduced to the posterior most tip of the PS (n=3) (Fig. 7B). This loss of Sox2 and T co-expressing zones is consistent with in vitro studies where Sox2+ progenitors, but not Sox2+T− progenitors, form in the absence of Wnt signaling in differentiating epiblast progenitors (Tsakiridis et al., 2014). It is, however, unknown...
what happens to the NMP population in the absence of Wnt signaling.

To test whether NMPs were being lost by cell death, we examined the PS of control (n=3) and Wnt3a Δ/Δ (n=3) embryos for apoptotic cells using an antibody to cleaved caspase 3. It has been previously shown that apoptosis occurs in the Wnt3a hypomorphic mutant, vestigial tail (vt), within the caudal axis (Shum et al., 1999). In Wnt3a Δ/Δ mutant embryos, cleaved caspase 3 was readily detected in cells of the epithelial PS, with some dying cells in the ingressed PS (Fig. 6C). Dying cells were observed along the lateral length of the mutant PS at E8.5 by whole-mount detection with LysoTracker Red (n=7) (supplementary material Fig. S6) (Fogel et al., 2012). To further explore whether NMPs were being lost in Wnt3a Δ/Δ mutants, we attempted to label this PS population with Tam-CreERT2, TcreERT2 transgenic lines, in concert with the R26R-lacZ reporter, allowed for a germ layer-specific characterization of cell lineages arising in the PS. The TcCreERT2 transgene permitted the permanent, temporally restricted tracing of T-expressing cell lineages across all three germ layers, including the neuroepithelium and the epithelial layer of the PS. TCreERT2 and Cited1-CreERT2 traced similar mesodermal and endodermal populations; however, Cited1-CreERT2 did not label the epithelial streak or neural tube. Thus, TCreERT2 uniquely labeled the neuroepithelium and PS epithelium. By subtracting the Cited1-CreERT2 labeled cells from the TCreERT2 population in wild-type and Wnt3a Δ/Δ embryos, we were able to deduce that cells capable of giving rise to both neural and paraxial mesoderm primarily reside in the epithelial PS. This result is consistent with TCreERT2 labeling a population of epithelial PS cells that include NMPs. As some Cited1-CreERT2-labeled Wnt3a Δ/Δ cells also contribute to the ectopic neural tube it suggests that some already-ingressed ‘mesodermal’ PS cells retain NMP potential, consistent with Sox2/T co-expression in cells of the ingressed PS. Additionally, we used Shh-Cre-based tracings to exclude notochord and endoderm as sources of cells with NMP potential. Thus, lineage tracing and Sox2/T marker analysis arrive at similar conclusions regarding the location of NMPs in the epithelial PS. It remains
formally possible that self-renewing progenitors with neural-restricted potential are also labeled in the TCreERT2 lineage analysis; however, we are unable to address this possibility with the currently available Cre drivers.

Previous studies have shown that axial progenitors display unique behavior in that they can be serially transplanted back into hosts of a younger developmental age and continue to contribute to the developing axis (Cambray and Wilson, 2002; Tam and Tan, 1992). Detailed retrospective clonal lineage analyses provide further evidence for a persistent pool of bipotent self-renewing NMPs (Wilson et al., 2009). Our data demonstrating that a single dose of TAM in TCreERT2 mice labeled NMPs that remained in the epithelial PS or tailbud for at least 3 days of development suggest that these posterior progenitors have self-renewing characteristics and are consistent with the concept that NMPs could be a stem cell population; however, a convincing demonstration that NMPs are consistent with the concept that NMPs could be a stem cell population; however, a convincing demonstration that NMPs could be a stem cell population.

Tracing the fate of PS cells in the Wnt3a−/− mutant background was essential to localize NMPs functionally, and to validate Sox2 and T co-expression as markers of NMPs within the PS. Based on Sox2/T co-expression at E8.5, the majority of the anterior epithelial PS cells are NMPs at this stage; however, we have not proven that all Sox2+T+ cells have the capacity to self-renew. Previous labeling experiments near this region show that it gives rise mainly to PM, while more posterior PS regions give rise to lateral and extra-embryonic mesoderm (Tam and Beddington, 1987). As PM is derived from the anterior PS, and nearly all anterior PS cells express Sox2 and T, it follows that the majority of trunk PM is derived from NMPs. Furthermore, a substantial proportion of the neural cells of the posterior trunk are derived from the anterior PS, and nearly all anterior PS cells express Sox2 and T, co-expression as markers of NMPs within the PS. Based on Sox2/T co-expression at E8.5, the majority of the anterior epithelial PS cells are NMPs at this stage; however, we have not proven that all Sox2+T+ cells have the capacity to self-renew. Previous labeling experiments near this region show that it gives rise mainly to PM, while more posterior PS regions give rise to lateral and extra-embryonic mesoderm (Tam and Beddington, 1987).

MATERIALS AND METHODS

Animals

Wnt3a−, TCreERT2, Cited1-CreERT2, Shh-Cre (Shh-GFPcre), R26R-lacZ, Ctnnb1tm2Kem (β-catenin LOF allele), Ctnnb1lox(ex3)β-catenin gain-of-function allele) and BATlacZ alleles have been described previously (Anderson et al., 2013; Boyle et al., 2008; Braut et al., 2001; Harada et al., 1999; Harfe et al., 2004; Soriano, 1999; Takada et al., 1994).
TCreERT2 or CreI-CreERT2-expressing progenitors in control and Wnt3a−/− mutants was initiated by injection of pregnant mothers with TAM at E7.5 as described previously (Anderson et al., 2013). Any reduction of TAM dose resulted in a reduction in labeled cells, indicating that TAM was not in excess. βcatLOF;TCreERT2 and βcatGOF;TCreERT2 cell lineage studies were initiated with TAM administered at E6.5. For determination of Wnt activation, the βcatGOF line was bred with BATlacZ (Nakaya et al., 2005). This study was carried out in compliance with the Guide for the Care and Use of Laboratory Animals of the National Institutes of Health (Frederick National Lab Animal Care and Use Committee Proposal #12-408). Rodents were euthanized by CO2 inhalation in accordance with the most recent AVMA Guidelines on Euthanasia.

Histology and immunohistochemistry
Antibody detection in frozen section was performed as previously described (Dunty et al., 2014) using primary antibodies to Sox2 (R&D Systems, mab2018, 1:100; Millipore, AB5603, 1:100-1:1000), T (R&D Systems, AF2085, 1:100), fibronectin (DakoCytomation, A0245, 1:500); MF20 (DSHB,1:20); neurofilament protein (DSHB 2H3-c,1:100); Tbx6 (R&D Systems, AF4744, 1:100), Pax7 (DSHB, 1:20), Sox9 (Millipore, Ab5535, 1:100), β-galactosidase (MP Biomedicals, 559761, 1:5000) and β-catenin (BD Bioscience, 610154, 1:400). For paraffin sectioning, embryos were dehydrated in ethanol, cleared in citrisolv (Fisher) for 2×20 min, embedded in 3×20 min changes of molten paraffin (Fisher) and then sectioned at 5 µm. Detection of p-Histone H3 (Cell Signaling, #97065, 1:200) and cleaved caspase 3 (Millipore, AB3623, 1:200) was performed on dewaxed/rehydrated slides boiled in 10 mM sodium citrate (pH 6) for 20 min followed by immunohistochemistry procedures used on frozen section. Aicin Blue 8GX (Sigma) was used to make a 1% staining solution in 3% acetic acid (final pH 2.5). Rehydrated paraffin sections were stained for 1 h, rinsed in tap water and then PBS, counterstained in nuclear Fast Red solution (0.1% NFR, 5% aluminum sulfate), and rinsed in PBS before ethanol dehydration and mounting with permount (Fisher).

In situ hybridization and βGal staining
Whole-mount in situ hybridization was performed using a modification (Biris et al., 2007) of previously described methods (Harland, 1991) with antisense RNA probes using digoxigenin- or fluorescein-labeled UTP (Roche) for synthesis of Sox2, T and Uncx4.1 RNA probes (Invitrogen). For color development, NBT/BCIP (dark purple, Roche), BCIP alone (light blue) or magenta phosphate (magenta, Goldbio) were used as substrates. Whole-mount βGal staining was performed as described previously (Ahn et al., 2013; Whiting et al., 1991). For lineage-tracing studies, labeled cells were identified by direct detection of βGal with specific antibodies (Fig. 2A) or through the enzymatic activity of βGal (Fig. 2B, supplementary material Fig. S2). LysoTracker Red was used as previously described (Fogel et al., 2012).

Imaging, quantification and analysis
Acquired digital images were prepared in Photoshop with adjustments to color, magnification, brightness and contrast applied equally to images under comparison. Cell counts for proliferation rates and CreI-CreERT2; Wnt3a−/− lineage studies represented the total cell count from tissues analyzed averaged over five consecutive sections per embryo. Sibling
embryos of similar size were compared in analysis containing Wnt3a−/− and TCreERT2;3catLOF mutants, as they lack a full complement of somites typically used to accurately stage match embryos. Variation in overall embryo proliferation rates was observed among sibling embryos regardless of genotype. To better compare embryos, we normalized the neural proliferation rate by dividing this rate with the proliferation rate of non-PM (intermediate and lateral plate mesoderm), as this tissue forms normally in Wnt3a−/− mutants. The resulting calculation produces a ratio used for comparisons of proliferation rate. For comparisons of lineage and cell fate, the quantifications are derived from sampled representative regions of 30 cells or more using DAPI-stained nuclei as a counting standard. Comparisons of lineage-traced cell fate were presented as a percent from each sampled embryo, and multiple samples were used to calculate average, standard deviations and t-test (two-tailed) statistical comparisons.

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Competing interests

The authors declare no competing or financial interests.

Author contributions

R.J.G. and T.P.Y. wrote the manuscript. R.J.G. created the genetic strains that were characterized by R.J.G., R.B.C., M.W.K. and L.C.C. with T.P.Y. guiding the study. L.C.C. drew the cartoons. M.L. shared mice strains, provided expertise and edited the manuscript.

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Supplementary material

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References


Brachyury is essential for posterior mesoderm formation. Development 129, 5425-5436.


