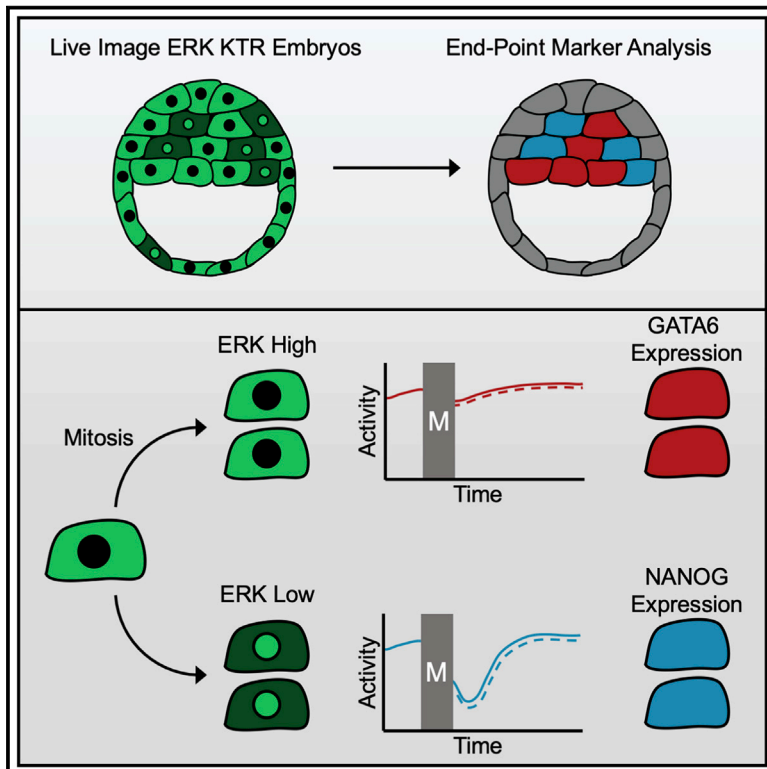


# Developmental Cell

## Cell-Cycle-Dependent ERK Signaling Dynamics Direct Fate Specification in the Mammalian Preimplantation Embryo

### Graphical Abstract



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### In Brief

Pokrass et al. discover a cell-cycle-dependent bifurcation of FGF-ERK signaling activity in cells selecting between fates in embryonic development. They show cells exiting mitosis exhibit either rapid ERK activation or pulsed ERK inhibition, and this signaling decision correlates with differentiation into the primitive endoderm or epiblast lineages, respectively.

### Highlights

- Kinase translocation reporters enable *in vivo* measurement of ERK signaling dynamics
- ERK activity bifurcates at mitotic exit in ICM blastomeres to specify EPI versus PrE
- Embryonic stem cells in culture recapitulate ERK bifurcation at mitotic exit
- Low ERK activity during G1 promotes rapid post-translational stabilization of NANOG



## Article

# Cell-Cycle-Dependent ERK Signaling Dynamics Direct Fate Specification in the Mammalian Preimplantation Embryo

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## SUMMARY

Despite the noisy nature of single cells, multicellular organisms robustly generate different cell types from one zygote. This process involves dynamic cross regulation between signaling and gene expression that is difficult to capture with fixed-cell approaches. To study signaling dynamics and fate specification during preimplantation development, we generated a transgenic mouse expressing the ERK kinase translocation reporter and measured ERK activity in single cells of live embryos. Our results show primarily active ERK in both the inner cell mass and trophectoderm cells due to fibroblast growth factor (FGF) signaling. Strikingly, a subset of mitotic events results in a short pulse of ERK inactivity in both daughter cells that correlates with elevated endpoint NANOG levels. Moreover, endogenous tagging of *Nanog* in embryonic stem cells reveals that ERK inhibition promotes enhanced stabilization of NANOG protein after mitosis. Our data show that cell cycle, signaling, and differentiation are coordinated during preimplantation development.

## INTRODUCTION

Mammalian embryogenesis initiates with a single fertilized oocyte that must rapidly increase in cell number and complexity to create a mature blastocyst of three lineages, the epiblast (EPI), primitive endoderm (PrE), and trophectoderm (TE) (Schrode et al., 2013). In contrast to other metazoan development, mammalian preimplantation development proceeds in the absence of any known maternal patterning cues, and thus, the initial fate specification events are determined from signals originating within the embryo itself (Chazaud and Yamanaka, 2016). While much is known about the signaling molecules and transcriptional regulators operating during this period (Ohnishi et al., 2014; Kang et al., 2013; Guo et al., 2010; Chazaud et al., 2006), a comprehensive understanding of the signaling and gene expression dynamics that allow robust selection of appropriate lineages is still missing.

Because of its capacity to self-organize and develop ex utero, the mouse blastocyst has emerged as an excellent model to understand how signaling coordinates cell fate determination (Simon et al., 2018). The first embryonic lineage segregation appears to be governed by polarity-mediated differential Hippo signaling that generates the TE, the apicobasal

polarized outer layer of cells defined by *Cdx2* expression, and the inner cell mass (ICM), defined by the pluripotency regulator *Sox2* (Frum et al., 2018; Wicklow et al., 2014; Hirate et al., 2013; Nishioka et al., 2009). Next, the ICM will further differentiate into the EPI and PrE lineages marked by NANOG and GATA6, respectively. While cells of the ICM initially coexpress the markers for both lineages, progenitors gradually upregulate the expression of one factor and concomitantly downregulate the other until all cells have been specified (Guo et al., 2010; Plusa et al., 2008). This bistability is achieved via cooperative feedforward and feedback mechanisms that allow both NANOG and GATA6 to increase transcription at their own promoter and inhibit expression of the opposing factor (Meng et al., 2018; Bessonard et al., 2014; Singh et al., 2007; Boyer et al., 2005). Individual ICM cells select between these lineages via fibroblast growth factor (FGF) signaling; however, the exact nature of the bifurcation of FGF-dependent signaling remains unclear.

Several lines of evidence support that ERK signaling downstream of fibroblast growth factor receptors (FGFRs) is essential for segregating these two lineages: (1) Inhibitors of FGFR or MEK and addition of recombinant FGF4 cause all ICM cells to select EPI or PrE fates, respectively (Saiz et al., 2016;

Yamanaka et al., 2010), (2) genetic ablation of FGF4, GRB2, or both FGFR1 and FGFR2 prevent the formation of the PrE lineage (Kang et al., 2017; Molotkov et al., 2017; Kang et al., 2013; Chazaud et al., 2006) and, (3) FGF4 and FGFR2 expression is mutually exclusive in ICM cells and these factors are early markers for the EPI and PrE lineages, respectively (Guo et al., 2010). Together, these findings supported a hypothesis that FGF4-expressing cells signal to neighboring FGFR2-expressing cells to generate bimodal ERK activity in the ICM essential to specify both lineages. However, recent studies reported that FGFR1 expression is required in all ICM cells to both generate the PrE and perhaps limit maximum NANOG concentrations in the EPI, indicating that a simple model consisting of cells either receiving or secreting the FGF4 stimulus is insufficient to describe the segregation of EPI and PrE lineages (Kang et al., 2017; Molotkov et al., 2017). Furthermore, ICM progenitors were observed to incrementally specify toward EPI or PrE lineages rather than all at once during a specific time in blastocyst development, suggesting a regulatory mechanism that could rely on other sources of cellular heterogeneity such as stochastic gene expression, asynchronous cell-cycle progression, or other signaling pathway activities (Saiz et al., 2016). While these experiments revealed the importance of ERK signaling in preimplantation development, the temporal patterns of ERK activity in single cells of the developing embryo have not been measured.

Live-cell imaging of biosensors has emerged as a powerful technique to study the dynamics of cellular decision making (Gaudet and Miller-Jensen, 2016). These approaches enable tracking of individual cells while obtaining high temporal resolution measurements of cellular parameters such as gene expression (Gu et al., 2018; Hafner et al., 2017; Wilson et al., 2017), metabolism (Lobas et al., 2019; Hung and Yellen, 2014), cell cycle (Sakaue-Sawano and Miyawaki, 2014), or signaling (Komatsu et al., 2011). Over the last decade, the use of biosensors in cultured cells, organoids, and even live organisms has shown that signaling dynamics can influence gene expression (Lane et al., 2017), collective cell behavior (Hino et al., 2020), and cell fate (Johnson and Toettcher, 2019; Muta et al., 2018). However, biosensors for kinase activity have not been used to understand mammalian development due to technical limitations including high phototoxicity and poor sensitivity. Here, we use kinase translocation reporter (KTR) technology (Regot et al., 2014) to study MAPK signaling dynamics during mammalian preimplantation development.

We generated ERK KTR mice to enable live single-cell analysis of kinase activity and validated them using multiple primary cell types as well as *in vivo* two-photon imaging. Focusing on preimplantation development, we characterized ERK signaling dynamics during cell specification and found that ICM cells exhibit a bifurcation of ERK activity after mitosis that ultimately drives daughter cells toward the EPI or PrE lineages. Furthermore, using endogenously tagged NANOG in embryonic stem (ES) cells, we show that NANOG is poised for rapid stabilization in G1. This work expands our understanding of how signaling dynamics and cell-cycle progression are coordinated to direct robust fate specification in the self-organizing mammalian blastocyst and ES cells.

## RESULTS

### Development and Validation of ERK KTR Mice

To study ERK signaling dynamics in primary mammalian cells and tissues, we generated a transgenic mouse line carrying the previously reported ERK KTR (Regot et al., 2014). The expression of the biosensor is restricted by a lox-stop-lox (LSL) cassette that, upon CRE-mediated recombination, allows spatiotemporally controlled expression of the sensor driven by the CAG promoter. Because KTR technology depends on accurate recognition of the nuclear boundary for quantitative kinase activity measurements, our vector contains a bicistronic ORF encoding both H2B-mRuby2 and KTR-mClover separated by the P2A peptide to allow equimolar expression of both proteins.

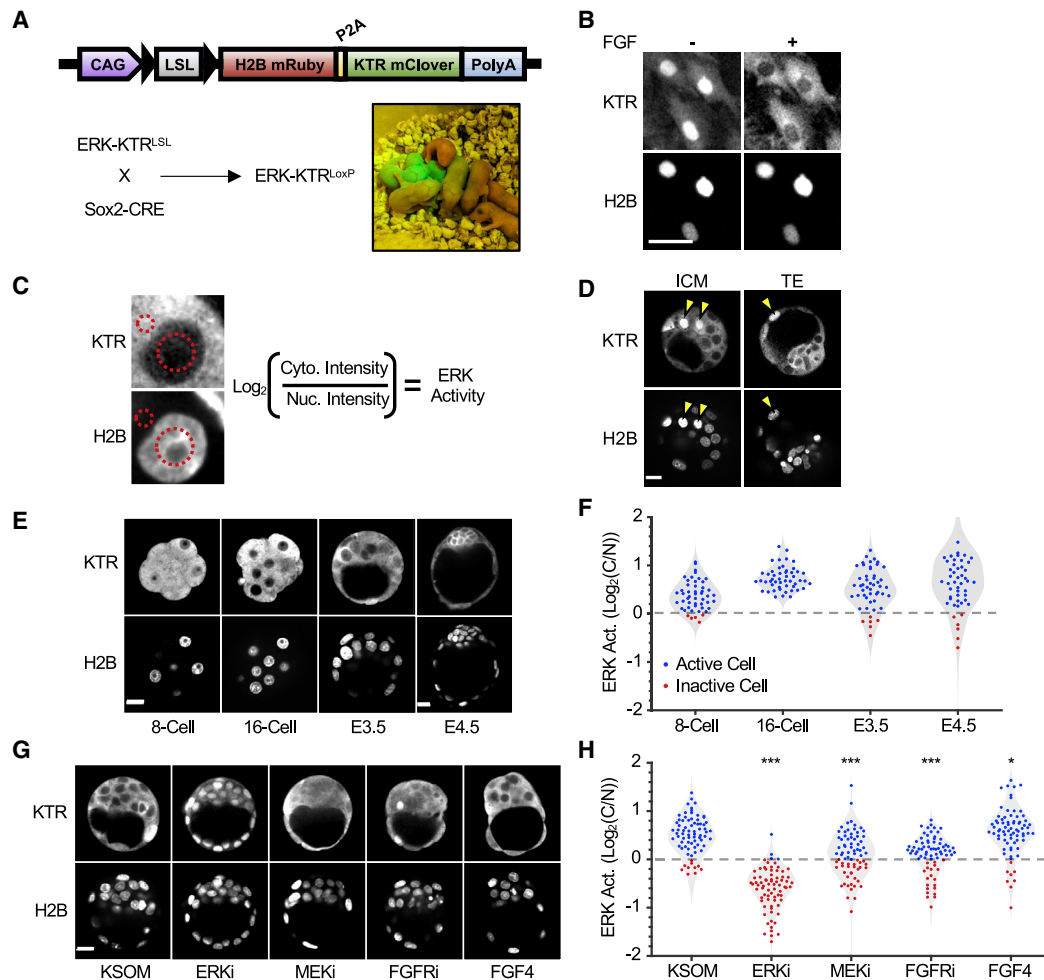
The construct was delivered to zygotes as linearized DNA by pronuclear injection and resulting transgenic integrants were determined by genotyping for the CAG promoter and termed ERK KTR<sup>LSL</sup>. Next, we crossed the founder lines with Sox2-CRE mice to accomplish germline recombination of the transgene and examined offspring using a UV flashlight system at p0–1 (Figure 1A) (Hayashi et al., 2003). In all cases, detection of the CAG promoter by PCR agreed with detectable green fluorescence at birth. These animals will be referred to as ERK KTR<sup>LoxP</sup>. Finally, we determined the precise genomic location of the transgene to be a non-coding region of Chr13 using nanopore Cas9-targeted sequencing (Gilpatrick et al., 2019) and validated it by PCR (Figures S1A and S1B).

To validate the dynamics of the biosensor, we isolated primary mouse embryonic fibroblasts from ERK KTR<sup>LoxP</sup> E13.5 pups and stimulated them with FGF. As expected, FGF stimulation triggered ERK KTR translocation to the cytoplasm (Figure 1B). Next, we wanted to investigate the utility of the KTR mice to study tissue homeostasis in organoids. To this end, we derived 2D gut enteroids from the ERK KTR<sup>LoxP</sup> mice as previously described (Thorne et al., 2018) and measured ERK signaling dynamics upon stimulation with epidermal growth factor (EGF). Live imaging was accompanied by an EdU incorporation assay to allow determination of crypt and villus domains and compare their signaling patterns. While all cells responded to the EGF stimulus, cells in the crypt region rapidly attenuated their ERK activity while villus cells exhibited prolonged ERK activation (Figure S1C). These data show that EGF elicits different ERK signaling dynamics in crypt or villus cells, highlighting the potential of live-cell biosensors to understand tissue homeostasis.

Lastly, we wanted to use the ERK KTR<sup>LoxP</sup> mice to measure MAPK signaling dynamics in single cells of live, unperturbed animals. We performed multiphoton intravital imaging of live mouse epidermis and dermis in ERK KTR<sup>LoxP</sup> mice capturing the basal layer of the skin as well as hair follicles, dermal fibroblast, and other tissue resident cells (Figures S1D and S1E). Our results showed waves of propagating ERK activation in the epidermis similar to those previously described (Hiratsuka et al., 2015) (Figure S1D). Taken together, these experiments support that transgenic KTR mice offer a versatile tool to study MAPK signaling dynamics in diverse *in vitro*, *ex vivo*, and *in vivo* models.

### ERK Signaling Patterns in Preimplantation Development

Although ERK signaling has been implicated in preimplantation development, immunostaining for the phosphorylated species



**Figure 1. ERK KTR<sup>LoxP</sup> Mice Reveal ERK Signaling Dynamics in Mouse Preimplantation Development**

(A) Schematic of KTR reporter construct for ERK KTR<sup>LoxP</sup> animals (upper) and mating scheme to generate mice with germline expression of transgene (lower). Representative image of p0 pups with visible GFP fluorescence is shown.

(B) MEFs were derived from ERK KTR<sup>LoxP</sup> animals and imaged before and after (30 min) stimulation with FGFb (5 ng/mL). Representative images are shown. Scale bar, 50  $\mu$ m.

(C) Schematic of ERK activity quantification method. For each cell, a nuclear and cytoplasmic region of interest was drawn and KTR intensity was measured. ERK activity was reported as the log<sub>2</sub> of the cytoplasmic/nuclear (log<sub>2</sub>(C/N)) intensity ratio.

(D) E3.5 blastocysts from ERK KTR<sup>LoxP</sup> animals were collected, cultured, and imaged as described in STAR Methods. Representative images containing ERK-low cells (arrows) are shown in ICM and TE compartments. Scale bar, 20  $\mu$ m.

(E) Embryos were isolated from ERK KTR<sup>LoxP</sup> animals at the 8-cell, 16-cell, E3.5 blastocyst, and E4.5 peri-implantation blastocyst stages and mounted for imaging in KSOM as described in STAR Methods. Representative images of single confocal planes are shown for clarity. Scale bar, 20  $\mu$ m.

(F) Single-cell ERK activities from embryos collected in (E) were quantified as described in STAR Methods. Cells with predominantly cytoplasmic (log<sub>2</sub>(C/N) > 0) or nuclear (log<sub>2</sub>(C/N) < 0) KTR intensity were represented as blue or red points, respectively. From left to right, N = 8 embryos, 55 cells; 4 embryos, 51 cells; 10 embryos, 339 cells; and 3 embryos, 190 cells.

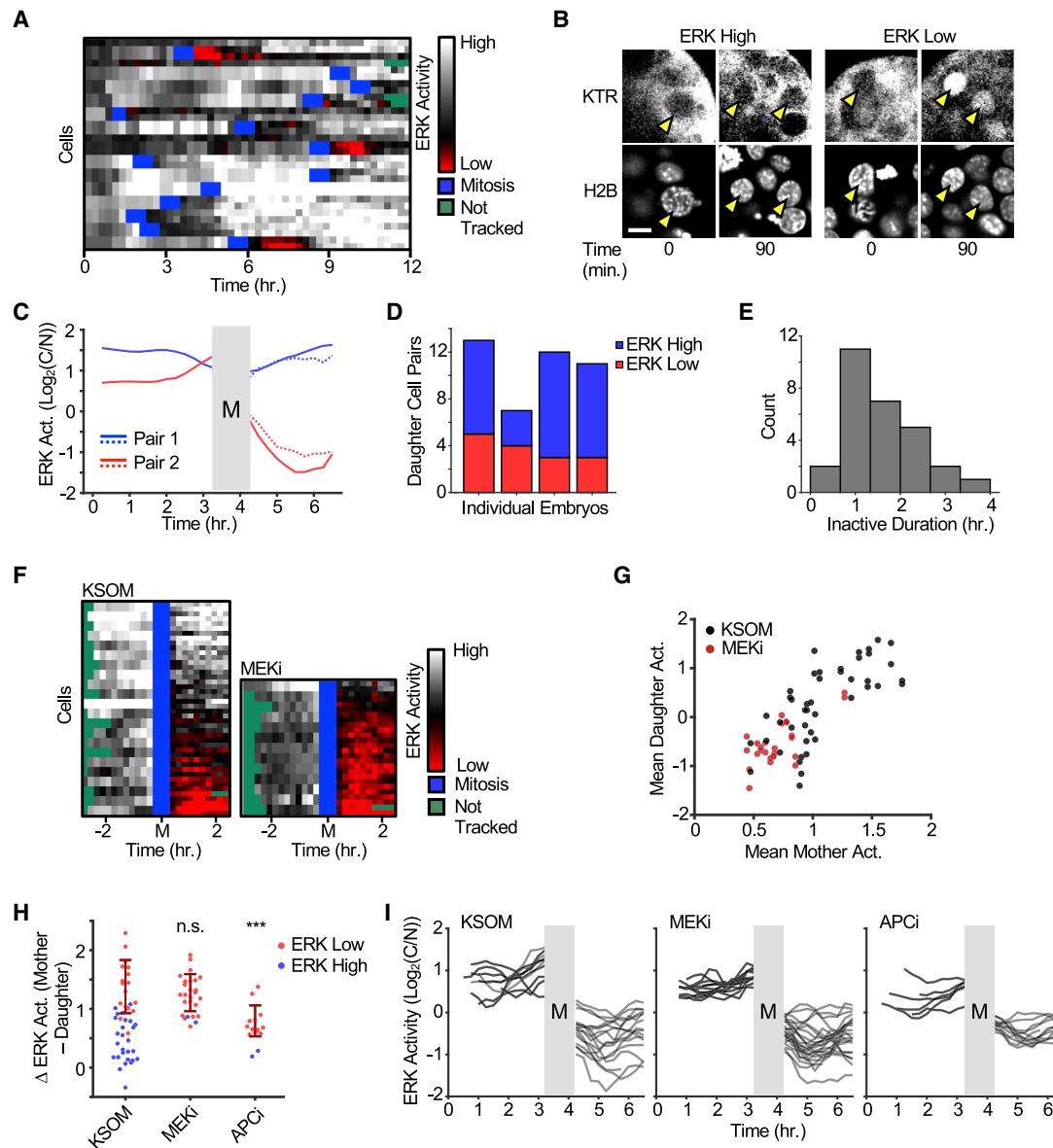
(G) E3.5 blastocysts were isolated from ERK KTR<sup>LoxP</sup> animals and treated with indicated inhibitors and growth factors (ERKi, 5  $\mu$ M Ulixertinib; MEKi, 1  $\mu$ M PD0325901; FGFRI, 1  $\mu$ M AZD4547, 1,000 ng/mL FGF4) for 45 min. Embryos were mounted for imaging in KSOM, maintaining the same concentration of inhibitors and growth factors, and embryos were imaged via confocal microscopy. Representative images of a single confocal plane are shown. Scale bar, 20  $\mu$ m.

(H) Single-cell ERK activities from embryos collected in (G) were quantified and plotted as in (F). From left to right, N = 10 embryos, 339 cells; 5 embryos, 216 cells; 6 embryos, 256 cells; 3 embryos, 103 cells; and 2 embryos, 105 cells. Untreated KSOM control corresponds to E3.5 blastocysts from (E). Significant differences between treated and untreated ERK activity distributions were determined by Kolmogorov-Smirnov test. \*p < 0.05 and \*\*\*p < 0.001.

of these molecules has been technically challenging (Azami et al., 2019; Bessonard et al., 2014). To determine ERK activity in single cells of the mammalian preimplantation embryo, we crossed pairs of ERK KTR<sup>LoxP</sup> mice, isolated embryos from different stages of development ranging from E2.5 to E4.5, and quantified activity in each cell by measuring cytoplasmic over

nuclear KTR ratios (Figures 1C, 1E, and 1F). We refer to ERK active or inactive cells if KTR signals are predominantly cytoplasmic (C > N) or nuclear (N > C), respectively.

Embryos at the 8–16 cell stages showed moderate levels of ERK activity that was sensitive to the ERK specific inhibitor ulixertinib (Figures 1E, 1F, and S2A) (Germann et al., 2017). Later, at



**Figure 2. ERK Activity Bifurcates after Mitosis in Single Cells of Developing Blastocysts**

(A) E3.5 blastocysts were isolated from ERK KTR<sup>LoxP</sup> animals, mounted in KSOM, and imaged every 15 min. for 12 h. Single-cell ERK activity was quantified and represented as a heatmap (see STAR Methods for details). Mitotic events and lost tracks are represented in blue and green, respectively. Predominantly cytoplasmic (log<sub>2</sub>(C/N) > 0) or nuclear (log<sub>2</sub>(C/N) < 0) values are depicted with gray or red scale colormaps, respectively. Black represents C=N intensity. Data depict traces from a single embryo representative of 2 independent experiments.

(B) Representative images of ERK-high and ERK-low mitotic events observed from the experiment in (A). For each example, the mother cell is shown immediately before entering mitosis and the daughter cells are shown 90 min after anaphase. Yellow arrows indicate featured cells. Scale bar, 10  $\mu$ m.

(C) Representative traces of ERK-high (pair 1) and ERK-low (pair 2) mitotic events are plotted. Gray box indicates mitosis. Traces left of mitosis indicate mother cell ERK activity and right of mitosis indicate corresponding daughter cell activities. Each sister cell pair is shown as a solid and dotted line pair.

(D) ERK KTR<sup>LoxP</sup> blastocysts were isolated and imaged as in (A). Daughter cell pairs were assigned as ERK high (C > N, blue) or ERK low (C < N, red). Based on 4 individual embryos.

(E) ERK KTR<sup>LoxP</sup> blastocysts were isolated and imaged as in (A). The duration of ERK inactivity in ERK-low divisions was measured as described in STAR Methods. Histogram reflects measurements from 28 ERK-low cells from 4 embryos obtained from 2 independent experiments.

(F) E3.5 blastocysts were isolated from ERK KTR<sup>LoxP</sup> animals, incubated in KSOM  $\pm$  MEKi (1  $\mu$ M PD0325901), and imaged every 15 min. for 9 h. ERK activity was quantified pre- and post-mitosis as described in STAR Methods. ERK activity is represented as in (A) and *in silico* synchronized to mitosis. Left and right sides of the heatmap indicate mother and daughter cell activities, respectively. Data represent 22 (KSOM) and 13 (MEKi) mitotic events from 4 and 3 embryos, respectively, from 3 independent experiments.

(G) Scatter plot of mean ERK activity 90 min. before (mother) and after (daughter) mitosis from data in (D). Significant linear correlations between mother and daughter cell activity were observed for both the KSOM and MEKi groups ( $p = 2.58 \times 10^{-6}$  and 0.00063 by F-test, respectively) with  $R^2$  values of 0.413 and 0.405, respectively.

(legend continued on next page)



the blastocyst stage, embryos showed predominantly ERK active cells. Interestingly, ERK inactive cells were rare in blastocysts, accounting for 11.8% of all cells at E3.5 and remained similar (12.1%) at E4.5. These inactive cells frequently occurred in pairs in the TE or ICM compartments, but some embryos had none. (Figures 1D and S3A). This overall rarity of ERK-low cells was surprising because we would expect these cells to specify the EPI (Bessonard et al., 2014).

Manipulation of FGF-ERK signaling in the blastocyst with ectopic agonists and inhibitors has been instrumental to study PrE and EPI fate determination (Yamanaka et al., 2010). However, it is unclear how these perturbations alter ERK activity in single cells. To address this question and validate our system, we cultured ERK KTR<sup>LoxP</sup> blastocysts in the presence of inhibitors or growth factors (Figure 1G). MEK and FGFR inhibitors achieve only moderate ERK inhibition compared with the specific ERK1/2 inhibitor ulixertinib, which appears to completely block ERK activity (Figure 1H). Conversely, FGF4 treated embryos showed consistently high ERK activity in most cells comparable to the highest levels of ERK activity detected in control embryos (Figures 1G and 1H). Taken together, these data indicate that KTRs enable quantification of single-cell ERK activity in live pre-implantation embryos.

### Cell-Cycle-Coordinated ERK Signaling Dynamics

Previous reports have shown differential FGF-ERK signaling requirements between the EPI and PrE lineages (Kang et al., 2013; Yamanaka et al., 2010). However, our analysis of embryos at multiple developmental stages showed that only a minority of ICM cells had distinctly lower ERK activity (Figures 1D–1F). Thus, we reasoned that blastomeres may experience transient regulation of the FGFR-ERK signaling axis during fate specification. To test this hypothesis, we performed time-lapse confocal imaging of ERK KTR<sup>LoxP</sup> embryos starting at E3.5 to visualize ERK signaling dynamics in single cells of developing blastocysts (Figures 2A and S2B; Video S1). Tracking and quantification of cells over 12 h confirmed that ICM cells show high levels of ERK activity for most of the imaging period, with some cells exhibiting pulsatile ERK activation and a general trend of increasing ERK signaling as cells approach mitosis. (Figures 2A, S2C, and S2D). Strikingly, ERK activity bifurcated in nascent daughter cells at mitotic exit, generating two groups of cells that either rapidly resume high ERK activity or that experience brief pulses of ERK inactivity (1–4 h) followed by high ERK activation (Figures 2B, 2C, and 2E; Videos S2 and S3). Of note, ERK inactivity pulses were only ever observed immediately after mitosis. Thus, we refer to ERK-high or ERK-low divisions depending on the daughter cell ERK activity. Comparison of multiple embryos revealed variability regarding the fraction of ERK-high versus ERK-low divisions, suggesting that the bifurcation may occur stochastically (Figure 2D).

Previous reports have suggested a limited supply of FGF4 in the blastocyst generates differential signaling to specify the EPI and PrE lineages (Krawchuk et al., 2013). In the case of ERK-low divisions, sister cells showed similar levels of ERK activity, but could be surrounded by neighbors that exhibited much higher ERK activation, suggesting that the symmetry breaking event is occurring in mother cells rather than reflecting local FGF4 concentration gradients (Figures 2C and S2E). Indeed, when cells were sorted according to their mean activity after mitosis, we noticed a correlation between mother and daughter cell ERK activation (Figures 2F and 2G). While all cells display high ERK activation for several hours preceding mitosis, mother cells with comparatively lower KTR ratios were more likely to generate daughter cells that exhibited ERK inactivity pulses (Figures 2F and 2G). This raised the possibility that the maternal signaling status before mitosis may influence signaling at mitotic exit. To test this, we incubated embryos in MEK inhibitor to promote modest ERK inhibition and again analyzed mother and daughter cell activity pre- and post-mitosis. Cells in MEK inhibitor still demonstrated a cytoplasmic enrichment of KTR localization prior to mitosis; however, nearly all daughter cells now experienced clear ERK inactivity after mitosis (Figures 2F and 2G).

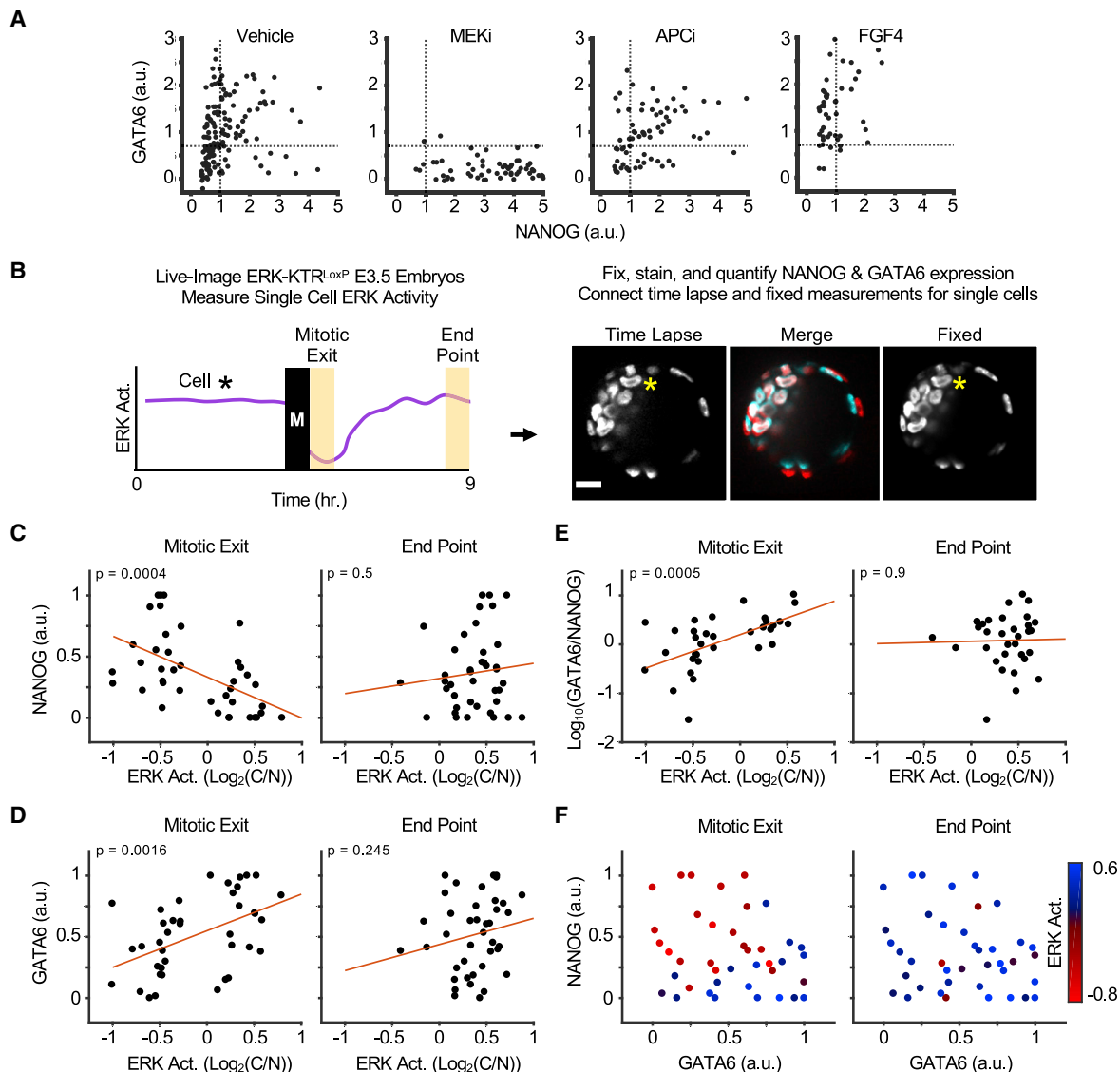
Further analysis revealed that nearly all daughter cells exhibit a decrease in ERK activity that depends on the mother cell activity in a non-linear fashion. We observed that individual mitotic events appear to segregate into two groups based on the magnitude of this decrease, with ERK-low divisions exhibiting a greater differential from maternal activity than ERK-high divisions (Figure 2H). Moreover, the ERK activity decrease in MEK inhibitor treated embryos closely resembles that of the ERK-low daughter cells in control embryos (Figures 2H and 2I). We then hypothesized that the E3 ubiquitin ligase anaphase-promoting complex (APC), which has been shown to regulate both mitosis and signaling (Kimata, 2019; Wan et al., 2017), may influence ERK activity at mitotic exit. We treated embryos with the APC<sup>CDH1</sup> inhibitor ProTAME and analyzed ERK activity changes as cells complete M phase (Zeng et al., 2010). Interestingly, even though APC<sup>CDH1</sup> inhibition led to modest ERK inhibition in mother cells, the relative decrease between mother and daughter cells was significantly reduced compared with ERK-low divisions from control or MEKi embryos (Figures 2H and 2I). Together these data suggest that APC inhibition impairs the regulation of ERK activity at mitotic exit.

### ERK Activity Bifurcation at Mitotic Exit Directs EPI Specification

Understanding the regulatory interactions between the cell cycle, FGF-ERK signaling, and gene expression is necessary to understand the mechanism underlying PrE and EPI establishment. Given that our imaging window was relatively short (9 h), we first tested the effect of inhibitor treatments of the same duration on

(H) E3.5 blastocysts were isolated from ERK KTR<sup>LoxP</sup> animals, incubated in KSOM and KSOM with MEKi (1  $\mu$ M PD0325901) or APCi (2  $\mu$ M ProTAME), and imaged every 15 min for 9 h.  $\Delta$ ERK activity was defined as the mean ERK activity post-mitosis subtracted from the mean ERK activity pre-mitosis. ERK-low (blue) and ERK-high (red) cells were defined as daughter cells whose mean ERK activity was less than 0.1 and greater than or equal to 0.1, respectively. Dark red lines indicate mean  $\pm$  standard deviation of measurements on ERK-low cells only. Data represent 22 mitotic events from 4 blastocysts (KSOM), 13 mitotic events from 3 blastocysts (MEKi), and 7 mitotic events from 2 blastocysts (APCi). \*\*\* $p < 0.001$  and NS  $p > 0.05$  by Student's *t* test comparing ERK-low cells in each group to the KSOM condition.

(I) Data obtained in (H) are shown as *in silico* synchronized ERK activity traces. ERK-low divisions only are shown for clarity. Gray vertical bar indicates mitosis.



**Figure 3. ERK Activity at Mitotic Exit Governs EPI and PrE Lineage Commitment**

(A) E3.5 blastocysts were isolated and cultured in KSOM plus indicated inhibitors or growth factors (MEKi, 1  $\mu$ M PD0325901; APCi, 2  $\mu$ M ProTAME; 1,000 ng/mL FGF4) for 9 h. Embryos were fixed, immunostained, imaged, and quantified as described in STAR Methods. TE cells were excluded from analysis for clarity (see Figures S4A and S4B for details). Data reflect 47 to 153 individual ICM cells from 3 to 8 embryos from 4 independent experiments.

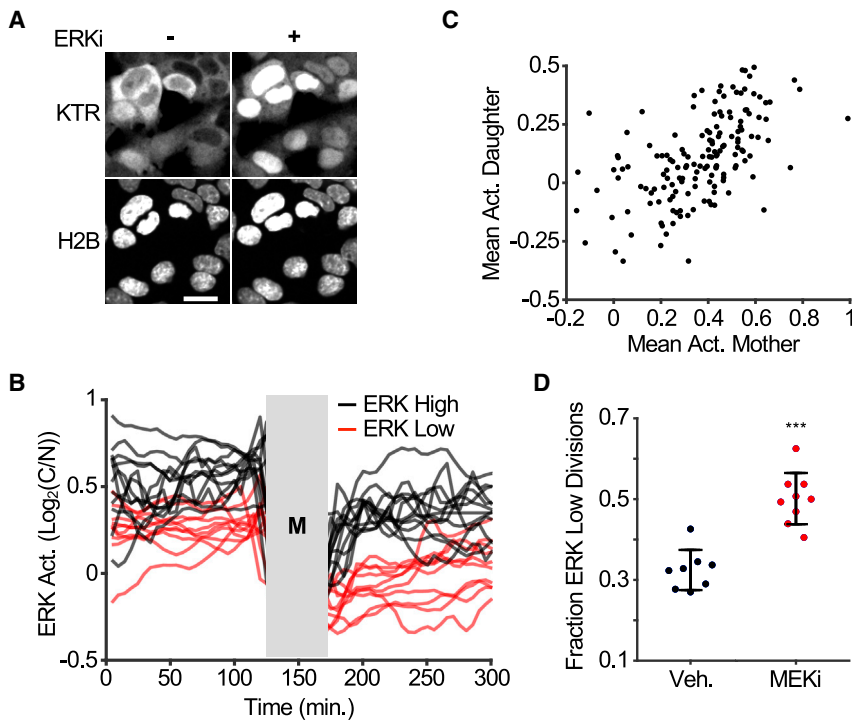
(B) Schematic of experimental workflow to investigate ERK signaling dynamics and fate selection in the same cells. E3.5 ERK KTR<sup>LoxP</sup> blastocysts were imaged in KSOM for 9 h, retrieved, fixed, and immunostained as described in STAR Methods. H2B-mCherry signal from the live imaging and fixed embryos were aligned *in silico* to correlate ERK signaling dynamics and marker expression in the same cells. For each cell that was unequivocally identified in both datasets, ERK activity was measured at mitotic exit (60 min) and at the end of the time-lapse period (end point) (60 min). GATA6 and NANOG signals were then measured and matched with ERK activity measurements for the corresponding cells (see STAR Methods for details). Scale bar, 20  $\mu$ m.

(C–E) ERK KTR<sup>LoxP</sup> blastocysts were imaged and immunostained as described in (B). ERK activity at mitotic exit (left) and end point (right) were plotted against normalized NANOG intensity (C), normalized GATA6 intensity (D), normalized  $\log_{10}(GATA6/NANOG)$  intensity ratio (E). *p* values of linear correlation by F-test are shown. Data represent measurements from 40 mitotic cells from 6 embryos from 3 independent experiments.

(F) Data from (C–E) were plotted as normalized NANOG versus normalized GATA6 with each point colored according to mean ERK activity at mitotic exit (left) or at end point (right).

ICM cell specification by immunostaining for NANOG and GATA6 (Figures 3A and S4C). Following automated measurement of all cells, ICM cells were identified based on their nuclear centroid position within the embryo (Figures S4A and S4B). MEK inhibitor and recombinant FGF4 were still able to bias the expression of NANOG and GATA6 over this interval in the expected patterns

(Figures 3A and S4C). Thus, relationships between signaling and cell fate can be perturbed and examined in this window of early blastocyst development. Interestingly, incubation with the APC inhibitor promoted NANOG/GATA6 double-positive cells relative to the controls, suggesting that APC<sup>CDH1</sup> inhibition impairs lineage commitment in ICM cells (Figures 3A and S4C).



**Figure 4. ERK KTR ES Cells Recapitulate ICM Signaling Dynamics**

(A) ES cells were derived from ERK KTR<sup>LoxP</sup> animals as described in STAR Methods and seeded into imaging plates coated with fibronectin with or without ERKi (5  $\mu$ M ulixertinib). Images were obtained every 5 min. Representative images before or after (15 min) inhibitor addition are shown. Scale bar, 20  $\mu$ m.

(B) ERK KTR ES cells were seeded to imaging plates coated with fibronectin and cultured in complete 2i growth media. Next day, MEK and GSK-3 $\beta$  inhibitors were removed, and cells were imaged every 5 min for 18 h. ERK activity in single cells was measured and quantified as described in STAR Methods. ERK-high and ERK-low cells were defined as cells whose mean ERK activity after mitosis was greater than  $-0.1$  or less than  $-0.1$ , respectively. Following this sorting, randomly selected traces were plotted from each group. Data represent 4 experimental replicates.

(C) Mean ERK activity before (mother) and after (daughter) mitosis (90 min) was calculated from ERK activity traces obtained in (B). Significant linear correlation was observed ( $p = 3.95 \times 10^{-12}$  and  $R^2 = 0.265$ ). Data represent 159 mitotic events.

(D) ERK KTR ES cells were seeded as described in (B) and imaged every 5 min for 20 h in growth media without inhibitors or growth media with MEKi (2  $\mu$ M PD0325901). Fraction of ERK-low divisions was calculated as described in STAR Methods. Data represent  $>700$  mitotic events from  $>8$  replicates for each condition. \*\*\* $p < 0.001$  by a Student's  $t$  test. Error bars represent mean  $\pm$  standard deviation.

To study the lineage commitment in cells coming from ERK-low or ERK-high divisions, we combined time-lapse microscopy of live ERK KTR<sup>LoxP</sup> embryos with fixed, endpoint analysis of cell fate markers. Embryos were imaged for 9 h, fixed, immunostained, and re-mounted onto new imaging plates to visualize protein expression (Figure 3B; Video S4). Embryos were only used for analysis if exceptional *in silico* alignment of H2B-mCherry nuclear signal was observed (Figure S4D). Thus, embryos that were deformed from the fixation or staining protocols, and embryos that were re-mounted in an orientation that precluded the unambiguous identification of matching cells were discarded. To maximize the likelihood of generating image stacks that could be aligned *in silico*, fixed embryos were gently reoriented on the imaging plate by pipetting and re-imaged (no more than three times).

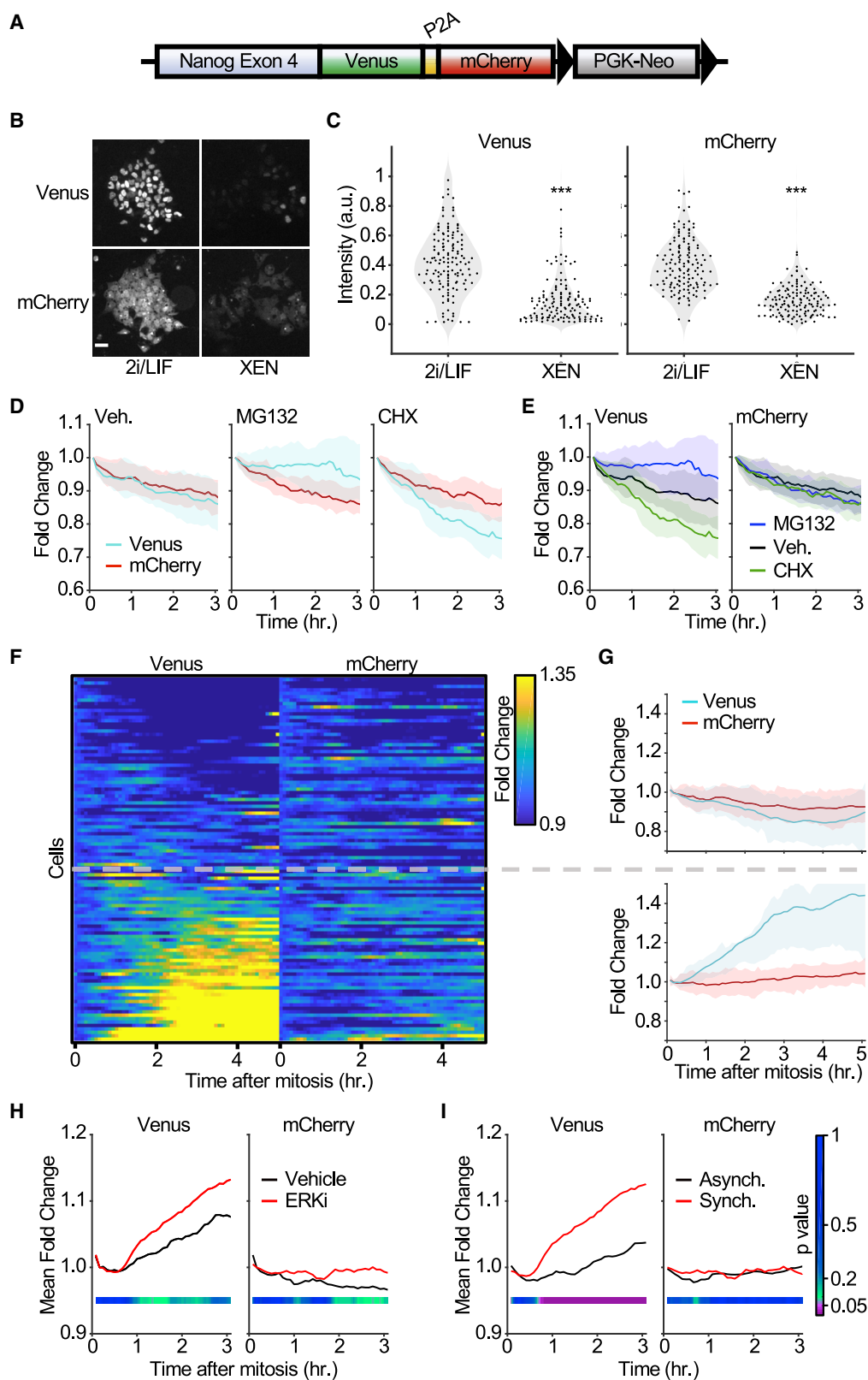
Because we observe the greatest heterogeneity in ERK activation at mitotic exit, we focused our analysis on cells that underwent mitosis during the time-lapse microscopy period. In particular, we measured mean ERK activity in nascent daughter cells in the hour immediately after mitosis and in the final hour of the time-lapse and looked for relationships between ERK signaling and protein expression of NANOG and GATA6 (Figure 3B). Cells that inhibit ERK at mitotic exit expressed elevated levels of NANOG and moderately lower levels of GATA6. Accordingly, cells that resume high ERK activity after mitosis displayed enhanced GATA6 and diminished NANOG expression (Figures 3C and 3D). While levels of each of these factors were independently correlated with ERK activity ( $r^2 = 0.233$  and

$r^2 = 0.285$  for ERK versus GATA6 and ERK versus NANOG, respectively), we found that analyzing the relationship between ERK and GATA6 over NANOG ratios resulted in a superior correlation than either marker alone ( $r^2 = 0.332$ ) (Figure 3E). 3D scatter plots with NANOG, GATA6, and ERK activity at mitotic exit showed an ERK sensitivity landscape for cells selecting between the EPI and PrE lineages (Figure 3F). Importantly, analyzing ERK activity of an equivalent duration, but in the final hour of the time course, and examining the relationships with these fate markers indicated no significant correlation between any of the variables described above (Figures 3C–3F). Together, these data suggest that ERK activity bifurcation at mitotic exit directs the GATA6/NANOG expression ratio in single cells.

### Modeling ICM Signaling Behavior in ES Cells

While the blastocyst is a powerful model to explore how signaling dynamics regulates cell fate specification in development, it is limited by its low phototoxicity tolerance and lack of tools for robust tracking of cells in 3D time-lapse datasets. In contrast, ES cells are accessible to rapid imaging and automated cell tracking while closely resembling the EPI of the preimplantation embryos (Fernández-de-Manuel et al., 2017; Boroviak et al., 2015; Schröter et al., 2015; Nichols and Smith, 2012). To further explore the mechanism underlying ERK signaling at mitotic exit, we derived ES cells from ERK KTR<sup>LoxP</sup> animals and validated the sensor using ERK inhibitor (Figure 4A). Time-lapse microscopy of ERK KTR ES cells revealed





(legend on next page)

a bifurcation of ERK activity at mitotic exit, sustained activity before mitosis, and a correlation between mother and daughter cell activity that recapitulate our observations in ICM cells (Figures 4B, 4C, and S5A). Incubation with MEK inhibitor again increased the frequency of ERK-low divisions (Figure 4D; Videos S6 and S7). Importantly, ERK inhibitor abolished cytoplasmic KTR localization for all but 5 min prior to mitosis, indicating that prolonged ERK activation before mitosis is not an artifact of nonspecific nuclear export before breakdown of the nuclear envelope (Video S5).

The role of ERK signaling in NANOG transcriptional and post-translational regulation has been previously investigated (Jin et al., 2016; Kim et al., 2014; Ochiai et al., 2014); however, the regulation of NANOG dynamics at mitotic exit and its dependence on ERK activity remain unexplored. In order to independently monitor NANOG protein stability and *Nanog* gene expression, we generated an ES cell line with a targeted genomic insertion at the *Nanog* locus of a Venus fluorescent protein followed by the self-cleaving peptide P2A and mCherry and refer to this line as NV2C (Nanog-Venus-2a-mCherry) cells (Figure 5A). Thus, while the Venus signal is linked to NANOG protein levels, the mCherry signal is independent of NANOG stability and can serve as a proxy for transcriptional activity. To validate our reporter line we cultured NV2C cells in pluripotency maintenance (2i/LIF) or extraembryonic endoderm (XEN) differentiation media for 24 h and measured Venus and mCherry levels in single cells (Niakan et al., 2013). As expected, both the NANOG-Venus and mCherry signals were significantly downregulated in XEN differentiation medium compared with 2i/LIF, reflecting attenuated *Nanog* expression (Figures 5B and 5C). Inhibition of translation or proteasomal degradation showed downregulation or upregulation of NANOG-Venus levels respectively while the mCherry signal remained stable (Figures 5D and 5E). These results indicate that NV2C cells independently report NANOG protein stability and gene expression.

Next, we live imaged NV2C cells and monitored NANOG-Venus and mCherry dynamics at mitotic exit. *In silico* synchronization of individual cells to mitotic exit revealed Venus levels were rapidly upregulated in a subset of cells after mitosis without a corresponding increase in mCherry intensity (Figures 5F and 5G; Video S8). This suggests heterogeneous post-translational regulation of NANOG at mitotic exit. Therefore, given the differential ERK activation we observed in ES cells and in blastocysts, we asked how ERK signaling regulates NANOG dynamics at mitotic exit. We incubated NV2C cells in ERK inhibitor to induce homogeneous ERK inhibition, *in silico* synchronized cells to mitotic exit, and analyzed Venus and mCherry dynamics (Figure 5H). Cells in ERK inhibitor exhibited enhanced Venus expression at mitotic exit while mCherry levels were relatively stable. Moreover, a comparison of synchronized and unsynchronized cells showed that, while ERK inhibition promotes NANOG-Venus expression in all cells, the effect of ERK inhibition is greatest at mitotic exit (Figure 5I). Taken together these data provide evidence that NANOG protein stability is regulated at mitotic exit and is enhanced by ERK inhibition.

## DISCUSSION

Development of multicellular organisms relies on cell fate choices that occur as a result of constant cross regulation between signaling and gene expression. The dynamic and noisy nature of these biological processes require approaches that enable single-cell high temporal resolution measurements in real time. Here, we describe the generation and validation of a mouse line that enables *in vivo* and *ex vivo* analysis of ERK signaling dynamics with single-cell resolution. Results show that this tool can be used in primary cells, multicellular organoids, or even live animals to interrogate the role of signaling dynamics and cell-cell communication in tissue homeostasis and development. Our ERK KTR mice are largely

### Figure 5. ERK Inhibition Promotes Rapid NANOG Stabilization at Mitotic Exit

- (A) Schematic representation of the NANOG NV2C reporter inserted at the endogenous *Nanog* locus in mouse ES cells (see STAR Methods for details).
- (B) NV2C ES cells were seeded to imaging plates and cultured in 2i/LIF media or XEN differentiation media for 24 h before imaging. Representative colonies from 16 replicates are shown. Scale bar, 30  $\mu$ m.
- (C) Quantification of Venus and mCherry intensities from single cells obtained in (B).  $n > 1,000$  cells for each condition from 16 replicates. \*\*\* $p < 0.001$  by Kolmogorov-Smirnov tests.
- (D) NV2C ES cells were seeded onto imaging plates and cultured in ES growth media without inhibitors. Cells were imaged every 5 min for 3 h and MG132 (10  $\mu$ M) or cycloheximide (CHX, 20  $\mu$ g/mL) were added after 35 min of imaging. Single-cell Venus and mCherry intensities were quantified and analyzed as described in STAR Methods. Mean fold change (solid lines) with 75<sup>th</sup> and 25<sup>th</sup> percentiles (shaded area) are plotted for 108, 73, and 85 cells from 4 replicates each for the vehicle, MG132, and CHX groups, respectively.
- (E) Data from (D) plotted as mean fold change (solid lines) and 75<sup>th</sup> and 25<sup>th</sup> percentiles (shaded area) of Venus (left) and mCherry (right) intensities from each condition.
- (F) NV2C ES cells treated as in (D) were imaged every 5 min for 20 h. Venus and mCherry intensities were quantified as described in STAR Methods and *in silico* synchronized to mitotic exit. Data are plotted as fold change in intensity for the first 10 min after anaphase. Overlaid gray dashed line indicates cells that up-regulate NANOG at mitotic exit (lower) and cells that maintain or downregulate NANOG expression (upper). Data represent 122 mitotic events from 6 experimental replicates.
- (G) Average traces are plotted for data in (F). Plots show mean fold change (solid lines) and 75<sup>th</sup> and 25<sup>th</sup> percentiles (shaded area) of Venus and mCherry intensities for cells that maintain or downregulate NANOG (upper) and cells that upregulate NANOG (lower).
- (H) NV2C ES cells were seeded as in (E) and treated with or without ERKi (5  $\mu$ M Ulixertinib). Single-cell traces of Venus and mCherry signals were quantified as in (F). Mitotic events were synchronized *in silico* and mean fold change of Venus and mCherry intensities for all cells analyzed are plotted. Data represent >122 cells from 6 replicates each. Student's *t* test was performed for means at each time point and *p* values are plotted according to color bar in Figure 5I.
- (I) ERKi (5  $\mu$ M ulixertinib)-treated cells obtained in (H) are plotted to compare average Venus and mCherry fold changes in asynchronous (black) and mitotic exit *in silico* synchronized (red) individual cells. Data represent >140 individual cells from 6 replicates for each condition. Student's *t* test was performed for means at each time point, and *p* values are plotted according to color bar.

equivalent to the previously reported Eisuke mice (EKAREV FRET sensor) (Hiratsuka et al., 2015); however, studies that require tissue specific expression of the biosensor will benefit from the CRE-dependent design of the ERK KTR line.

ERK signaling has been implicated in specifying the mammalian preimplantation embryo; however, the signaling dynamics leading to cell fate choices had not been explored. Our data show that ERK activity is surprisingly high throughout all stages of pre-implantation development, with a small subset of cells experiencing transient ERK inhibition in the blastocyst stage. Interestingly, ERK activity at the blastocyst stage can be completely blocked with ERK specific inhibitors but other conditions previously used to bias cell fate specification (MEK or FGFR inhibitors) only partially inhibit ERK activity, indicating that even minor perturbation of FGF-ERK signaling axis is sufficient to prevent proper lineage specification in the ICM (Figure 1H). While we observed increased ERK activation upon FGF4 stimulation, many FGF4-treated blastomeres exhibited comparable ERK activity to cells in control embryos, suggesting that FGF4 is locally present in saturating amounts to achieve maximal FGFR signaling output to ERK. A recent study reported pERK staining in nearly all cells of blastocysts and differential pERK staining within the ICM, in agreement with our measurements (Azami et al., 2019).

Further analysis of ERK activity dynamics in single cells of the blastocyst revealed that an apparently stochastic number of cell divisions result in 1–4 h pulses of ERK inactivity that then transitions back into high levels of ERK activation. Sister cells exhibited similar signaling patterns that could be quite different from non-sister neighbor cells, resembling previously reported ERK signaling dynamics in ES cells (Deathridge J et al. Development 2019) (Figure S5A). Interestingly, cells that experienced ERK inactivity pulses at mitotic exit exhibited higher NANOG and lower GATA6 expression (Figure 3F). This finding is in agreement with several studies that suggest low ERK activity in ICM cells specifies the EPI lineage, but to our knowledge no previous report postulates a model incorporating transient inactivation of ERK. This correlation suggests a coordination between ERK signaling and the cell cycle that is important for cell fate determination; however, studies utilizing live-cell biosensors for NANOG and GATA6 alongside kinase activity biosensors in embryos are necessary to understand the temporal relationships between mitotic exit signaling and ICM progenitor specification.

The amplitude of the ERK activity decrease at mitotic exit correlated with maternal signaling before mitosis and was dampened by APC<sup>CDH1</sup> inhibition. An increasing body of evidence supports cell-cycle independent, APC-mediated function in cell fate determination through the degradation of developmental signaling pathway components, including the ERK pathway via downregulation of Raf (Kimata, 2019). Moreover, ERK itself can downregulate APC activity constituting a positive feedback loop that could explain the dependency on mother cell activity (Wan et al., 2017). We report that APC inhibition led to an increase in uncommitted ICM progenitors. This could be a result of the conflicting roles of this treatment: APC inhibition increased the frequency of ERK-low divisions, which may limit GATA6 expression, but also attenuated the amplitude of the inhibition to promote only modest NANOG expression. Additional work is required to understand the regulatory interactions between

APC activity, ERK signaling, and fate determination in embryos. Though the molecular mechanism behind the ERK activity bifurcation remains to be characterized, it is tempting to speculate that feedback regulation between the mitotic machinery and ERK pathway components may desensitize pairs of daughter cells to FGF signals and ultimately specify the EPI.

Characterization of ES cells derived from our ERK KTR mouse line revealed that ERK activity bifurcation at mitotic exit also occurs in ES cells and depends on maternal signaling levels. Moreover, we found that NANOG protein stability at mitotic exit is particularly sensitive to ERK inhibition. A model where NANOG levels are sustained via protein stabilization rather than transcriptional regulation may explain why inhibiting transcription in blastocysts with flavopiridol does not promote rapid loss of NANOG expression (Bessonard et al., 2017). These findings provide a potential explanation for the significance of ERK activity at mitotic exit to specify the EPI and PrE lineages via modulation of NANOG levels and are in good agreement with recent reports suggesting that cell-cycle status plays a critical role in regulating pluripotent cell differentiation. Human and mouse embryonic stem cells (ESCs) are both more responsive to differentiation cues during G1 compared with S and G2, suggesting a mechanism for coordination of the cell-cycle program with the signaling network (Singh et al., 2015; Pauklin and Vallier, 2013). As ERK has been reported to regulate several pluripotency factors, further experimentation is required for a complete understanding of the consequences of ERK signaling at mitotic exit (Ma et al., 2016). We anticipate that a better understanding of the cross regulation between cell cycle, signaling, and differentiation will ultimately facilitate the use of ES and iPS cells for therapeutic purposes.

### Limitations

The findings of this study possess some notable limitations. The experiments including the APC<sup>CDH1</sup> inhibitor, ProTAME, showing an effect on both ERK signaling at mitotic exit and ICM cell fate commitment are not sufficient to show a direct regulatory mechanism between ERK and APC. Though there is literature to support that APC could directly regulate ERK and vice versa, we acknowledge that a pharmacological inhibitor of this essential regulator of mitosis could regulate both ERK activity and ICM differentiation via a number of indirect mechanisms. A more detailed analysis with additional cell-cycle component inhibitors would help identify whether and how this regulatory control functions.

The connections between ERK activity, cell cycle, and NANOG/GATA6 protein levels in embryos and ES cells are still unclear. We present correlative evidence that: (1) ERK activity is uniquely heterogeneous immediately following mitosis, (2) mitotic exit signaling correlates with NANOG and GATA6 expression, (3) NANOG protein is stabilized in a subset of ES cells after mitosis, and (4) ERK inhibitors promote NANOG stabilization specially after mitosis. However, due to spectral overlap between sensors and other current restrictions, we were unable to directly measure NANOG and GATA6 with the ERK KTR in live embryos or ES cells. Thus, we cannot conclusively claim that ERK signaling at mitotic exit is the cause of (1) cells selecting between EPI and PrE fates or (2) the rapid NANOG stabilization in

ES cells. The mechanistic links between cell-cycle status, ERK signaling, and NANOG dynamics still remain an open question.

## 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.devcel.2020.09.013>.

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## AUTHOR CONTRIBUTIONS

M.P. and S.R. conceived the study, analyzed the data, and wrote the manuscript. M.P. and K.R. performed all experiments except nanopore sequencing and intravital imaging. B.P. and W.T. performed nanopore sequence. T.X. and V.G. performed intravital imaging of ERK KTR mice. S.R. supervised the study and secured funding.

## DECLARATION OF INTERESTS

The authors declare no competing interests.

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# STAR★METHODS

## KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
<b>Antibodies</b>		
Rabbit monoclonal anti-NANOG	Cell Signaling Technology	Cat# 8822; RRID:AB_11217637
Goat polyclonal anti-GATA6	R&D Systems	Cat# AF1700; RRID:AB_2108901
Mouse monoclonal anti-CDX2	BioGenex	Cat# MU392; RRID:AB_2335627
Donkey anti-rabbit IgG (Alexa Fluor 405)	Abcam	Cat# ab175649; RRID:AB_2715515
Donkey anti-mouse IgG (Alexa Fluor 488)	Molecular Probes	Cat# A21202; RRID:AB_141607
Donkey anti-Goat IgG (Alexa Fluor 647)	Molecular Probes	Cat# A21447; RRID:AB_141844
<b>Chemicals/Peptides/Recombinant Proteins</b>		
CHIR99021	Sigma-Aldrich	Cat# SML1046
PD0325901	Selleckchem	Cat# S1036
Ulixertinib	Selleckchem	Cat# S7854
AZD4547	Selleckchem	Cat# S2801
proTAME	R&D Systems	Cat# I-440-01M
Y-27632	Selleckchem	Cat# S1048
LDN-193189	Selleckchem	Cat# S2618
Retinoic Acid	Sigma-Aldrich	Cat# R2625
MG132	Selleckchem	Cat# S2619
Cycloheximide	Sigma-Aldrich	Cat# C4859
Mitomycin C	Sigma-Aldrich	Cat# M4287
ESGRO	Milipore Sigma	Cat# ESG1106
rhFGF4	Thermo Fisher Scientific	Cat# PHG0154
EGF	PeproTech	Cat# AF-100-15
FGFb	Sigma-Aldrich	Cat# GF003
Activin A	R&D Systems	Cat# 338-AC-010
Heparin	Sigma-Aldrich	Cat# H3149
N-acetyl-L-cysteine	Sigma-Aldrich	Cat# A7250
Mouse R-Spondin-1	Thermo Fisher Scientific	Cat# 3474RS050
Matrigel Matrix Growth Factor Reduced	Corning	Cat# 354230
5 ethynyl 2 deoxyuridine (EdU)	Thermo Fisher Scientific	Cat# A10044
Human Plasma Fibronectin	Milipore Sigma	Cat# FC010
EmbryoMax Gelatin	Milipore Sigma	Cat# ES-006-B
2-Mercaptoethanol	Thermo Fisher Scientific	Cat# 21985023
Dithiothreitol	Thermo Fisher Scientific	Cat# R0861
GlutaMAX	Thermo Fisher Scientific	Cat# 35050061
<b>Critical Commercial Assay</b>		
Nanobind Tissue Big DNA Kit	Circulomics	Cat# NB-900-701-01
Short Read Eliminator	Circulomics	Cat# SS-100-101-01
Oxford Nanopore Sequencing 1D	Nanoporetech	Cat# SQK-LSK109
<b>Experimental Models: Organisms/Strains</b>		
Mouse: ERK KTR-LSL	This Paper	JAX#035566
Mouse: ERK KTR-LoxP	This Paper	N/A
<b>Oligonucleotides</b>		
R Chkactin: 5'-GGGGGAGATGGGGAGAGTGAAGCAGAA-3'	This paper	N/A
F CMV: 5'-GGGTGGAGTATTTACGGTAACTGCCC-3'	This paper	N/A
oMP106 5'-CTCCAGCATGACTAGAGTCAGGCTCTGG-3'	This paper	N/A

(Continued on next page)

### Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
oMP109 5'-CATTAAATGGTTTCAGGTGGATGTCTCCTCCC-3'	This paper	N/A
oMP115 5'-GGGTAGATGAGGGATTAGTTGAAATTCATCATGAGG-3'	This paper	N/A
Recombinant DNA		
pSpCas9(NANOG)-2A-GFP	This Paper	N/A
pSpCas9(BB)-2A-GFP	<a href="#">Ran, et al., 2013</a>	Addgene Repository #48138
Nanog-2a-mCherry	<a href="#">Faddah, et al., 2013</a>	Addgene Repository #59995
pMP108 Nanog-Venus-2a-mCherry	This Paper	N/A
pOO16 Step2 Rosa26 Vector ERK	This Paper	N/A
Software and Algorithms		
MATLAB	mathworks	<a href="http://www.mathworks.com">www.mathworks.com</a>
Samtools	<a href="#">Li et al., 2009</a>	<a href="http://www.htslib.org/download">www.htslib.org/download</a>
TE and ICM cell ID	This Paper	<a href="https://github.com/MichaelPokrass/EmbryoScripts">https://github.com/MichaelPokrass/EmbryoScripts</a>
Deposited Data		
Integrated time-lapse and end-point datasets of ERK KTR Embryos	This Paper	Biolmage Archive: <a href="https://www.ebi.ac.uk/biostudies/studies/S-BIAD28">https://www.ebi.ac.uk/biostudies/studies/S-BIAD28</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, Sergi Regot ([sregot@jhmi.edu](mailto:sregot@jhmi.edu)).

### Material Availability

The ERK KTR<sup>LSL</sup> mouse line is available from The Jackson Laboratory as JAX#035566. The Nanog targeting vector is available from Addgene.

### Data and Code Availability

Live imaging of embryo and corresponding fixed embryo data sets is available at the Biolmage Archive (<https://www.ebi.ac.uk/biostudies/studies/S-BIAD28>). MATLAB code generated to assign ICM versus TE cell identity on basis of cell position is available from: <https://github.com/MichaelPokrass/EmbryoScripts>.

## EXPERIMENTAL MODEL AND SUBJECT DETAILS

### Animals

All mice were housed, handled, and included in experiments according to the approved Institutional Animal Care and Use Committee protocol M019M487 of the Johns Hopkins University School of Medicine. The ERK KTR<sup>LSL</sup> line was generated by injection into embryos from mice of B6SJL background and bred into the CD1 background. To generate the ERK KTR<sup>LoxP</sup> mice, male ERK KTR<sup>LSL</sup> mice were crossed with female Sox2-CRE mice and GFP+ offspring were bred into the CD1 background. Wild-type embryos were collected from CD1 background mice.

### Embryos

Mouse preimplantation embryos were cultured in EmbryoMax Advanced potassium-supplemented simplex optimized medium (KSOM) (Milipore Sigma) under mineral oil (Sigma Aldrich) on polystyrene plates. Embryos were maintained in a humidified incubator at 37°C with 5% CO<sub>2</sub> and 3% O<sub>2</sub>.

### Primary Cell Culture

Primary mouse embryonic fibroblasts (MEFs) were maintained in MEF culture medium (DMEM with 15% fetal bovine serum (FBS), 100 U/ml penicillin and 100 U/ml streptomycin) and cultured on polystyrene plates at 37°C with 5% CO<sub>2</sub> and 3% O<sub>2</sub>. Primary mouse 2D gut enteroid cultures were maintained in organoid culture medium (advanced DMEM-F-12 medium with 100 U/ml penicillin and streptomycin, 10 mM HEPES buffer, 1x Glutamax, 1X B27, 1 mM N-acetyl-cysteine, 100 µg/ml primocin, 50 ng/ml EGF, 10% Noggin-conditioned medium, and 20% R-spondin conditioned medium) and cultured on

Matrigel-coated optical polymer plates at 37°C with 5% CO<sub>2</sub>. Primary mouse ES cells were maintained in either ES culture medium (advanced DMEM-F-12 medium with 100 U/ml penicillin and streptomycin, 15% ES-qualified FBS, 1x GlutaMAX, 1X MEM non-essential amino acids (Thermo), 55 μM 2-mercaptoethanol (BME, Gibco), and 1000 U/ml leukemia inhibitor factor (LIF) (ESGRO, Milipore Sigma)) on Mitomycin C-inactivated feeder MEFs or feeder-free with ES culture medium supplemented with 1 μM PD0325901 and 3 μM CHIR99021 (2i) on 0.1% gelatin-coated (Milipore Sigma) polystyrene plates at 37°C with 5% CO<sub>2</sub> and 3% O<sub>2</sub>.

## METHOD DETAILS

### Mouse Generation

The ERK KTR reporter construct was cloned into a Rosa26 targeting vector (gift from Dr. Jeremy Nathans). Cas9 protein, gRNA, and purified linear DNA insert was introduced to B6SJL embryos by pronuclear injection and transferred to pseudopregnant ICR females. Litters were screened for integrants by PCR amplification of the CAG promoter with F CMV and R Chkactin oligos. CAG+ males were crossed with Sox2-CRE females (gift from Dr. Jeremy Nathans) to generate animals with germline expression of reporters. Offspring were screened for visible GFP expression by blue light flashlight (Nightsea). Founder lines were bred into CD1 background.

### Targeted Nanopore Sequencing

Genomic DNA extracted from mouse tail using Nanobind Tissue Big DNA kit (Circulomics #NB-900-701-01). Mouse tail was incubated with proteinase K and Buffer CLE3 for 90 min on a ThermoMixer at 55 degrees C and 900 rpm. DNA was incubated in elution buffer overnight. Size selection was performed using Short Read Eliminator (Circulomics #SS-100-101-01) followed by a DNA clean-up using 0.9 AMPure XP (Beckman Coulter, A63881) beads. DNA quantification was done using the Qubit fluorometer (Thermo) before sequencing. Targeted Nanopore Sequencing with Cas9 was performed as previously described (Gilpatrick et al. 2020). Library prep was performed using Oxford Nanopore Sequencing 1D sequencing kit (SQK-LSK109). Each sample was sequenced using a 9.4.1 version flow cell using the MinION sequencer. Basecalling was performed using the GUPPY algorithm (Version 3.2.1) to generate FASTQ reads from the electrical data. Reads were aligned to a synthetic reference created by fusing the mouse reference genome (mm10) and the known insert sequence using Minimap2 (Li, 2018). Analysis of the insert coverage was performed using Samtools (Li, et al., 2009) and insert coverage was visualized using IGV and Ribbon. Insertion sites were verified by PCR amplification of sequence spanning junction of reference genome and insert sequence with multiplexed reaction including oMP106, oMP109, and oMP115 oligos.

### MEF Derivation and Culture

Primary MEFs were derived from ERK KTR<sup>LoxP</sup> and wild type mice as previously described (Xu, 2005). Following timed matings, embryos were dissected from mice at E13.5. Embryos were washed in ice-cold, sterile PBS and, using forceps, the brain, liver, and heart were manually removed and discarded. Embryos were cut into fine pieces and transferred to ice-cold 0.25% trypsin-EDTA and incubated at 4°C overnight. Next, the embryos were incubated at 37°C for 30 min and the tissue was vigorously pipetted in MEF culture medium. When tissue was separated into a uniform cell suspension, the cells were plated onto tissue culture flasks in MEF medium. For generation of feeder cells for ES experiments, MEFs were expanded to passage 3 and inactivated by 2.5 hr. incubation with 10 μg/mL mitomycin C (Milipore Sigma). Adherent cells were expanded and cultured in MEF culture medium. For imaging experiments, 7 x 10<sup>5</sup> MEFs were seeded to fibronectin (EMD-Milipore) coated 96-well glass-bottom plates. MEFs were serum starved 16 h before stimulus addition.

### Gut Enteroid Culture

Primary 2D gut enteroids were derived as previously described (Thorne, et al., 2018). Briefly, mouse jejunae were dissected from 1-6 month old mice. Jejunae were cut open lengthwise, washed with PBS, and incubated 30 min in ice-cold PBS containing 100 U/ml penicillin and streptomycin, 1.5 mM DTT, 2 mM EDTA, and 10 μM Y-27632. The jejunae were transferred to ice-cold PBS with 2 mM EDTA and shaken for 90-120 seconds to release crypts. Intestinal tissue was discarded and the crypt suspension was washed three times in DMEM with 10% FBS by centrifugation at 300 x g for 3 min. The crypts were resuspended in DMEM with 10% FBS and filtered through a 500 μm strainer followed by a 70 μm strainer. The crypts were pelleted at 200 x g for 3 min and resuspended in attachment media, which consisted of basal organoid media (advanced DMEM-F-12 medium with 100 U/ml penicillin and streptomycin, 10 mM HEPES buffer, and 1x Glutamax) supplemented with 1 mM N-acetyl-cysteine, 1x B-27 supplement, 50 ng/ml EGF, 100 nM LDN-193189, 1 μg/ml R-spondin 1, 10 μM CHIR99021, and 10 μM Y-27632 and seeded to optical polymer 96-well plates coated with 0.8 mg/ml growth factor reduced Matrigel diluted in basal organoid media. Crypts were incubated in attachment media overnight before media was exchanged for organoid culture medium. Enteroids received fresh organoid culture medium every 24 h and were cultured up to one week.

Prior to EGF addition experiments, organoid culture medium was exchanged for equivalent medium without EGF overnight. Labeling of S-phase cells was performed alongside EGF addition. EdU was added to media to a final concentration of 10 μM for a 3-hour incubation followed by fixation in ice-cold methanol. Cells were washed and labeled via click chemistry with Fluor Azide 488 (Thermo Fischer Scientific). Regions of EdU positive cells were used to designate crypt-like regions and EdU negative cells defined the villus-like regions (Thorne, et al., 2018).

### Imaging

Images were acquired with a Metamorph-controlled Nikon Eclipse Ti microscope. For epifluorescence imaging, data were acquired using 10X and 20X air objectives and a Hamamatsu sCMOS camera. For confocal imaging, data were acquired using 20X air, 40X oil, or 60X oil objectives, a Yokogawa CSU-W1 spinning disk confocal unit, and a Photometric Prime 95B sCMOS camera. Live specimens were maintained at 37°C in 5% CO<sub>2</sub> in a humidified environmental control chamber from OKO Labs.

### Intravital Imaging

Multiphoton microscopy was performed as previously described (Pineda, et al., 2015). Mice were maintained on anesthesia with nose cone supply of vaporized isoflurane and placed on a warming pad. Skin was mounted on a custom-stage and covered by a glass coverslip. A LaVision TriM Scope II (LaVision Biotec) microscope equipped with a Chameleon Vision II (Coherent) 2-photon laser (using 940nm) and a Chameleon Discovery (Coherent) two-photon laser (using 1120nm) was used to acquire images through a Nikon 40x/1.15 water immersion objective at an interval of every 6.25 min for 3.5 h.

### Preimplantation Embryo Culture

Embryos were collected at indicated developmental stages following timed matings of mice maintained in a 12-hour light-dark cycle according to previously described protocols (Behringer, 2014). For live imaging experiments, embryos were transferred to glass-bottom microwell dishes (MatTek) and maintained in an environmental control chamber (OKO Lab) and imaged every 15 min for 9–12 h, sampling a depth 60 μm at 5 μm steps. For inhibitor incubations, embryos were transferred to EmbryoMax Advanced KSOM medium with the specified concentrations of inhibitors and incubated for at least 20 min before placing the glass-bottom dish in the environmental control chamber. For FGF4 addition experiments, zona pellucidae were removed by brief incubation in Acid Tyrode's solution and FGF4 was co-administered with 1 μg/ml heparin (Sigma Aldrich).

### Embryo Immunofluorescence

Embryos were cultured in EmbryoMax Advanced KSOM medium with specified inhibitor or growth factor conditions prior to fixation in 4% formaldehyde (Sigma Aldrich). Fixed embryos were washed and then permeabilized in 0.5% Triton X-100 and 100 mM glycine in PBS. Embryos were blocked in 2% FBS and incubated with primary antibodies at 4°C diluted in blocking buffer overnight. Embryos were washed, blocked again, and incubated with secondary antibodies at RT for 1 h. Embryos were washed, and mounted to glass-bottom plates in PBS.

### ES Cell Culture and Editing

Embryonic stem cells were derived as previously described (Behringer, 2014). Blastocysts were dissected from mice at E3.5 and cultured overnight in advanced KSOM to allow embryos to hatch. Once hatched, blastocysts were transferred to a gelatin-coated tissue culture dish with ES culture medium and monitored for attachment and ES cell outgrowth. Outgrowths were disaggregated by incubation with 0.25% trypsin-EDTA for 5 min at 37°C and gently pipetted up and down in ES culture medium to reduce the mass of cells to 3–4 cell aggregates. The aggregates were transferred to tissue culture dishes with feeder cells and inspected for primary ES cell colony formation daily. If individual ES cell lines generated cultures with extensive differentiated cell types they were discarded. Cultures that contained primarily ES cell colonies were expanded and cultured with daily media exchanges and were passaged every 2–3 days depending on confluence and growth rate.

For imaging experiments, ES cells were incubated in 0.05% trypsin-EDTA, pipetted to disaggregate colonies into single cells, and 1–3 × 10<sup>3</sup> cells were seeded to fibronectin-coated 96-well glass-bottom plates. Single cells were cultured in ES culture medium with 2i for 48 h to allow colony formation before imaging. ES cells were imaged in ES culture medium (without 2i) unless otherwise specified. For differentiation experiments, ES cells were seeded to glass-bottom plates and then incubated in XEN differentiation media consisting of DMEM-F12 with 15% FBS, 100 U/ml penicillin and streptomycin, 100 μM BME, 10 nM retinoic acid, and 10 ng/ml Activin A for 24 h (Niakan, et al., 2013).

The Nanog-Venus-2a-mCherry targeting plasmid was cloned from a previously described mouse targeting vector, Nanog-2a-mCherry (gift from Rudolf Jaenisch, Addgene plasmid # 59995) (Faddah, et al., 2013). In brief, a short linker sequence followed by the mVenus fluorescent protein was inserted downstream of the *Nanog* coding sequence and immediately before the 2a-mCherry cassette. Approximately 1 kb each of the original 2 kb homology arms were retained and flanked with sgRNA recognition sequences identical to the target sequence in the mouse genome to generate a “double cut HDR donor” repair template plasmid (Zhang, et al., 2017). Nanog-Venus-2a-mCherry repair template was co-transfected (Lipofectamine 2000, Invitrogen) with PX458 (gift from Feng Zhang, Addgene plasmid # 48138) to ES cells derived from wild type animals (Ran, et al., 2013). Cells were cultured on neomycin resistant feeder MEFs (Milipore) and selected with G418 to obtain cells with targeted integration of repair template.

## QUANTIFICATION AND STATISTICAL ANALYSIS

### Image Analysis

Primary images from 2D time lapse data sets of cells with H2B-mCherry nuclear marker (ERK KTR ES cells, ERK KTR<sup>LoxP</sup> epidermis) were used to segment nuclei and measure fluorescence intensity with custom Cell Profiler pipelines. Nuclear masks were used to



track cells through time series (custom software) and calculate KTR C/N ratios as previously described. ERK high or ERK low divisions in ERK KTR ES cells were manually scored as cells that either have C>N KTR intensity after mitosis or cells that have N<C intensity after mitosis for at least 30 consecutive minutes. Some experiments included cells without a designated nuclear marker (NV2C ES cells) or that grow too densely for reliable segmentation (2D gut enteroids) and cells from these experiments were tracked and quantified by manually constructing regions of interest and measuring image intensity in Fiji.

For 3D embryo image sets, single cell ERK activity was tracked and quantified manually. Briefly, H2B-mCherry signal was used to define the nuclear boundary in each cell. KTR-mClover intensity was quantified in the nucleus and in a smaller region of the cytoplasm just outside the nuclear boundary. Measurements were made in the plane where the largest region of the nucleus was sampled. Both single time and time lapse KTR ratio measurements were made on live embryos and quantified in this way. For quantification of NANOG, GATA6, and CDX2 intensities, 3D image sets were analyzed by MINS as previously described (Lou, et al., 2014). In experiments where CDX2 staining was not possible, TE vs ICM cell definition was assigned by cell position within the embryo. XYZ coordinates from MINS output were used to construct a convex hull of embryo volume in MATLAB (custom software). Cells on the surface of hull were assigned TE identity and cells within hull were assigned ICM identity.

### Statistical Analysis

All graphs were generated and all statistical analysis was performed in MATLAB. Data are represented as mean  $\pm$  standard deviation or with 75<sup>th</sup> and 25<sup>th</sup> percentiles as indicated. Student's t-test or Kolmogorov-Smirnov tests were used to determine significance as indicated in legends. Significant correlation was determined by fitting data to linear regression model and using *F*-test. Sample sizes, including the number of cells, mitotic events, and embryos in each experiment and p-values are indicated in figures and figure legends.