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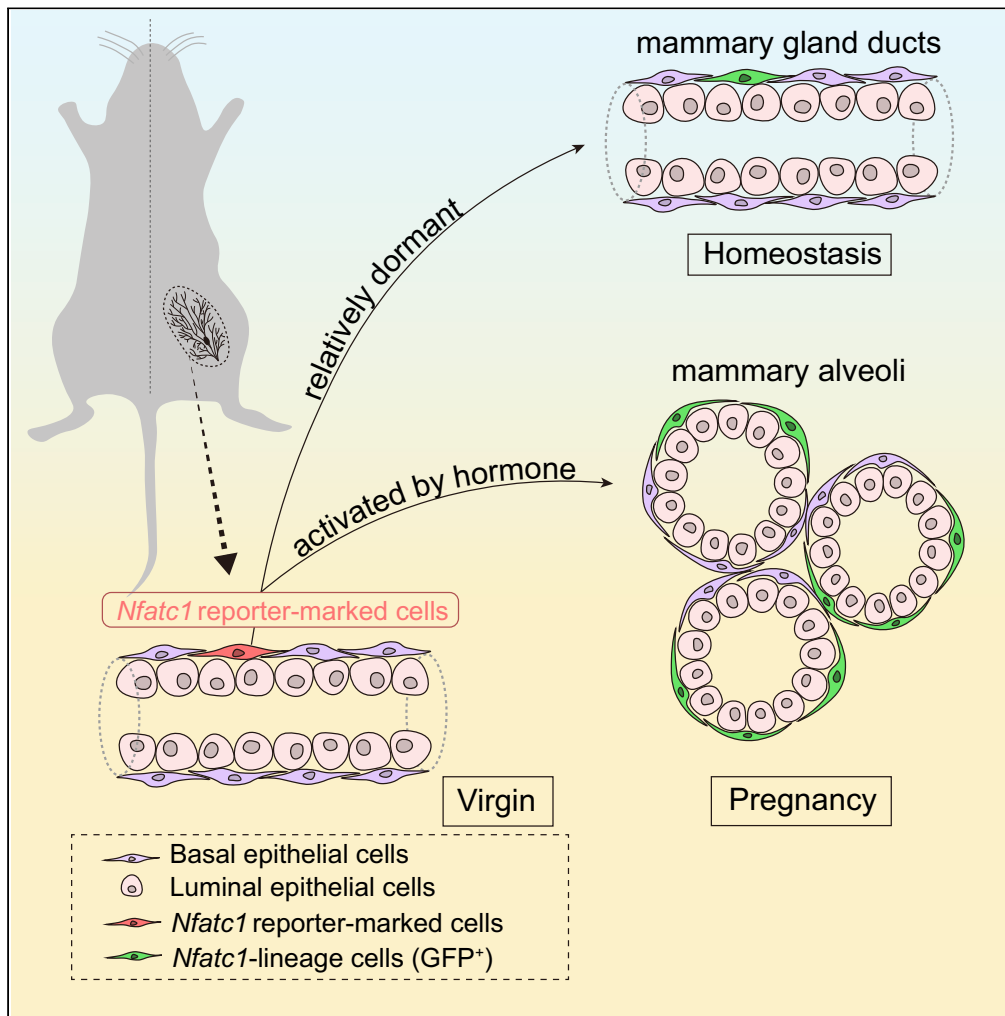
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Highlights

Nfatc1 marked a subpopulation of basal dormant stem/progenitor cells

Nfatc1-marked cells have limited contribution to the homeostasis of mammary gland

Nfatc1 reporter-marked cells divide rapidly during pregnancy

Nfatc1 reporter-marked cells are preferentially used for multiple pregnancies

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Article

Dormant *Nfatc1* reporter-marked basal stem/progenitor cells contribute to mammary lobuloalveoli formation

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SUMMARY

The Mammary gland undergoes complicated epithelial remodeling to form lobuloalveoli during pregnancy, in which basal epithelial cells remarkably increase to form a basket-like architecture. However, it remains largely unknown how dormant mammary basal stem/progenitor cells involve in lobuloalveolar development. Here, we show that *Nfatc1* expression marks a rare population of mammary epithelial cells with the majority being basal epithelial cells. *Nfatc1* reporter-marked basal epithelial cells are relatively dormant mammary stem/progenitor cells. Although *Nfatc1* reporter-marked basal epithelial cells have limited contribution to the homeostasis of mammary epithelium, they divide rapidly during pregnancy and contribute to lobuloalveolar development. Furthermore, *Nfatc1* reporter-marked basal epithelial cells are preferentially used for multiple pregnancies. Using single-cell RNA-seq analysis, we identify multiple functionally distinct clusters within the *Nfatc1* reporter-marked cell-derived progeny cells during pregnancy. Taken together, our findings underscore *Nfatc1* reporter-marked basal cells as dormant stem/progenitor cells that contribute to mammary lobuloalveolar development during pregnancy.

INTRODUCTION

The mammary gland is unique in that the majority of its epithelial morphogenesis occurs postnatally (Richert et al., 2000). It undergoes complicated epithelial remodeling at multiple distinctive stages from embryonic and pubertal development to reproductive life (Macias and Hinck, 2012). After birth, the gland just rests with a rudimentary structure until puberty; with the onset of puberty, the increase of estrogen and growth hormone levels promote the rapid proliferation of mammary epithelial cells leading to the formation of a mammary ductal tree that fills the fat pad (Inman et al., 2015; Macias and Hinck, 2012). During pregnancy, massive tissue changes occur within the mammary gland in response to progesterone and prolactin, resulting in the formation of milk-secreting lobuloalveoli in preparation for lactation. In this process, both basal and luminal epithelial cells divide rapidly and globally within the ductal branches and developing alveoli, and the luminal alveolar cells form a sphere-like single-layer structure surrounded by a basket-like architecture of basal contractile myoepithelial cells (Hens and Wysolmerski, 2005). However, although it has been known that the numbers of basal myoepithelial cells increase 11-fold in the basal compartment during pregnancy (Asselin-Labat et al., 2010), the cellular mechanism underlying basal mammary stem/progenitor cells' role in driving contractile basket-like architecture remains to be fully understood.

Increasing evidence indicates that mammary stem cells (MaSCs) are diverse (Fu et al., 2020). MaSCs have been shown to reside in the basal layer and can reconstitute the entire mammary gland when transplanted into a cleared fat pad (Shackleton et al., 2006; Stingl et al., 2006). Subsequently, lineage tracing assays with *K5*, *Lgr5*, and *Axin2* promoter drivers have demonstrated the existence of long-lived basal-restricted unipotent progenitor cells that drive postnatal morphogenesis and replenishment of the basal layer during homeostasis in the adult stage (van Amerongen et al., 2012; Van Keymeulen et al., 2011; Wuidart et al., 2016). Bipotent basal cells have also been identified in the adult mammary gland in lineage tracing assays using the same or more restricted gene promoter drivers (*K5*, *Lgr5*, *Procr*, and *Dll1*) (Chakrabarti et al., 2018; Rios et al., 2014; Wang et al., 2015). In addition, *Lgr5*⁺*Tspan8*^{high} cells and *Bcl11b*^{high} cells have been

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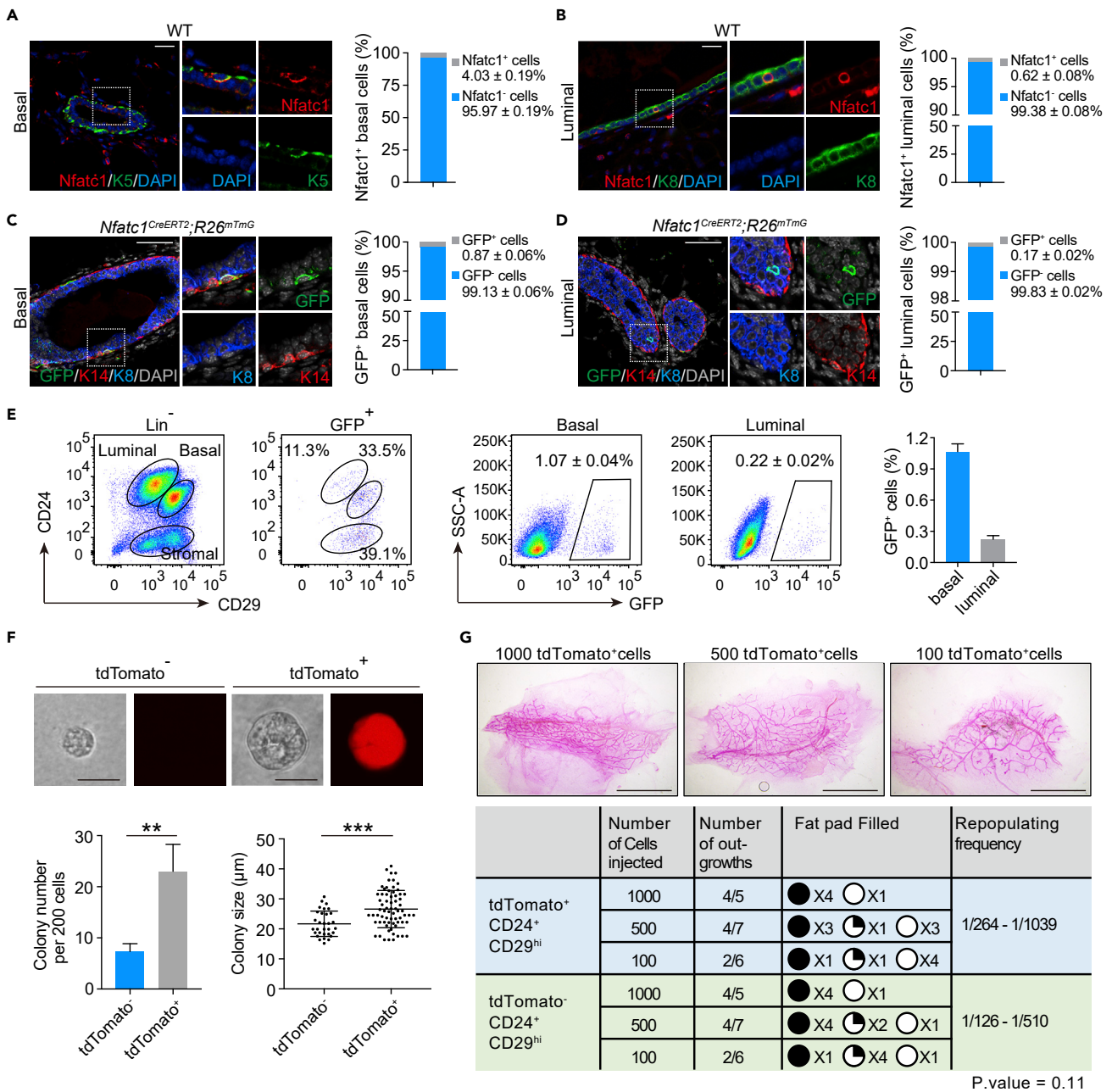


Figure 1. *Nfatc1* expression labels a small population of mammary stem/progenitor cells (see also Figure S1)

(A and B) Co-immunofluorescence assay for *Nfatc1* and K5 (A) or *Nfatc1* and K8 (B) in eight-week-old WT mammary glands. $n = 3$ mice. Scale bar, 25 μm . High-magnification images are shown on the right. The percentages of *Nfatc1*⁺ basal cells among total basal cells (A) and *Nfatc1*⁺ luminal cells among total luminal cells (B) were quantified. A total of 1,952 basal cells and 8,335 luminal cells from three mice were counted

(C and D) Co-immunofluorescence assay for GFP (green), K14 (red), and K8 (blue) in mammary glands from *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice at the age of eight weeks. GFP⁺ cells were present in the basal (C) and luminal (D) layers. $n = 3$ mice. Scale bar, 50 μm . Quantification analysis shows the percentages of GFP⁺ basal cells among total basal cells (C) and GFP⁺ luminal cells among total luminal cells (D). A total of 1,614 basal cells and 5,314 luminal cells from three mice were quantified

(E) Flow cytometry assay of GFP⁺ cells from *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice at the age of eight weeks. The mice were administered TAM three times, and the samples were collected 48 h after the last induction. Quantification analysis showing the percentages of GFP⁺ basal cells among total basal cells and GFP⁺ luminal cells among total luminal cells. Another analysis approach is also shown. Total GFP⁺ cells were applied to CD24 and CD29 gates, showing the distribution of GFP⁺ cells in the basal and luminal layers. $n = 3$ mice

(F) Colony formation efficiency and colony size in the Matrigel culture. For tdTomato⁺ and tdTomato⁻ basal cells, a total of 69 and 31 clones were analyzed from three independent experiments. Scale bar, 25 μm

Figure 1. Continued

(G) Transplantation of tdTomato⁺ and tdTomato⁻ basal epithelial cells by limiting dilution. Data are pooled from two independent experiments. Scale bar, 5 mm. The repopulation frequency was calculated using the method in <http://bioinf.wehi.edu.au/software/elda/>. Data represent the mean value \pm SD ***p* < 0.01, ****p* < 0.001

identified as quiescent MaSCs in the basal compartment, with *Lgr5*⁺*Tspan8*^{high} cells residing predominantly in the proximal region (Chakrabarti et al., 2018) and *Bcl11b*^{high} cells localizing throughout the mammary gland (Cai et al., 2017). It has been implicated that a striking expansion of basal epithelial cells during pregnancy is fueled by distinct MaSC subsets (Chakrabarti et al., 2018; Rios et al., 2014; Van Keymeulen et al., 2011; Wang et al., 2015). However, our understanding of the contribution of mammary stem/progenitor cells to lobuloalveolar development remains limited.

Nfatc1 is a transcription factor the expression of which labels the quiescent bulge stem cells of the hair follicle, and it maintains the stem cells in a quiescent state as a downstream factor of BMP signaling (Horsley et al., 2008; Keyes et al., 2013). Interestingly, *Nfatc1* regulates prolactin receptor, *Prlr*, in hair follicles during pregnancy (Goldstein et al., 2014). As prolactin/*Prlr* signaling is a key driver of mammary lobuloalveolar development and milk protein production (Sternlicht, 2006), we entertained the possibility that *Nfatc1* may also mark mammary epithelial cell subsets that are important for lobuloalveolar development during pregnancy. In this study, we demonstrated that *Nfatc1* marked a rare population of mammary epithelial cells as dormant long-lived stem/progenitor cells. *Nfatc1* reporter-marked stem/progenitor cells show limited contribution to the homeostasis of mammary epithelial cells, whereas they contribute to mammary lobuloalveolar development during pregnancy and are preferentially used for multiple pregnancies.

RESULTS***Nfatc1* expression labels a small population of mammary stem/progenitor cells**

To identify and characterize *Nfatc1*⁺ epithelial cells in the mammary gland, we first performed immunofluorescence assay to detect the *Nfatc1* expression pattern in the mammary gland. We found that a small population of *Nfatc1*⁺ epithelial cells was sporadically distributed in both basal and luminal layers of mammary epithelium (Figures 1A and 1B). Quantification analysis showed that $4.03 \pm 0.19\%$ of the basal cells were *Nfatc1*⁺ (Figure 1A), and $0.62 \pm 0.08\%$ of the luminal cells were *Nfatc1*⁺ (Figure 1B).

In order to further characterize *Nfatc1*⁺ mammary epithelial cells, we generated a tamoxifen-inducible Cre (*Cre*^{ERT2}) knock-in allele targeted immediately downstream of the eighth exon of the endogenous *Nfatc1* locus (Figure S1A), thereby targeting all isoforms. The targeted allele was validated by PCR (Figure S1B). After crossing, *Nfatc1*^{CreERT2}; *R26*^{mTmG} or *Nfatc1*^{CreERT2}; *R26*^{tdTomato} reporter mice were obtained for labeling *Nfatc1*⁺ cells and their progenies (Figure S1C). To validate these *Nfatc1* reporter mice, we administered one single pulse of tamoxifen (TAM) into *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice and analyzed the mice 48 h after TAM induction for hair follicle labeling. As expected, we found *Nfatc1* reporter-marked cells in the bulge regions of the hair follicles at both telogen and anagen stages that also stained positive for *Nfatc1* protein (Figure S1D), indicating the validity of the *Nfatc1* reporter mice as a useful lineage tracing tool.

To detect the distribution of reporter-marked cells in the mammary gland, eight-week-old *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice were induced with three pulses of TAM, which enhances the labeling efficiency relative to one pulse of TAM, and the mice were analyzed 48 h after the last induction (Figure S1E). In agreement with *Nfatc1* protein expression, *Nfatc1* reporter labeled rare mammary epithelial cells in both the basal and luminal layers, with $0.87 \pm 0.06\%$ of basal cells and $0.17 \pm 0.02\%$ of luminal cells being GFP⁺ (Figures 1C and 1D). We also quantified the proportions of *Nfatc1* reporter-marked basal and luminal cells by flow cytometry, which revealed $1.07 \pm 0.04\%$ and $0.22 \pm 0.02\%$ GFP⁺ cells in basal and luminal cells, respectively (Figure 1E). These values are lower than those of *Nfatc1*⁺ cells assayed by immunofluorescence, which is most likely due to incomplete labeling efficiency. These results were further corroborated using the *Nfatc1*^{CreERT2}; *R26*^{tdTomato} mice (Figures S1F and S1G). Taken together, our findings show that *Nfatc1* labels a rare population of mammary epithelial cells with the majority of them being in the basal compartment.

Next, we sought to examine whether *Nfatc1* reporter-marked basal cells are mammary stem/progenitor cells. We sorted *Nfatc1* reporter-marked basal epithelial cells (tdTomato⁺/CD24⁺/CD29^{hi}) using *Nfatc1*^{CreERT2}; *R26*^{tdTomato} mice and cultured the cells in Matrigel. *Nfatc1* reporter-marked (tdTomato⁺)

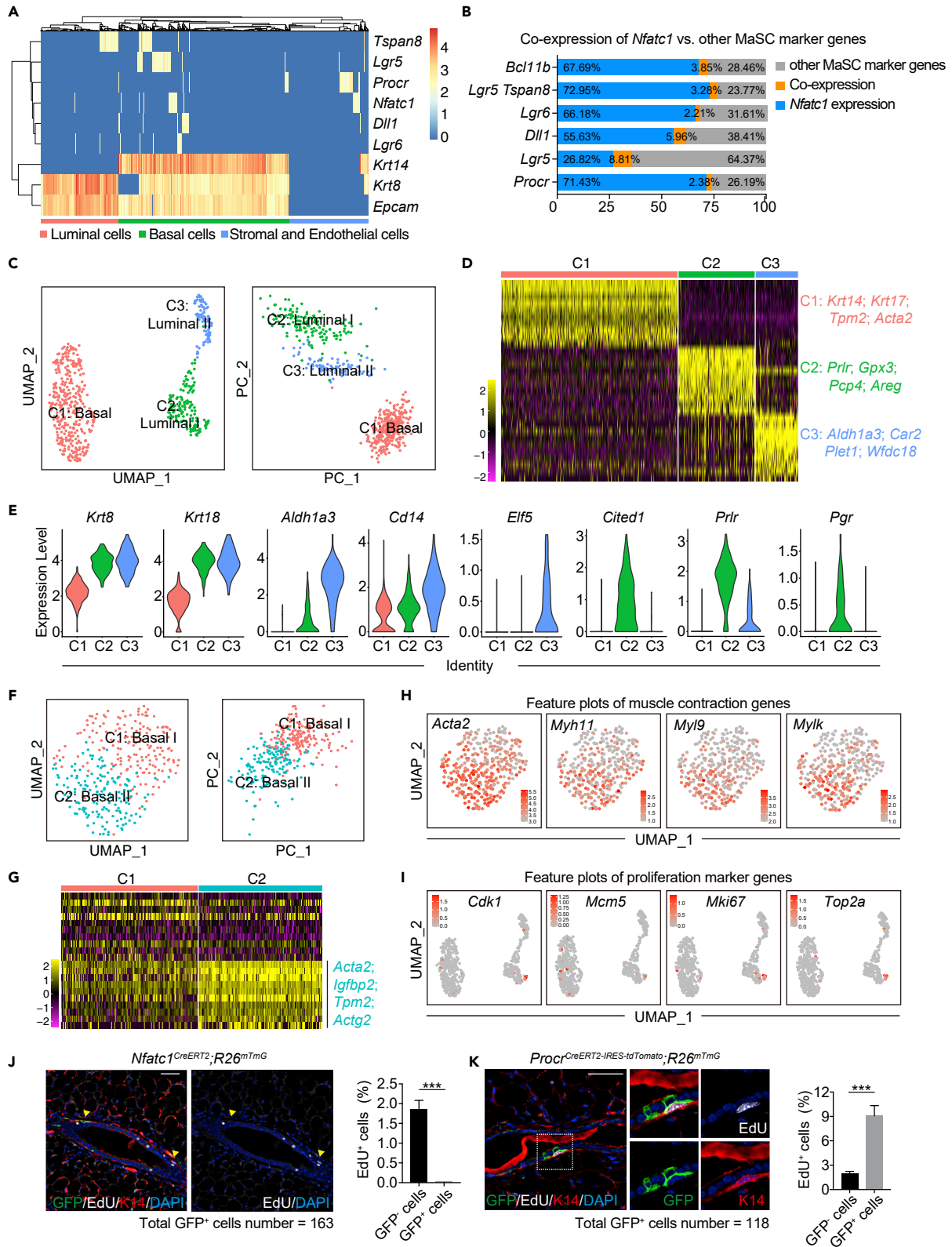


Figure 2. *Nfatc1* reporter-marked stem/progenitor cells are dormant and distinct from other known MaSCs (see also Figure S2)

(A) Heatmap analysis of the scRNA-seq database shows the expression patterns of *Nfatc1* and other MaSC marker genes (*Tspan8*, *Lgr5*, *Procr*, *Dll1*, and *Lgr6*), as well as *Krt14* (*K14*), *Krt8* (*K8*), and *Epcam*. The scRNA-seq database for mammary epithelial cells was adapted from a previously published study in Nature and downloaded from <https://tabula-muris.ds.czbiohub.org>. A nUMI count >150 was used as a cutoff for low-expression cells

(B) Quantification for the percentages of basal cells expressing *Nfatc1* or the MaSC marker genes *Bcl11b*, *Tspan8/Lgr5*, *Lgr6*, *Dll1*, *Lgr5*, and *Procr* in (A). Blue indicates *Nfatc1*⁺ cells; grey indicates other MaSC marker genes; and orange indicates double-positive cells

(C) UMAP and PCA plots reveal cellular heterogeneity of 558 tdTomato⁺ mammary epithelial cells that were sorted from *Nfatc1*^{CreERT2}; *R26*^{tdTomato} mice

(D) Heatmap showing differentially expressed genes in each cluster for (C). Selected signature genes are shown on the right

(E) Violin plots show marker genes of common luminal cells (*Krt8/K8* and *Krt18/K18*), milk lineage cells (*Cd14*, *Aldh1a3*, and *Elf5*), and ER⁺ lineage cells (*Cited1*, *Prlr*, and *Pgr*) for each cluster in (C)

(F) UMAP and PCA plots showing the reclustering results for 333 tdTomato⁺ basal cells. Luminal epithelial cells were removed

(G) Heatmap of differentially expressed genes in each cluster for (F). Selected signature genes are shown on the right

(H) Feature plots showing expression distribution of genes functioning in muscle contraction

(I) Feature plots showing expression distribution of genes functioning in cell proliferation in each cluster for (C)

(J and K) Co-immunofluorescence for GFP, EdU, and K14 in *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice (J) and *Procr*^{CreERT2-IRES-tdTomato}; *R26*^{mTmG} mice (K) at 48 h post-TAM induction. Scale bar, 50 μm. The samples were collected 3 h after one dose of EdU injection. Quantification analysis shows the percentages of GFP⁺EdU⁺ cells versus GFP⁺ cells and GFP⁻EdU⁺ cells versus GFP⁻ cells in *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice (J) and *Procr*^{CreERT2-IRES-tdTomato}; *R26*^{mTmG} mice (K). For *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice, 163 GFP⁺ cells and 1766 GFP⁻ cells were quantified, respectively. For *Procr*^{CreERT2-IRES-tdTomato}; *R26*^{mTmG} mice, 118 GFP⁺ cells and 1949 GFP⁻ cells were quantified, respectively. *n* = 3 mice for each mouse model. Data represent the mean value ±SD ****p* < 0.001

basal epithelial cells gave rise to 3-fold more colonies than their tdTomato⁻ (tdTomato⁻/CD24⁺/CD29^{hi}) counterparts (Figure 1F), and the average size of the colonies derived from tdTomato⁺ basal epithelial cells was significantly larger than that from tdTomato⁻ basal epithelial cells (Figure 1F). These data indicate that tdTomato⁺ basal epithelial cells possess higher colony-forming and proliferative ability than tdTomato⁻ basal epithelial cells. Furthermore, We sorted *Nfatc1* reporter-marked basal epithelial cells (tdTomato⁺/CD24⁺/CD29^{hi}) and *Nfatc1* reporter-negative basal epithelial cells (tdTomato⁻/CD24⁺/CD29^{hi}) using *Nfatc1*^{CreERT2}; *R26*^{tdTomato} mice and transplanted them into cleared fat pads. We found that the tdTomato⁺ basal epithelial cells were able to reconstitute the whole mammary gland, although the repopulating frequency of tdTomato⁺ basal epithelial cells is generally identical to tdTomato⁻ basal epithelial cells (Figure 1G). It suggests that tdTomato⁺ basal epithelial cells contain mammary stem cells with high colony-forming capacity and that their activation might be context-dependent.

***Nfatc1* reporter-marked stem/progenitor cells are dormant and distinct from other known MaSCs**

Next, we wanted to distinguish between tdTomato⁺ basal epithelial cells and other mammary stem/progenitor cells identified by known marker genes including *Procr*, *Lgr5*, *Lgr6*, and *Dll1* (Blaas et al., 2016; Chakrabarti et al., 2018; Fu et al., 2017; Wang et al., 2015). We thus compared the *Nfatc1* expression pattern to those known mammary stem/progenitor marker genes in mammary epithelial cells at a single-cell level. We analyzed an scRNA-seq database (<https://tabula-muris.ds.czbiohub.org>) for mammary epithelial cells from adult female mice, which is adapted from a previously published literature in Nature (Tabula Muris Consortium et al., 2018). With a cutoff of nUMI count >150, quantification of the cells positive for each marker gene showed that the proportions of *Nfatc1*⁺, *Procr*⁺, *Lgr5*⁺, *Lgr6*⁺ and *Dll1*⁺ basal epithelial cells out of total basal epithelial cells were 7.09%, 2.75%, 14.57%, 3.51% and 5.11%, respectively (Figure S2A). These values were generally consistent with the proportions for each marker gene (*Procr*⁺, 3%; *Lgr5*⁺, 11.4%; *Lgr6*⁺, 4.7%; *Dll1*⁺, 12%) that were analyzed by flow cytometry in previously published studies (Blaas et al., 2016; Chakrabarti et al., 2018; Fu et al., 2017; Wang et al., 2015). Thus, we utilized this cutoff to perform heatmap analysis, and found that *Nfatc1*⁺ basal epithelial cells rarely overlap with *Lgr6*⁺, *Tspan8*⁺, *Procr*⁺, and *Dll1*⁺ basal epithelial cells, and the overlap between *Nfatc1*⁺ and *Lgr5*⁺ basal epithelial cells is 8.81% out of *Nfatc1*⁺ and *Lgr5*⁺ cells (Figures 2A and 2B). The conclusion of low overlap was further supported by feature t-SNE assays (Figure S2B). Therefore, the *Nfatc1* reporter-marked mammary stem/progenitor cells are generally distinct from other known mammary stem/progenitor cells.

To further characterize the identity of *Nfatc1* reporter-marked epithelial cells, we performed unbiased single-cell transcriptomic analysis on tdTomato⁺ mammary epithelial cells that were isolated from eight-week-old female *Nfatc1*^{CreERT2}; *R26*^{tdTomato} mice 48 h after three pulses of TAM induction (Figure S2C). After quality control, 2,952 cells were used for downstream analysis (Figure S2D). Unsupervised clustering identified seven distinct clusters (Figure S2E). Based on known gene signatures, cluster C4 was identified as basal epithelial cells, cluster C6 as luminal epithelial cells, clusters C1 and C3 as endothelial cells, cluster C2 as stromal cells, cluster C5 as muscle cells, and cluster C7 as immune cells (Figures S2E–S2G). After

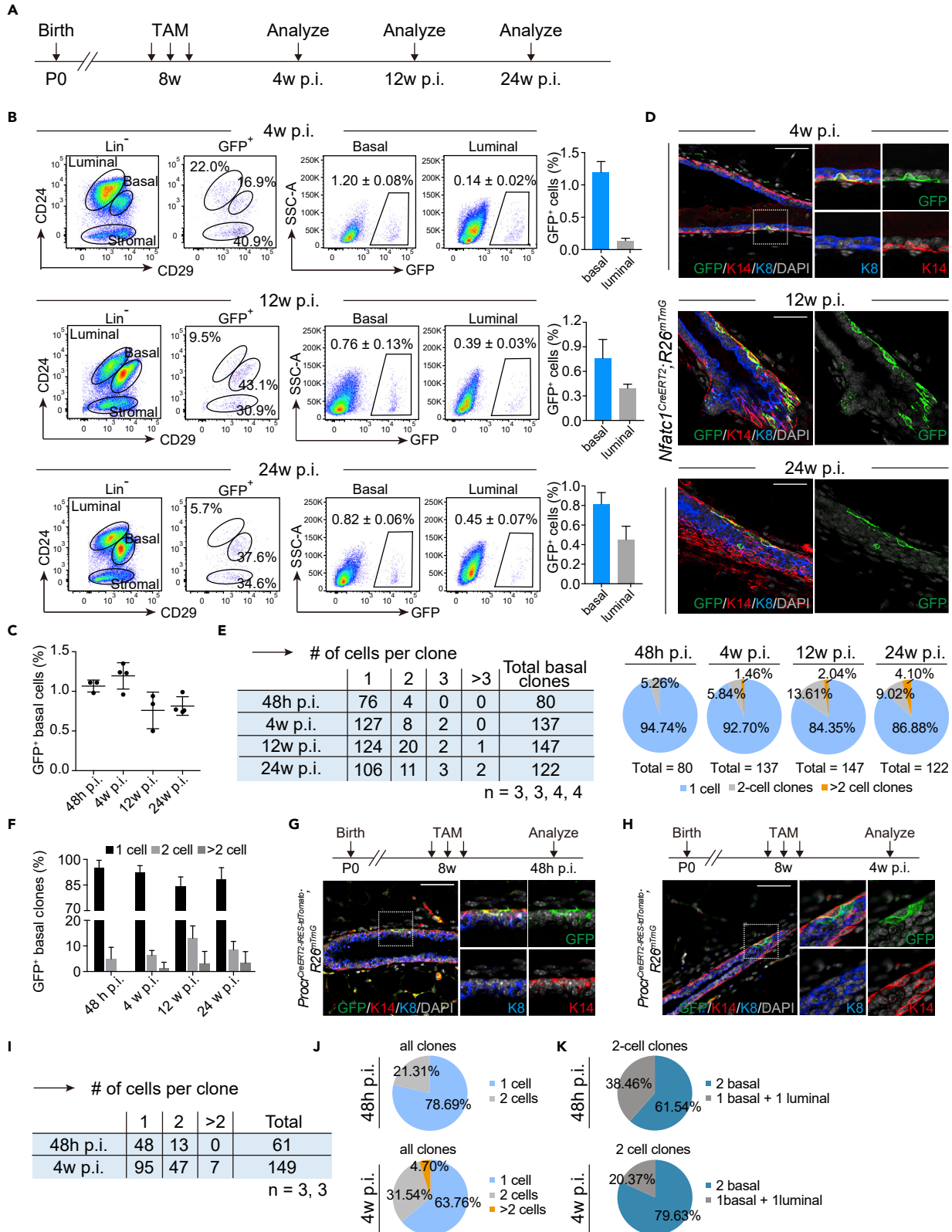


Figure 3. *Nfatc1* reporter-marked basal epithelial cells have restrictive contribution to mammary epithelium during homeostasis (see also Figures S3 and S4)

(A) Experimental strategy for successive lineage tracing experiments at different timepoints
(B) Flow cytometry for GFP⁺ epithelial cells from 8-week-old female *Nfatc1*^{CreERT2};*R26*^{mTmG} mice 4, 12, and 24 weeks post-TAM induction (p.i.). Another approach to analysis was also shown. Total GFP⁺ cells were applied to CD24 and CD29 gates, showing the distribution of GFP⁺ cells in the basal and luminal layers. Quantification analysis showing the percentages of GFP⁺ basal cells per total basal cells and GFP⁺ luminal cells per total luminal cells for each timepoint. *n* = 4, 3, and 4 mice at 4, 12, and 24 weeks post-induction, respectively
(C) Quantification analysis of the percentages of GFP⁺ basal cells per total basal cells in *Nfatc1*^{CreERT2};*R26*^{mTmG} mice 48 h (*n* = 3), 4 weeks (*n* = 4), 12 weeks (*n* = 3), and 24 weeks (*n* = 4) after TAM induction
(D) Co-immunofluorescence assay for GFP (green), K14 (red), and K8 (blue) in *Nfatc1*^{CreERT2};*R26*^{mTmG} mice at 4, 12, and 24 weeks post-TAM induction. Scale bar, 50 μm
(E and F) Quantification analysis shows the numbers (left panel of (E)) and the percentages (right panel of (E)) of single-cell, two-cell, and multicell basal clones in *Nfatc1*^{CreERT2};*R26*^{mTmG} mice at indicated timepoints (E). Another statistical result is also shown in panel (F). The total numbers of basal clones were 80, 137, and 147 and 122 at 48 h, 4, 12, and 24 weeks post-induction, respectively. *n* = 3, 3, 4, and 4, respectively
(G and H) Co-immunofluorescence assay for GFP (green), K14 (red), and K8 (blue) from *Procr*^{CreERT2-IRES-tdTomato};*R26*^{mTmG} mice 48 h (G) and 4 weeks (H) post-induction. Scale bar, 50 μm
(I–K) Quantification analysis shows the numbers (I) and the percentages of single-cell, two-cell, and multicell clones in *Procr*^{CreERT2-IRES-tdTomato};*R26*^{mTmG} mice at the indicated timepoints (J). The percentages of basal-only and bilineage two-cell clones was also quantified, shown on (K). *n* = 3 for each timepoint. A total of 61 and 149 clones were quantified at 48 h and 4 weeks post-induction, respectively. Data represent the mean value ± SD

removing non-epithelial cells, tdTomato⁺ epithelial cells were regrouped into three distinct clusters (Figures 2C and 2D). Basal epithelial cells belonged to a single cluster (C1), whereas luminal epithelial cells were divided into two clusters, C2 and C3. Among them, the cells in C3 were identified as milk lineage cells, as they highly express marker genes of luminal progenitor cells, *Aldh1a3*, *Cd14*, and *Elf5* (Bach et al., 2017; Pervolarakis et al., 2020; Shehata et al., 2012) (Figure 2E). The cells in C2 were identified as ER⁺ lineage cells, as they highly express differentiated cell marker genes, *Prlr*, *Pgr*, and *Cited1* (Bach et al., 2017; Shehata et al., 2012) (Figure 2E). To better define the identity of tdTomato⁺ basal epithelial cells, we removed luminal epithelial cells and then performed unsupervised clustering on these cells. tdTomato⁺ basal epithelial cells can be further divided into two distinct subgroups based on the expression levels of genes functioning on muscle contraction (Figures 2F–2H). We also examined the proliferative status of tdTomato⁺ mammary epithelial cells. Cell cycle genes were barely expressed in tdTomato⁺ basal epithelial cells (Figure 2I), suggesting a dormant status. To confirm this idea, we performed an EdU incorporation assay using *Nfatc1*^{CreERT2};*R26*^{mTmG} mice at 48 h post induction, then quantified EdU⁺GFP⁺ cells versus GFP⁺ cells and EdU⁺GFP⁻ cells versus GFP⁻ cells in mammary gland at the age of eight weeks. It showed that no *Nfatc1* reporter-marked basal cells were found in S phase, whereas 1.85 ± 0.13% of *Nfatc1* reporter-negative cells were in S phase (Figure 2J). In comparison, a parallel experiment showed that approximately 9.06 ± 0.73% of *Procr* reporter-marked cells were in the S phase at the same stage, whereas 1.93 ± 0.18% of *Procr* reporter-negative cells were in the S phase (Figure 2K). These data suggest that *Nfatc1* reporter-marked basal epithelial cells are dormant relative to both *Nfatc1* reporter-negative cells and *Procr*-reporter marked cells.

***Nfatc1* reporter-marked basal epithelial cells have restrictive contribution to mammary epithelium during homeostasis**

To test the contribution of *Nfatc1* reporter-marked basal epithelial cells to the homeostasis of mammary epithelium *in vivo*, we performed lineage tracing assay using female *Nfatc1*^{CreERT2};*R26*^{mTmG} mice at successive timepoints. The mice were administered three doses of TAM at the age of 8 weeks and then were analyzed 4, 12, and 24 weeks after the last induction (Figure 3A). Using flow cytometry, the proportions of basal GFP⁺ cells 48 h, 4, 12 and 24 weeks after TAM induction were found to be 1.07 ± 0.04%, 1.20 ± 0.08%, 0.76 ± 0.13% and 0.82 ± 0.06%, respectively (Figures 1E and 3B). This result implies that basal GFP⁺ epithelial cells existed for a long time, whereas the proportion of GFP⁺ basal epithelial cells was not significantly altered with time after TAM induction (Figures 3B and 3C). Immunofluorescence assay further showed that GFP⁺ basal-only clones exist at distinct timepoints, whereas no GFP⁺ bilineage clones were observed (Figure 3D). Quantification of GFP⁺ clones showed that the proportions of two-cell basal clones 48 h, 4, 12, and 24 weeks after TAM induction were 5.26, 5.84, 13.61, and 9.02%, respectively, and those of multicell basal clones (≥ 3 cells) were 0, 1.46, 2.04, and 4.10%, respectively (Figures 3E and 3F). The data demonstrate that the proportions of two-cell and multicell basal clones increased with time. Taken together with findings above, these data suggest that *Nfatc1*-reporter marked basal epithelial cells, whereas relatively dormant, still contribute to the homeostasis of the mammary epithelium. In parallel experiments, we also administered the same TAM induction in *Procr*^{CreERT2-IRES-tdTomato};*R26*^{mTmG} mice at the age of 8 weeks and analyzed them 48 h and 4 weeks after initial lineage labeling. We found that the

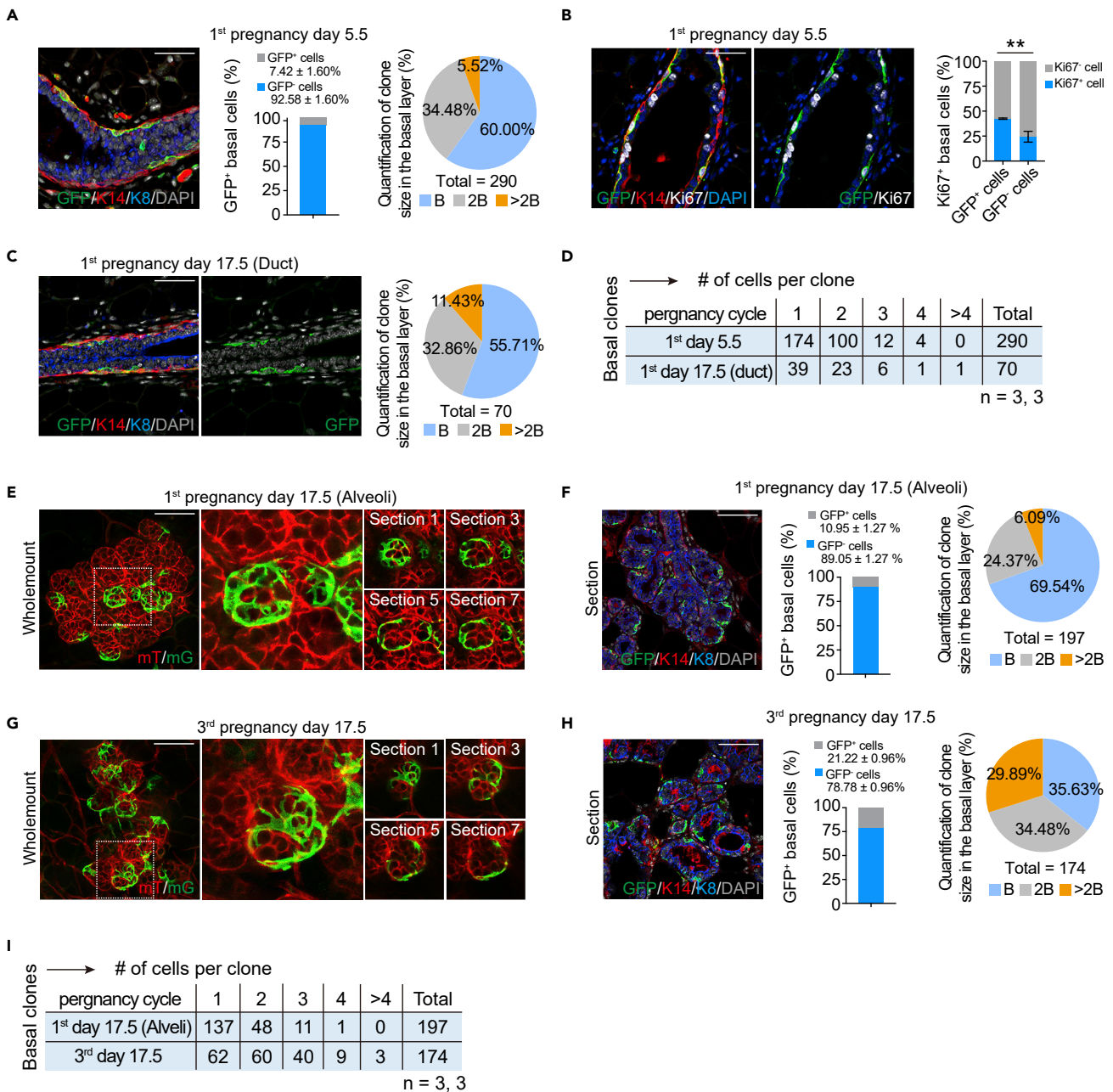


Figure 4. *Nfatc1* reporter-marked basal epithelial cells significantly contribute to lobuloalveolar development during pregnancy (see also Figure S5)

(A) Co-immunofluorescence assay of GFP (green), K14 (red), and K8 (blue) in mammary ducts from female *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice at pregnancy day 5.5. Scale bar, 50 μ m. Quantification analysis showing the percentage of GFP⁺ basal cells among total basal cells. A total of 2369 basal cells from three mice were quantified. Quantification analysis showing the percentages of single-cell, two-cell, and multicell basal clones. n = 3 mice. A total of 290 basal GFP⁺ clones were quantified

(B) Co-immunofluorescence assay of GFP (green), K14 (red), and Ki67 (white) in mammary glands from female *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice at pregnancy day 5.5. Scale bar, 50 μ m. Quantification analysis showing the percentage of GFP⁺Ki67⁺ cells among GFP⁺ cells and the percentage of GFP⁻Ki67⁺ cells among GFP⁻ cells. A total of 175 GFP⁺ cells and 1,444 GFP⁻ cells were quantified from three mice

(C) Co-immunofluorescence assay of GFP (green), K14 (red), and K8 (blue) in mammary ducts from female *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice at pregnancy day 17.5. Scale bar, 50 μ m. Quantification analysis showing the percentages of single-cell, two-cell and multicell basal clones. n = 3 mice for each timepoint. The total number of basal GFP⁺ clones was 70

(D) Quantification analysis showing the numbers of single-cell, two-cell, and multicell basal clones for *Nfatc1*^{CreERT2}; *R26*^{mTmG} mice at indicated timepoints. n = 3 mice for each timepoint

Figure 4. Continued

(E and G) Wholemout confocal image for *Nfatc1*^{CreERT2};*R26*^{mTmG} mice at the first pregnant day 17.5 (E) and third pregnant day 17.5 (G). High-magnification images were shown in the middle. Optical serial sections were shown on the right. Scale bar, 50 μ m

(F and H) Co-immunofluorescence assay for GFP (green), K14 (red), and K8 (blue) in sections from *Nfatc1*^{CreERT2};*R26*^{mTmG} mammary alveoli at the first (F) and third (H) pregnant day 17.5, respectively. Scale bar, 50 μ m. Quantification analysis showing the percentage of GFP⁺ basal cells among total basal cells. A total of 1,949 basal cells in (F) and 988 basal cells in (H) from three mice were quantified. Quantification analysis shows the percentages of single-cell, two-cell and multicell basal clones for *Nfatc1*^{CreERT2};*R26*^{mTmG} mice at the indicated timepoints. $n = 3$ mice for each timepoint. The total number of basal GFP⁺ clones was 197 and 174 at the first (F) and third (H) pregnant day 17.5, respectively

(I) Quantification analysis showing the numbers of single-cell, two-cell and multicell basal clones for *Nfatc1*^{CreERT2};*R26*^{mTmG} mice at the indicated timepoints. $n = 3$ mice for each timepoint. Data represent the mean value \pm SD * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$

proportions of two-cell clones were 21.31 and 31.54% 48 h and 4 weeks after initial lineage labeling and those of multicell clones were 0 and 4.7%, respectively (Figures 3G–3J). In comparison, 38.46 and 20.37% bilineage two-cell clones were detected in the mammary duct from *Procr*^{CreERT2-IRES-tdTomato};*R26*^{mTmG} mice (Figure 3K). These findings show that, compared with *Procr* reporter-marked MaSCs, *Nfatc1* reporter-marked basal epithelial cells are unipotent and have limited contribution to the homeostasis of mammary epithelium.

Regarding the luminal layer, we found that the proportion of luminal GFP⁺ cells slightly decreased four weeks after initial lineage labeling, relative to 48 h after TAM induction, and then markedly increase 12 and 24 weeks after TAM induction (Figure S3A). Immunofluorescence assays revealed GFP⁺ multicell clones in mammary luminal epithelium at distinct timepoints (Figure S3B), and the proportions of two-cell and multicell clones increased with time (Figures S3C and S3D). These results support the idea that *Nfatc1* reporter-marked luminal epithelial cells contain a fraction of luminal progenitor cells, which is consistent with the above scRNA-seq data.

Next, we examined the contribution of *Nfatc1* reporter-marked epithelial cells to mammary gland development at the age of 4 weeks when it is puberty. At this stage, the mammary gland undergoes a period of expansive proliferation that is primarily regulated by estrogen and growth hormones (Macias and Hinck, 2012). Similar to the adult stage (eight-week old), GFP⁺ basal epithelial cells contributed to the turnover of mammary epithelium with low efficiency (Figures S4A–S4E). It further suggests a limited contribution to mammary homeostasis.

***Nfatc1* reporter-marked basal epithelial cells significantly contribute to lobuloalveolar development during pregnancy**

We next sought to explore the behaviors of *Nfatc1* reporter-marked epithelial cells during pregnancy (Figure S5A). *Nfatc1*^{CreERT2};*R26*^{mTmG} mice were administered three doses of tamoxifen at the age of 8 weeks, and then bred with WT male mice until they are pregnant. We found that basal-only GFP⁺ multicell clones were extensively distributed in the basal layer at pregnancy day 5.5, whereas no bilineage clones were observed (Figure 4A). Quantification for GFP⁺ cells of basal cells shows that $7.42 \pm 1.60\%$ of basal cells were GFP⁺ (Figure 4A). This result suggests that *Nfatc1* reporter-marked basal epithelial cells are rapidly expanding at the early pregnancy stage. Quantification of GFP⁺ clones showed that the percentage of two-cell clones was 34.48% at this stage, and the proportion of multicell clones (≥ 3 cells) was 5.52% (Figure 4A). To validate this idea, we examined the proliferative status of GFP⁺ basal epithelial cells. We found that $42.43 \pm 0.01\%$ of GFP⁺ basal epithelial cells were Ki67⁺ at this stage, whereas $24.34 \pm 0.03\%$ of GFP⁻ basal epithelial cells were Ki67⁺ (Figure 4B). Collectively, these data suggest that *Nfatc1* reporter-marked cells and their progeny cells are more rapidly cycling relative to *Nfatc1* reporter-negative basal epithelial cells at an early pregnancy stage.

At pregnancy day 17.5, we detected basal-only GFP⁺ clones in both mammary ducts and alveoli (Figures 4C–4F). In the mammary duct, quantification analysis showed that the proportion of two-cell GFP⁺ clones was 32.86% at this stage, and the proportion of multicell GFP⁺ clones was 11.43%. Compared with pregnancy day 5.5, the proportion of multicell GFP⁺ clones becomes more robust (Figures 4A and 4C). In the alveoli, GFP⁺ clones were clearly visible using wholemout confocal imaging (Figure 4E). Immunostaining assays of mammary tissue sections showed that most GFP⁺ cells were located in the basal layer of the alveoli (Figures 4F and S5B). Quantification for GFP⁺ cells of basal cells shows that $10.95 \pm 1.27\%$ of basal cells were GFP⁺ in the mammary alveoli (Figure 4F). The proportion of two-cell GFP⁺ clones was 24.37% and that of multicell GFP⁺ clones was 6.09% in the alveoli (Figures 4F and S5B). Compared with the duct, the slight decrease in the proportion of multicell GFP⁺ clones is likely owing to the sphere

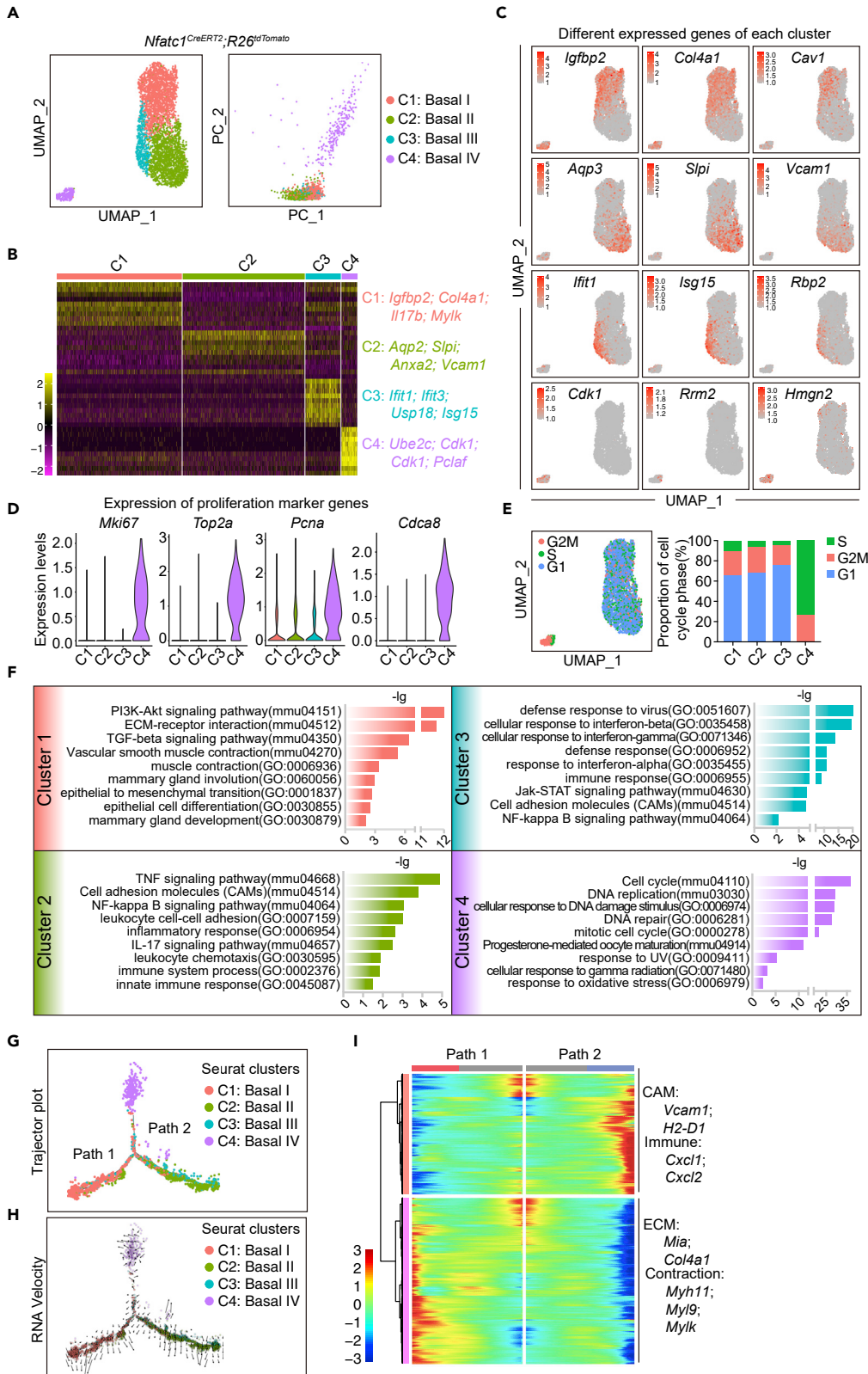


Figure 5. *Nfatc1*-lineage basal epithelial are heterogeneous during pregnancy (see also Figure S6)

- (A) UMAP and PCA plots reveal cellular heterogeneity of 4,203 tdTomato⁺ cells that were sorted from *Nfatc1*^{CreERT2}; *R26*^{tdTomato} mammary glands at the first pregnancy day 14.5
- (B) Heatmap showing differentially expressed genes in each cluster. Selected signature genes are shown on the right
- (C) Feature plots showing expression distribution of selected signature genes for each cluster
- (D) Violin plots showing genes functioning in cell proliferation
- (E) Cell cycle analysis based on the UMAP plot of *Nfatc1*-lineage basal epithelial cells at pregnancy day 14.5. The percentages of cells in the S, G2/M and G1 phases were quantified in each cluster
- (F) KEGG and GO analysis of each cluster using the top 250 differentially expressed genes in each cluster
- (G) Pseudotime ordering of *Nfatc1*-lineage basal epithelial cells at pregnancy day 14.5.
- (H) RNA velocity of *Nfatc1*-lineage basal epithelial cells on pregnancy day 14.5
- (I) scEpath analysis identifying two gene clusters of branching genes

morphology of mammary alveoli. These results suggest that *Nfatc1* reporter-marked basal epithelial cells are activated during early pregnancy and contributed to lobuloalveolar development.

Next, we examined the contribution of *Nfatc1* reporter-marked basal epithelial cells to lobuloalveolar development during the third round of pregnancy after TAM induction. Interestingly, more basal-only GFP⁺ clones were found surrounding the alveoli, as assayed by wholemount confocal imaging (Figure 4G). Co-immunofluorescence for GFP, K14, and K8 showed that most GFP⁺ clones were located in the basal layer (Figures 4H and S5C). Quantification for GFP⁺ cells of basal cells shows that 21.22 ± 0.96% of basal cells were GFP⁺ in the mammary alveoli (Figure 4H). The percentage of GFP⁺ cells increased about two-fold than that at the same stage during first pregnancy (Figures 4F and 4H). Quantification of GFP⁺ basal clones showed that the percentage of two-cell GFP⁺ basal clones was 34.48% and that of multicell GFP⁺ basal clones was 29.89% at the third pregnancy day 17.5, whereas they were 24.37 and 6.09%, respectively, at the same stage during the first pregnancy (Figures 4F, 4H, and 4I). Taken together, these data provide strong evidence that *Nfatc1* reporter-marked basal epithelial cells have stronger contribution to lobuloalveolar formation during third pregnancy than first pregnancy.

We also assessed the frequencies of *Nfatc1* reporter-marked luminal cells during pregnancy. We found that the proportion of luminal GFP⁺ cells was less than 1% at both pregnancy day 17.5 in the first round of pregnancy and pregnancy day 17.5 in the third round of pregnancy (Figures S5D and S5E). This suggests that the contribution of *Nfatc1* reporter-marked luminal cells to lobuloalveolar formation is limited. However, it is interesting that a small number of large luminal GFP⁺ clones containing dozens of cells were observed at pregnancy day 17.5 during both the first and third rounds of pregnancy (Figures S5D–S5G). This suggests that it is a minimal number of *Nfatc1* reporter-marked luminal cells rather than most *Nfatc1* reporter-negative luminal cells that are capable of generating luminal alveolar cells during pregnancy.

***Nfatc1*-lineage basal epithelial cells are heterogeneous during pregnancy**

To further understand the contribution of *Nfatc1* reporter-marked epithelial cells to lobuloalveolar development, we performed scRNA-seq analysis on *Nfatc1* reporter-marked cell progeny at pregnancy day 14.5. *Nfatc1*^{CreERT2}; *R26*^{tdTomato} mice were administered three doses of TAM at the age of eight weeks and then bred with WT male mice until pregnancy. We isolated Lin[−]CD24⁺tdTomato⁺ cells from the mammary gland on pregnancy day 14.5 and subjected them to droplet-enabled 3'-end scRNA-seq. After quality control (Figure S6A), a total of 6,745 cells were used for downstream analysis. Unsupervised clustering identified seven clusters (Figure S6B), which included mesenchymal cells, endothelial cells, muscle cells, immune cells, luminal epithelial cells, and two clusters of basal epithelial cells based on signature genes (Figures S6C and S6D). After removing non-epithelial cells and luminal epithelial cells, tdTomato⁺ basal epithelial cells were regrouped into four clusters (Figure 5A). We utilized the differentially expressed gene signatures to assign putative cell type identities to these clusters (Figures 5B and 5C). PCA analysis showed that C4 is obviously distinct from C1 to C3 (Figure 5A). The cells in cluster 4 were enriched for the proliferating cell marker genes, *Mki67*, *Top2a*, and *Pcna* (Figure 5D). In agreement, cell cycle analysis showed that a majority of C4 cells were in S and G2/M phases, whereas most cells in C1–C3 were in G0/G1 phase (Figure 5E). GO and KEGG analyses showed that the pathways of cell cycle, DNA replication and DNA repair were enriched in C4 cells (Figure 5F). Thus, C4 cells are referred to “cycling basal epithelial cells.” C1 cells were enriched for genes functioning in muscle contraction, the PI3K-Akt signaling pathway and the TGFβ signaling pathway, suggesting a potentially contractile feature in myoepithelial cells. C2 cells were enriched for

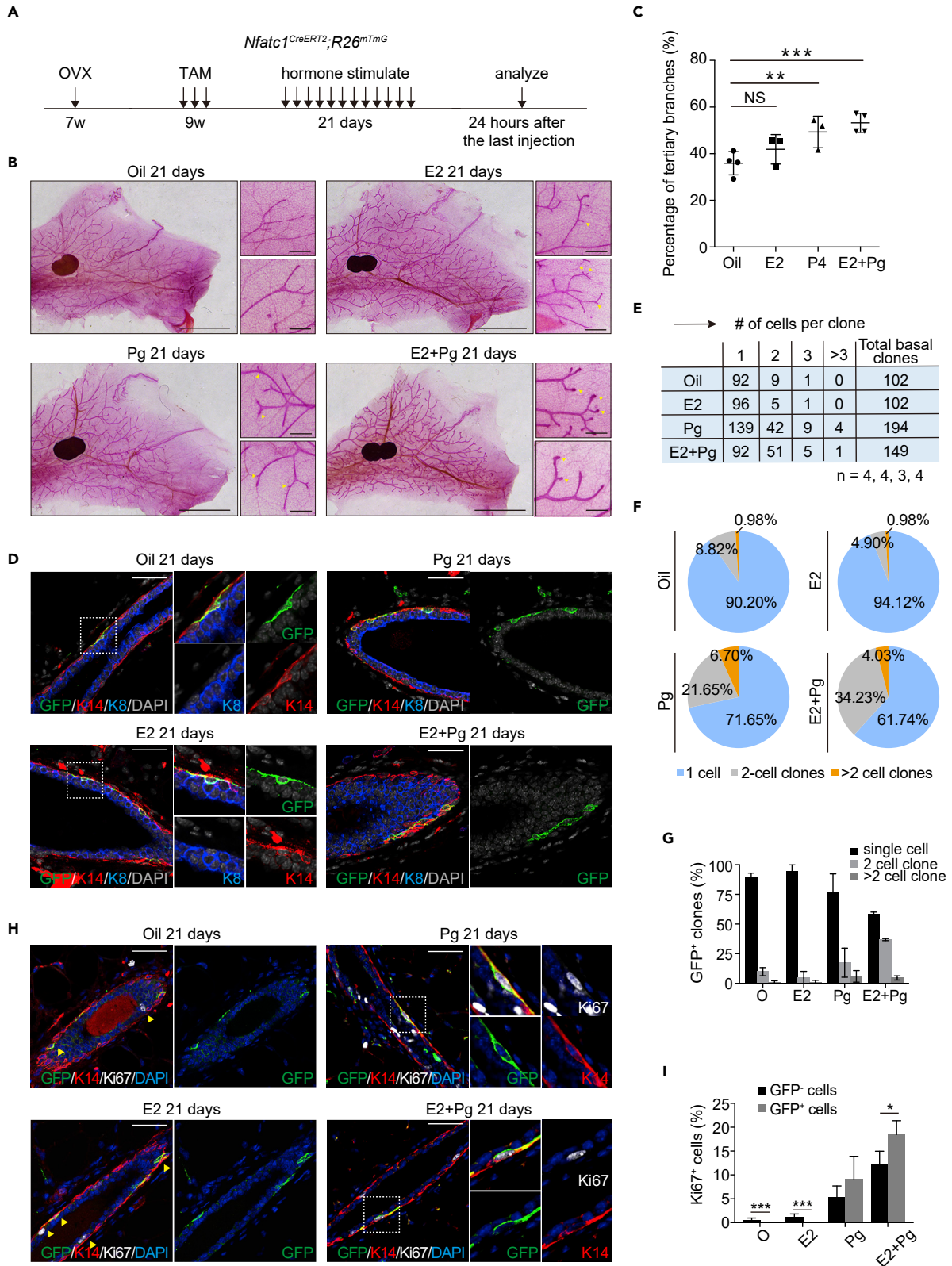


Figure 6. *Nfatc1* reporter-marked basal epithelial cells are activated by progesterone signal in ovariectomized (OVX) mice

(A) Experimental strategy for hormone treatment and the lineage tracing

(B) Wholmount images showing mammary glands of *Nfatc1*^{CreERT2};R26^{mTmG} OVX mice that were stimulated for 21 days with hormones (oil, E2, Pg, and E2+Pg). Scale bar, 5 mm. For each panel, high-magnification images are shown on the right. Arrowheads indicated tertiary branches. Scale bar, 500 μ m

(C) Quantification of tertiary branches in the ducts of *Nfatc1*^{CreERT2};R26^{mTmG} OVX mice after 21 days of stimulation with hormones (oil, E2, Pg, and E2+Pg); $n = 4, 3, 3,$ and 4 mice, respectively

(D) Co-immunofluorescence assay of GFP (green), K14 (red), and K8 (blue) in mammary glands from *Nfatc1*^{CreERT2};R26^{mTmG} OVX mice after 21 days of stimulation with hormones (oil, E2, Pg, and E2+Pg). Scale bar, 50 μ m

(E–G) Quantification analysis shows the number (E) and the percentage (G) of single-cell, two-cell, and multicell clones in *Nfatc1*^{CreERT2};R26^{mTmG} OVX mice after 21 days of stimulation with hormones (oil, E2, Pg, and E2+Pg); $n = 4, 4, 3,$ and 4 mice, respectively. Another statistical method is also shown in (F). The total numbers of basal clones were 102, 102, 194, and 149

(H and I) Co-immunofluorescence of GFP (green), K14 (red), and Ki67 (white) in mammary glands from *Nfatc1*^{CreERT2};R26^{mTmG} OVX mice after 21 days of stimulation with hormones (oil, E2, Pg, and E2+Pg) (H). Scale bar, 50 μ m. Quantification analysis shows the percentages of Ki67⁺GFP⁺ cells versus GFP⁺ cells and Ki67⁺GFP⁻ cells versus GFP⁻ cells under hormone treatment. In total, 83 (Oil), 91 (E2), 180 (Pg), and 101 (E2+Pg) GFP⁺ cells and 567 (Oil), 1,482 (E2), 2,045 (Pg), and 1,783 (E2+Pg) GFP⁻ cells were quantified, respectively (I). $n = 3$ mice for each treatment. Data represent the mean value \pm SD NS, not significant, * $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$

genes in the TNF signaling pathway, cell adhesion molecules (CAM), and the NF- κ B signaling pathway, suggesting a putative immune function. C3 cells were specifically enriched for gene functioning in interferon response. Therefore, *Nfatc1*-lineage basal epithelial cells include four functionally distinct clusters during pregnancy.

To examine the potential lineage relationships among these basal cell clusters, we performed pseudotime analysis, which showed that the cells in C4 were distributed in a major pseudotime trajectory that bifurcated to contractile basal epithelial cells (C1) and immunity-related basal epithelial cells (C2), whereas C3 cells were evenly distributed in the two directions (Figures 5G and S6E). We proposed a hypothesis that cycling basal progenitor cells give rise to functionally distinct differentiated cells. This idea was further supported by RNA velocity (Figure 5H). A large number of differentially expressed genes were presented along the pseudotime trajectory. We identified 1327 “branching” genes that are potentially important for the differentiation of contractile basal epithelial cells versus immunity-related basal epithelial cells (Figure 5H). In agreement, we found that genes related to vascular smooth muscle contraction (*Myh11*, *Myl9*) and ECM-receptor interaction (*Col4a1*, *Col4a2*) were upregulated in path 1 (Figures 5H and S6F). In contrast, genes related to CAM (*Vcam1*, *H2-D1*) and immune function (*Cxcl1*, *Cxcl2*) were upregulated in path 2 (Figures 5I and S6F). Taken together, our computational analysis suggests a two-branch lineage differentiation trajectory for the four distinct basal epithelial cell clusters during pregnancy.

***Nfatc1* reporter-marked basal epithelial cells become activated in ovariectomized (OVX) mice in response to progesterone**

Finally, we asked whether *Nfatc1* reporter-marked basal epithelial cells can be activated in response to hormones. To this end, we surgically removed the ovaries from *Nfatc1*^{CreERT2};R26^{mTmG} female mice as previously described (Zhao et al., 2010), to exclude the effect of endogenous hormones. After three doses of TAM induction, we intraperitoneally injected exogenous hormones (oil, E2 only, Pg only, or E2+Pg) to OVX mice every day for 21 days (Figure 6A). Upon E2+Pg treatment, enlarged terminal ducts (TDs) were extensively present in the mammary gland and the number of tertiary branches was significantly increased relative to vehicle control (Figures 6B and 6C), suggesting that E2+Pg treatment can mimic the hormone conditions of pregnancy. After 21 days of lineage tracing, the proportions of GFP⁺ two-cell and multicell basal clones markedly increased in response to the Pg or E2+Pg treatments compared to the vehicle control and E2 only (Figures 6D–6G). Moreover, co-immunofluorescence assay for GFP, Ki67, and K14 showed that the percentage of Ki67⁺GFP⁺ cells versus GFP⁺ cells is significantly higher than that of Ki67⁺GFP⁻ cells versus GFP⁻ cells under E2+Pg treatment. In contrast, the percentage of Ki67⁺GFP⁺ cells versus GFP⁺ cells is dramatically lower than that of Ki67⁺GFP⁻ cells versus GFP⁻ cells under conditions of no hormone treatment or only E2 treatment (Figures 6H and 6I). Taken together, these data suggest that *Nfatc1* reporter-marked basal cells are activated to rapidly divide in response to E2+Pg treatment and contribute to the expansion of mammary epithelial cells more efficiently than GFP⁻ basal epithelial cells.

In summary, we have identified *Nfatc1* reporter-marked basal mammary epithelial cells as dormant stem/progenitor cells. They divide more rapidly than *Nfatc1* reporter-negative basal cells during pregnancy and contribute to the lobuloalveolar development of the mammary gland during pregnancy.

DISCUSSION

Here we have identified *Nfatc1* reporter-marked basal epithelial cells as dormant unipotent stem/progenitor cells in virgin mammary gland. These cells show limited contribution to mammary epithelium homeostasis, but they expand more rapidly than *Nfatc1* reporter-negative cells during pregnancy to fuel mammary lobuloalveolar development. As such, this work points to the function of dormant mammary stem/progenitor cells in contributing to pregnancy-induced lobuloalveoli formation. The behaviors of *Nfatc1* reporter-marked cells are reminiscent of quiescent mammary stem cells such as *Lgr5*⁺*Tspan8*^{high} cells and *Bcl11b*⁺ cells. Fu et al. have identified that *Lgr5*⁺*Tspan8*^{high} cells as quiescent MaSCs that located at proximal region and can be activated by ovarian hormones during pregnancy (Fu et al., 2017). Cai et al. have demonstrated that the expression level of *Bcl11b*, an intrinsic regulator of mammary stem cell quiescence, is markedly downregulated during pregnancy, whereas a small population of quiescent mammary stem cells express high levels of *Bcl11b* at homeostasis (Cai et al., 2017), supporting the idea that quiescent MaSCs are activated during pregnancy. Interestingly, scRNA-seq analysis showed that *Nfatc1* is barely overlapped with *Lgr5*, *Tspan8*, and *Bcl11b*. It appears that *Nfatc1* reporter-marked basal cells represent a subpopulation of dormant mammary stem cells.

In addition to quiescent MaSCs, it has been reported that WNT signaling-responsive *Procr*⁺ and *Axin2*⁺ MaSCs can also contribute to alveoli formation (van Amerongen et al., 2012; Wang et al., 2015). *Procr*⁺ cells were identified as multipotent mammary stem cells that is critical for both the homeostasis of mammary gland development and the alveolar formation during pregnancy (Wang et al., 2015). In comparison, *Nfatc1* reporter-marked basal cells have limited contribution to the homeostasis of mammary gland development, but have bias to make contributions to the alveolar formation during pregnancy. It is worth noting that the percentage of *Procr*- and *Axin2*-marked cells remained unchanged after multiple pregnancies, but the percentage of *Nfatc1* reporter-marked cells increases three times after three rounds of pregnancies. Therefore, although both active and dormant mammary stem/progenitor cells contribute to mammary lobuloalveoli formation, the dormant *Nfatc1* reporter-marked stem/progenitor cells are preferentially used for multiple pregnancies. We speculate that this might represent a protective mechanism to ensure robust formation of functional mammary lobuloalveoli that secrete milk for feeding offspring, as the genomic DNA in dormant mammary stem/progenitor cells is relatively stable.

Recently, scRNA-seq analysis has been extensively utilized for characterizing mammary epithelial cells. Bach et al. demonstrated that the identity of basal epithelial cells is different at distinct developmental stages, whereas basal epithelial cells are homogeneous at the virgin or gestation stage (Bach et al., 2017). Pal et al. have also shown the same idea that basal epithelial cells are homogeneous at virginal stage (Pal et al., 2017). Interestingly, we found that *Nfatc1* reporter-marked basal epithelial cells are generally homogeneous at the adult virginal stage, but they can be divided into two distinct groups based on the expression levels of genes functioning in muscle contraction, suggesting that there are differences in contractile capacity in different basal epithelial cells. Notably, *Nfatc1*-lineage cells are heterogeneous during pregnancy, including four clusters that appear functionally distinct. It appears that cycling progenitor cells bifurcate into two distinct groups, contractile basal epithelial cells, and immunity-related basal epithelial cells. We noticed that *Rrm2* and *Hmgn2*, which are specifically and highly expressed in the population of cycling progenitor cells, function as important effectors of progesterone signaling to induce cell proliferation (Lei et al., 2012; Li et al., 2017), raising the possibility that these cycling progenitor cells are responsive to progesterone signaling. It is well known that mammary basal epithelial cells, also called myoepithelial cells, have contractile functions, which are essential for milk ejection (Haaksma et al., 2011; Sopel, 2010). However, GO and KEGG analyses indicate that a subset of basal epithelial cells might be important for immune function including leukocyte cell-cell adhesion and leukocyte chemotaxis. In agreement with this idea, the high levels of *Vcam1* in this immunity cluster can mediate the accumulation of IgA antibody secreting cells (Low et al., 2010). Thus, our findings provide evidence that, in addition to the known heterogeneity that there are different subsets of MaSCs, mammary basal epithelial cells contain functionally distinct clusters.

Our work uncovers a specific population of basal mammary epithelial cells marked by the *Nfatc1* expression. These cells are dormant basal stem/progenitor cells that contribute to mammary lobuloalveoli formation during pregnancy and are preferentially used for multiple pregnancies. The study provides significant insights for the cellular mechanisms of mammary lobuloalveoli formation.

Limitations of study

Our study aims to study the function of *Nfatc1*⁺ mammary stem/progenitor cells in contributing to lobuloalveoli formation during pregnancy. We label *Nfatc1*⁺ cells using *Nfatc1*^{CreERT2};*R26*^{mTmG} mice upon

tamoxifen induction. However, it is hard to determine whether all of *Nfatc1* reporter-marked basal mammary epithelial cells are *Nfatc1*⁺ cells. The first reason is that *Nfatc1* reporter-marked cells might contain a small population of *Nfatc1*⁺ cell-derived progeny cells, as the mammary samples were analyzed after three-time tamoxifen induction. Another reason is that only one allele of *Nfatc1* can express its protein in *Nfatc1*^{CreERT2;R26^{mTmG}} mice owing to the knocking of *CreERT2*, leading to a remarkable reduction of *Nfatc1* protein. Given that the reduced *Nfatc1* protein is not detectable, it is hard to determine whether *Nfatc1* reporter-marked epithelial cells are *Nfatc1*⁺ cells using double immunofluorescence for GFP and *Nfatc1*. Thus, we use “*Nfatc1* reporter-marked mammary epithelial cells” in this study.

STAR★METHODS

Detailed methods are provided in the online version of this paper and include the following:

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SUPPLEMENTAL INFORMATION

Supplemental information can be found online at <https://doi.org/10.1016/j.isci.2022.103982>.

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AUTHOR CONTRIBUTIONS

Conceptualization: J.S, Z.Y., and C.L.; methodology: R.L., C.L., and H.H.; investigation: R.L., M.M., Z.X., X.B., P.L., and H.H.; resources: H.H. and C.F.G.-J.; writing – original draft: R.L.; visualization: R.L.; writing – review and editing: Z.X., C.L., Z.Y., J.S., X.D., and M.V.P.; funding acquisition: J.S. and Z.Y.; supervision, J.S, Z.Y., and C.L.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Rabbit anti-keratin 8	Abcam	Cat# ab59400; RRID: AB_942041
Mouse anti-keratin 14	Abcam	Cat# ab9220; RRID: AB_307087
Chicken anti-GFP	Abcam	Cat# ab13970; RRID: AB_300798
Rabbit anti-keratin 5	Abcam	Cat# ab52635; AB_869890
Mouse anti-Ki67	Abcam	Cat# ab16667; RRID: AB_302459
Mouse anti-NFAT2	Abcam	Cat# ab2796; RRID: AB_303308
Mouse anti-RFP	Invitrogen	Cat# MA5-15257; RRID: AB_10999796
Anti-Rat CD45-APC	eBioscience	Cat# 17-0451-82; RRID: AB_469392
Anti-Rat CD31-APC	eBioscience	Cat# 17-0311-82; RRID: AB_657735
Anti-Rat TER 119-APC	eBioscience	Cat# 17-5921-82; RRID: AB_469473
Anti-Mo/Rat CD24-PeCy7	eBioscience	Cat# 25-0291-82; RRID: AB_1234962
Anti-Mo CD24-BV711	BD Bioscience	Cat# 563450; RRID: AB_2738213
Anti-Mo/Rat CD29-PeCy7	eBioscience	Cat# 25-0242-82; RRID: AB_10853806
Anti-Rat CD29-FITC	BD Bioscience	Cat# 561796; RRID: AB_10894590
Chemicals, peptides, and recombinant proteins		
Tamoxifen (TAM)	Sigma-Aldrich	Cat# T5648
5-ethynyl-20-deoxyuridine (EdU)	Thermo Fisher	Cat# A10044
Collagenase IV	Gibco	Cat# 17104019
Hyaluronidase	Merck	Cat# H1115000
Dispase	Stem Cell Technologies	Cat# 07913
Ammonium chloride solution	Stem Cell Technologies	Cat# 07850
DNase I	Sigma-Aldrich	Cat# D7291
Fixable Viability Dye	eBioscience	Cat# 65-0863-14
Matrigel	Corning	Cat# 356231
Epicult-B medium	Stem Cell Technology	Cat# 05610
Human Recombinant EGF	Stem Cell Technology	Cat# 78006.1
Human Recombinant bFGF	Stem Cell Technology	Cat# 78003.1
Heparin solution	Stem Cell Technology	Cat# 07980
Critical commercial assays		
Click-iT Edu Alexa Flour 594 Kit	Beyotime	Cat# C00785
Single cell 3 'Library and Gel Bead Kit V2	10x Genomics	Cat# 120237
Deposited data		
Single-cell RNA sequencing data	This paper	GEO: GSE175432
Experimental models: Organisms/strains		
Mouse: <i>Nfatc1</i> ^{CreERT2}	This paper	N/A
Mouse: <i>R26</i> ^{mTmG}	The Jackson Laboratory	JAX# 007676
Mouse: <i>R26</i> ^{tdTomato}	The Jackson Laboratory	JAX# 007914
Mouse: <i>Procr</i> ^{CreERT2-IRES-tdTomato}	Zeng's laboratory	N/A

(Continued on next page)

Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Oligonucleotides		
Genotyping primer, <i>Nfatc1</i> ^{CreERT2} -P1: CCTCCGGCCAATTCACAATC	This paper	N/A
Genotyping primer, <i>Nfatc1</i> ^{CreERT2} -P2: AGGCTTCGCTTTTCTTTGAG	This paper	N/A
Genotyping primer, <i>Nfatc1</i> ^{CreERT2} -P3: GGTCAGTAAATTGGAGCCCA	This paper	N/A
Software and algorithms		
GraphPad Prism 7	GraphPad	N/A
Cellranger 2.0.1		
	10x Genomics	https://support.10xgenomics.com/singlecell-gene-expression/software/downloads/latest
Seurat 3.2.2		
	Stuart et al., 2019	https://satijalab.org/seurat
Monocle 2.10.1	Qiu et al., 2017	http://cole-trapnell-lab.github.io/monocle-release/
Velocity.R	La Manno et al., 2018	https://github.com/velocity-team/velocity.R
FlowJo	BD	N/A

RESOURCE AVAILABILITY**Lead contact**

Further information and requests for resources and reagents should be directed to and will be fulfilled by the lead contact, Cong Lv (lvc@cau.edu.cn)

Materials availability

Mouse lines generated in this study are available upon request to Lead Contact provided the requestor covers shipping costs.

Data and code availability

All scRNA-seq data from this study are available at NCBI Gene Expression Omnibus (GEO). The accession number for data reported in this paper is GEO: GSE175432. Microscopy data reported in this paper will be shared by the lead contact upon request. All generic and custom R scripts are available on reasonable request. Any additional information required to reanalyze the data reported in this paper is available from the lead contact upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

All mouse experimental procedures and protocols were evaluated and authorized according to the Beijing Regulations for Laboratory Animal Management. They were strictly in accordance with the guidelines of the Institutional Animal Care and Use Committee of China Agricultural University (approval number: SKLAB-2019-04-03). *Nfatc1*^{CreERT2} mice were generated at Shanghai Biomodel Organism Science & Technology Development Co., Ltd. *R26*^{mTmG} mice and *R26*^{tdTomato} mice were purchased from Jackson Laboratories (stock number: 007676, 007914). *Procr*^{CreERT2-IRES-tdTomato} mice were obtained from Zeng's laboratory at the Chinese Academy of Sciences, Shanghai. Three-week old female NOD-SCID mice were purchased from Beijing Vital River Laboratory Animal Technology Co., Ltd. Female age-matched mice were utilized for all experiments. Eight-week old female mice were utilized for short-term labeling, long-term tracing and pregnancy experiments. Seven-week old female mice were used for ovariectomized and hormone stimulation.

METHOD DETAILS**Lineage tracing**

To label *Nfatc1*⁺ cells at homeostasis, the mice were intraperitoneally injected with tamoxifen (Sigma-Aldrich, T5648) diluted in corn oil (Sigma-Aldrich, C8267) at a dose of 4 mg pre 25 g body weight. Female mice

were administered tamoxifen 3 times every other day at the age of 8 weeks. For the pregnancy experiment, female mice were administered tamoxifen 3 times every other day at the age of 8 weeks and then mated to WT mice at 2 weeks post-induction.

Hormone stimulation

The protocol for ovariectomy and hormone stimulation was described in previous reports (Zhao et al., 2010). In brief, female mice were bilaterally ovariectomized at the age of 7 weeks and then recovered for 2 weeks. After 3 doses of TAM induction every other day, the mice were intraperitoneally injected with 0.1 µg 17β-estradiol (Sigma, E2758) and/or 0.1 mg progesterone (Sigma, P0130) diluted in 100 mL corn oil every other day for 21 days. Mammary glands were harvested for further analysis 24 hours after the last induction.

Immunofluorescence (IF)

Mammary glands were fixed overnight in 4% paraformaldehyde, washed with PBS and then processed and embedded in paraffin. Paraffin-embedded tissues were sectioned at 4-µm. Tissue sections deparaffinized in xylene, and then rinsed in ethanol and washed in water. Antigen retrieval was performed by heating the slides in 0.01 M citrate buffer (pH 6.0) in a microwave. The sections were cooled naturally to room temperature (RT) and then blocked for 1 hour at RT with blocking solution. The sections were incubated with primary antibodies overnight at 4 °C and incubated with secondary antibodies (Life Technologies) for 1 hour at RT. The slides were then washed 3 times with PBS and stained with DAPI for 5 min. The sections were covered with mounting medium. Antibodies against the following proteins were used: keratin 8 (Abcam, Ab59400, 1:400), keratin 14 (Abcam, Ab9220, 1:400), GFP (Abcam, Ab13970, 1:1000), keratin 5 (Covance, 905501, 1:800), Ki67 (Abcam, Ab16667, 1:1000), NFAT2 (Abcam, Ab2796, 1:400), and RFP (Invitrogen, MA5-15257).

Wholemout confocal imaging

For wholemount confocal imaging, samples were dissected under a Leica M165FC stereomicroscope to yield portions containing a large area of ductal network (approximately 3 mm x 3 mm). Adipose tissue was removed using dissecting scissors (Rios et al., 2014). Small pieces of tissue were cleared in 80% glycerol overnight and then covered with mounting medium.

EdU labeling

EdU (Thermo Fisher, A10044) (0.2 mg per 10 g body weight) was intraperitoneally injected into 8-week-old wild-type (WT) female WT mice. The mammary glands were harvested 3 hours after injection. Co-staining of EdU and K14 was performed according to the manufacturer's instructions using the Click-iT EdU Alexa Fluor 594 kit (Beyotime, C0078S).

Primary cell preparation

Single MECs were obtained from mammary glands following a published protocol (McNally and Stein, 2017). Mammary glands from female mice were isolated at indicated timepoints. The minced tissue was placed in culture medium (DMEM/F12 (Gibco) with 2% fetal bovine serum, 300 U/mL collagenase IV (Gibco, 17104019), and 300 U/mL hyaluronidase (Merck, H1115000)) and digested for 2 hours at 37 °C. After lysis of the red blood cells with ammonium chloride solution (Stem Cell Technologies, 07850), the suspension was incubated with 0.25% trypsin-EDTA at 37 °C for 2 min. A single-cell suspension was obtained via incubation with 1 U/mL dispase (Stem Cell Technologies, 07913) plus 0.1 mg/mL DNase I (Sigma, D7291) for 3 min with gentle pipetting, followed by filtration through 40 µm cell strainers.

Cell labeling and flow cytometry

Antibody incubation was performed on ice for 15 min in PBS with 2% fetal bovine serum. The primary antibodies employed were anti-CD24-PeCy7 (eBiosciences, 25-0291-82, 1:1000), anti-CD24-BV711 (BD Biosciences, 563450, 1:1000), anti-CD29-PeCy7 (eBiosciences, 25-0242-82, 1:1000), anti-CD29-FITC (BD Biosciences, 561796, 1:1000), anti-CD45-APC (eBiosciences, 17-0451-82, 1:1000), anti-CD31-APC (eBiosciences, 17-0311-82, 1:1000), anti-TER119-APC (eBiosciences, 17-5921-82, 1:1000) and anti-Dyeflour450 (eBiosciences, 65-0863-14, 1:1000). Flow cytometry analysis was performed using Aria SROP (BD). The data were analyzed using FlowJo software. For sorting cells, an Aria III (BD) instrument was used.

In vitro colony formation assay

Colony formation assays were performed according to the manufacturer's protocol (Stem Cell Technologies, 05610). Freshly sorted tdTomato⁺ and tdTomato⁻ cells were embedded in 50% Matrigel (Corning, 356231) and cultured in mouse Epicult-B medium (Stem Cell Technology, 05610) containing 5% FBS, 20 ng/mL Human Recombinant EGF (Stem Cell Technology, 78006.1), 20 ng/mL Human Recombinant bFGF (Stem Cell Technology, 78003.1), and 4 µg/mL heparin solution (Stem Cell Technology, 07980). FACS-sorted cells were resuspended at a density of 80,000 cells/mL in chilled 100% growth-factor-reduced Matrigel (BD Bioscience). The medium was changed every 2 days, and colonies were scored after 7~8 days. Colony size was analyzed with Adobe Photoshop CC 2018.

Mammary fat pad transplantation

Sorted tdTomato⁺ cells (100, 500 or 1000 cells) were resuspended in 50% Matrigel and injected into the cleared fat pads of 3-week-old female NOD-SCID mice in a volume of 10 µL. Reconstituted mammary glands were harvested 8-10 weeks post-surgery. Outgrowths were detected under a dissection microscope (Leica M165FC) after Carmine staining. The repopulation frequency was calculated using the method at <http://bioinf.wehi.edu.au/soft-ware/elda/> (Hu and Smyth, 2009).

Wholemount carmine staining

Inguinal mammary glands were spread on glass slides and fixed in Carnoy's fixative (6:3:1, 100% ethanol: chloroform: glacial acetic acid) for 24 hours at room temperature, and then were washed in 70% ethanol for 15 min and rinsed with graded alcohol followed by distilled water for 5 min. The mammary gland was stained in carmine alum for 30 min and then dehydrated in 70%, 95% and 100% ethanol. Finally, the mammary gland was cleared in xylene and mounted with neutral gum.

Microscope image acquisition

All stained samples were mounted in Antifade Mounting Medium (P0128S; Beyotime), microscopy was performed at room temperature. For general immunofluorescence, image were captured by a microscope (LEICA DM6 B) with LEICA DFC9000 GT camera. 20× U-Plan-Apochromat/0.70 NA and 40× U-Plan-Apochromat/ 0.85 NA objective lenses were used. Wholemount confocal images were acquired using a confocal laser scanning microscope (Nikon A1). Fluorochromes included Alexa Fluor 488, Alexa Fluor 594, Alexa Fluor 647, and all nuclei were detected by DAPI staining. For wholemount carmine staining, image were captured by LEICA M165 FC with LEICA DFC450 C camera.

Quantification of lineage-specific cells and size of GFP⁺ clones

In *Nfatc1*^{CreERT2;R26^{mTmG}} mammary glands, clones were defined as cell clusters containing one or more GFP⁺ cells that contacted each other, as described in previous studies (Wang et al., 2015). A minimum of 3 mice with GFP⁺ clones on more than 20 sections were analyzed at each timepoint. For each clone, the number of cells was scored according to co-staining of K14 and K8. Images of representative clones were captured by microscopy (Leica DM6-B).

Single-cell mRNA sequencing

A single-cell suspension of the mammary gland epithelium was prepared as described above. TdTomato⁺ cells were sorted into EP tubes in single-cell mode using an Aria III (BD Biosciences) instrument. The collected cells were held on ice until they were loaded onto the GemCode single-cell platform (10×). Chromium Single Cell 3' v2 libraries were sequenced with a NovaSeq 6000 sequencer with the following sequencing parameters: read 1, 150 cycles; i7 index, 8 cycles; and read 2, 150 cycles.

For the sample of cells isolated from 8-week-old *Nfatc1*^{CreERT2;R26^{tdTomato}} mice 48 hours after three pulses of TAM induction, raw Illumina data were demultiplexed and processed using Cell Ranger (10× Genomics version Cell Ranger 2.0.1) and then mapped to the mouse mm10 reference genome with transcriptome version mm10-1.2.0. The alignment metrics based on the web summary were as follows: total number of reads: 484,349,675; reads confidently mapped to the transcriptome: 68.8%; estimated number of cells: 2,604; mean reads per cell: 186,002; median genes per cell: 1,964; total genes detected: 18,299; and median UMI counts per cell: 5,516.

For the sample of *Nfatc1* reporter-marked cell progeny at pregnancy day 14.5, raw Illumina data were demultiplexed and processed using Cell Ranger (10× Genomics version Cell Ranger 2.0.1) and then mapped to the mouse mm10 reference genome with transcriptome version mm10-3.0.0. The alignment metrics based on the web summary were as follows: total number of reads: 467,349,027; reads mapped to the genome: 91.9%; reads mapped confidently to the genome: 88.9%; reads confidently mapped to the transcriptome: 74.3%; estimated number of cells: 8,095; mean reads per cell: 57,733; median genes per cell: 1,936; total genes detected: 19,245; and median UMI counts per cell: 5,823.

Cluster identification using seurat

Seurat version 3.2.2 was used for filtering and subsequent clustering (Stuart et al., 2019). To remove partial cells and doublets, quality control was performed, in which cells with less than 500 genes or more than 6,000 genes were removed. Additionally, considering that a high proportion of mitochondrial expression in cells is indicative of cell stress/damage during isolation, cells with more than 10% unique mitochondrial molecular identifiers (UMIs) were removed from both of the samples. Gene-cell matrices were normalized and scaled in Seurat using the default parameters for UMIs. Variable genes/features were identified using a total of 2,000 features. The data were then scaled using the *ScaleData* function. Principal component analysis (PCA) was performed using the highly variable genes identified. Neighbors and clusters were identified using the *FindNeighbors* and *FindClusters* functions, respectively, with dimensions specified by the user and visualized using UMAP. We used the *VlnPlot* function to highlight the expression of known marker genes for basal (*Krt5/K5*, *Krt14/K14*) and luminal (*Krt8/K8*, *Krt18/K18*) epithelial cells. Differentially expressed genes are shown using the *Featureplot* function. Cell cycle analysis was carried out in Seurat using a list of cell cycle genes. The gene list was obtained from the Regev laboratory (Kowalczyk et al., 2015). Differentially expressed genes were identified using the *FindAllMarkers* function with the following parameters enabled: *logfc.threshold* = 0.25 and *min.pct* = 0.25. The top 250 differentially expressed genes in each cluster were used for KEGG and GO analysis. Functional enrichment of gene sets with different expression patterns was performed using the Database for Annotation on <http://kobas.cbi.pku.edu.cn/kobas3> (Xie et al., 2011).

Reconstructing differentiation trajectories using monocle

To analyze the differentiation trajectories of *Nfatc1* reporter-marked basal epithelial cells and their progeny, Monocle version 2.10.1 was used on cells filtered from Seurat to infer differentiation trajectories (Qiu et al., 2017). An expression threshold of 0.01 was applied. The highly variable genes identified from Seurat were used as the ordering filter. DDRTree was used for dimension reduction. For the ordering_genes function, differentially expressed genes with *p_val* < 1e−10 were chosen to plot the cell trajectory. Branch analysis was also performed for differentially expressed genes with *p_val* < 1e−10.

Inferring cellular dynamics based on RNA velocity

We applied RNA velocity technology to infer the dynamic change process of *Nfatc1* reporter-marked basal epithelial cells and their progeny. In the preparation phase, the Python version of Velocity 0.17 was applied to generate loom files (<https://velocity.org/velocity.py/index.html>) (La Manno et al., 2018). The “run10x” parameter was set because the sequencing protocol was 10×. The “mm10_rmsk.gtf.gtf” file for removing expression duplicate units was from <https://genome.ucsc.edu/cgi-bin/hgTables>. The parameters for generating the “*.gtf” file were as follows: *clade* = “Mamal”; *genome* = “Mouse”; *assembly* = “Dec. 2011 (GRCm38/mm10)”; *group* = “All Tracks”; *track* = “RepeatMasker”; *table* = “rmsk”; *region* = “genome”; *output format* = “GTF-gene transfer format (limited)”; *output file* = “mm10_rmsk.gtf.” In the velocity estimation phase, velocity.R version v0.6 was applied to estimate RNA velocity using gene-relative slopes. The embeddings were derived from the pseudotime differentiation trajectory of basal cells through Monocle. We also used this approach to estimate the distance of the cells. “fit.quantile” was set to the default value of 0.02 to perform a gamma fit to the upper/lower magnitudes of expression. The amount of time to project the cells forward “deltaT” was set to 2. The number of 10 nearest neighbors (NN) was used in the slope calculation smoothing, i.e., *kCells* = 10. Default values were used for other parameters.

QUANTIFICATION AND STATISTICAL ANALYSIS

Most of the experiments were repeated at least three times and the exact *n* is stated in the corresponding figure legend. Quantification data were analyzed using GraphPad Prism software and are presented as the mean ± standard deviation (SD) (as indicated in the figure legends). Unpaired Student’s *t*-tests were performed, and asterisks indicate the levels of statistical significance (**P* < 0.05; ***P* < 0.01; ****P* < 0.001).