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In vitro capture and characterization of embryonic rosette-stage pluripotency between naive and primed states

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Following implantation, the naive pluripotent epiblast of the mouse blastocyst generates a rosette, undergoes lumenogenesis and forms the primed pluripotent egg cylinder, which is able to generate the embryonic tissues. How pluripotency progression and morphogenesis are linked and whether intermediate pluripotent states exist remain controversial. We identify here a rosette pluripotent state defined by the co-expression of naive factors with the transcription factor OTX2. Downregulation of blastocyst WNT signals drives the transition into rosette pluripotency by inducing OTX2. The rosette then activates MEK signals that induce lumenogenesis and drive progression to primed pluripotency. Consequently, combined WNT and MEK inhibition supports rosette-like stem cells, a self-renewing naive-primed intermediate. Rosette-like stem cells erase constitutive heterochromatin marks and display a primed chromatin landscape, with bivalently marked primed pluripotency genes. Nonetheless, WNT induces reversion to naive pluripotency. The rosette is therefore a reversible pluripotent intermediate whereby control over both pluripotency progression and morphogenesis pivots from WNT to MEK signals.

Pluripotent cells generate all embryonic tissues and arise in the blastocyst in an immature naive state. These cells become responsive to lineage-inductive signals only after maturation into the transcriptionally and epigenetically different primed pluripotent state of the post-implantation epiblast[1]. Following implantation, the naive cells polarize and arrange into a rosette[2]. Negatively charged sialomucins, including podocalyxin (PODXL), are exposed and repulse the apical cell membranes, which form a central lumen[2,3]. Pluripotency therefore progresses through several developmental intermediates before acquiring its true potential[1,4,5]. How these transitions are regulated is unclear, and whether intermediate pluripotent states occur during this transition remains controversial. WNT and MEK signals mediate the naive-to-primed pluripotency transition in vitro[6,7], and the combination of a WNT agonist, the GSK3 inhibitor CHIR99021 (CHIR), and the MEK inhibitor PD325901 (PD; also known as 2i)[7], together with the cytokine LIF maintains naive pluripotency in embryonic stem cells (ESCs). Here, we show that WNT and MEK signals control successive pluripotency and morphogenetic transitions in the embryo that are centred on the rosette. Simultaneous WNT and MEK inhibition stabilizes rosette pluripotency in vitro. Our study demonstrates a genuine intermediate pluripotent state with its markers and the signals that control its entrance and exit (Fig. 1a).

Results

We analysed peri-implantation embryos to pinpoint the naive-primed transition in vivo. By checking copulation plugs every 3 h, we obtained blastocysts attached to the uterine wall around embryonic day 4.9 (E4.9) to E5.0. Rosettes were buried in implantation crypts around E4.9–5.2, and embryos displayed a lumen around E5.1–5.25. Several attached embryos also displayed a rosette. We identified rosettes by their central aggregation of actin fibres (Fig. 1c,f, arrows (not always visible due to the sectioning plane)), while lumens were surrounded by PODXL (Fig. 1d,g, arrows). The naive markers KLF4, NANOG and ESRRB were expressed in the blastocyst and in most rosettes, but were absent from lumen-stage embryos (Fig. 1b–e,g–j; Extended Data Fig. 1a–d, ESRRB levels slightly lower in rosettes). Importantly, rosettes expressed the primed marker OTX2, which indicates a more advanced pluripotent state (Fig. 1f; Extended Data Fig. 1b). Half of the rosettes displayed some cells lacking NANOG, some of which expressed OCT6, another primed marker (Fig. 1j, $n=12$). Lumen-stage epiblasts coexpressed OCT6 with OTX2 in all cells (Fig. 1g). Analysis

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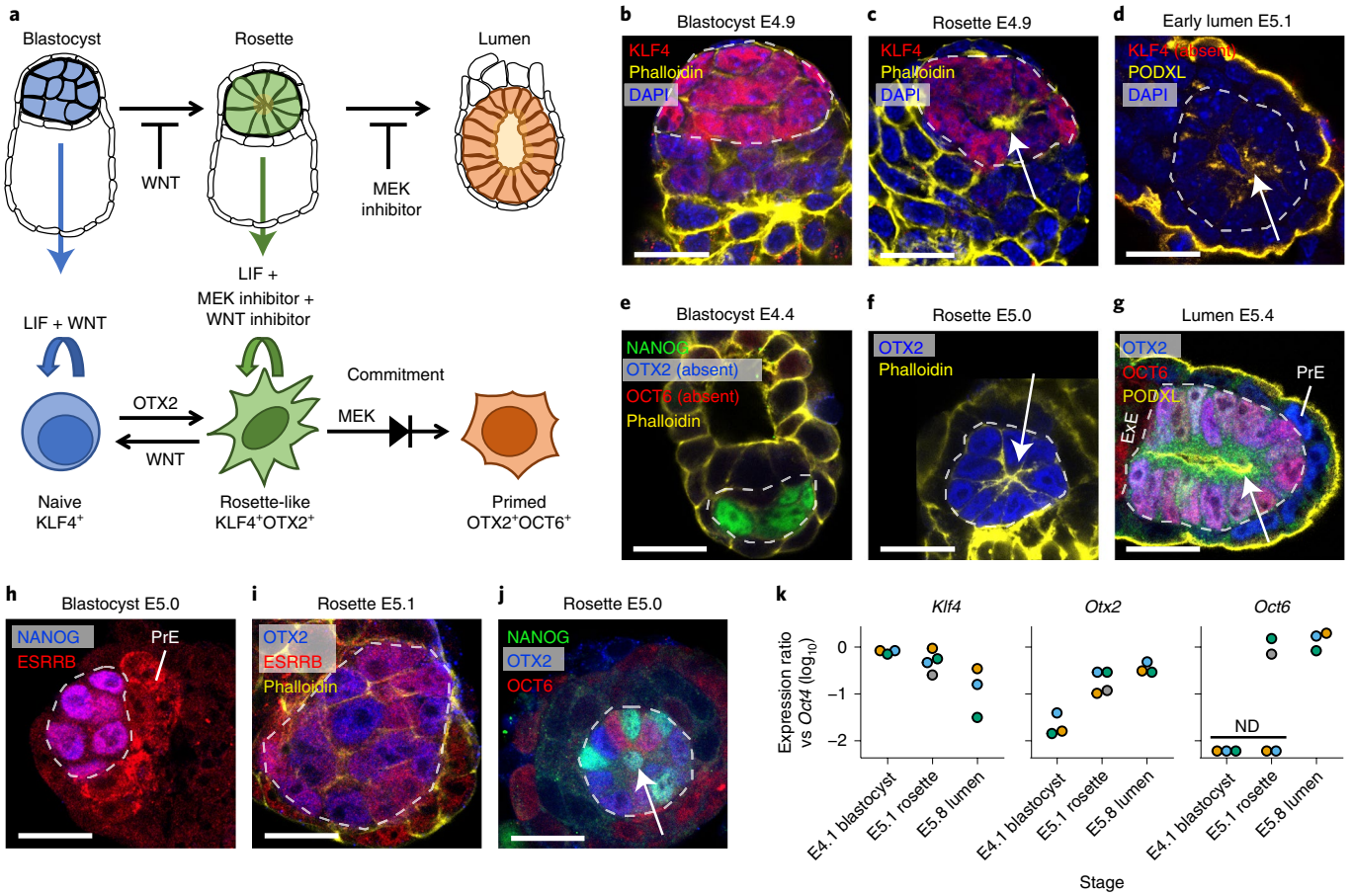


Fig. 1 | Naive and primed markers in the embryonic rosette. a, Summary of the main findings indicating the logic of signalling and embryonic and in vitro relationships. Downregulation of WNT allows the induction of OTX2, the transition from naive to rosette pluripotency and the acquisition of MEK responsivity. MEK signals drive commitment, lumenogenesis and further progression to primed pluripotency. Combined WNT and MEK inhibition captures rosette-stage pluripotency in vitro. WNT reverts rosette cells to naive pluripotency. **b–j**, Embryos of the indicated stages (blastocysts (**b**,**e**,**h**), rosettes (**c**,**f**,**i**,**j**) and early lumen (**d**,**g**)) stained for the indicated markers. The epiblast is encircled by a dashed white line. Arrows indicate the rosette centre (**c**, **f** and **j** visualized by phalloidin staining of the apical actin accumulation (in **c** and **f**)) or the lumen surrounded by PODXL (**d** and **g**). KLF4, NANOG, ESRRB, OTX2 and OCT6 were detected in 17 out of 17, 12 out of 12, 5 out of 5, 3 out of 17 and 0 out of 6 blastocysts, respectively, and in 14 out of 14, 18 out of 20, 7 out of 8, 38 out of 38 and 6 out of 12 rosettes, respectively. KLF4, NANOG, OTX2 and OCT6 were detected in 0 out of 15, 0 out of 4, 9 out of 9 and 22 out of 22 lumen-stage embryos, respectively. OCT6 was also detected in extraembryonic ectoderm (ExE), ESRRB in primitive endoderm (PrE) and OTX2 in lumen-stage PrE. **k**, Expression of *Klf4*, *Otx2* and *Oct6* relative to *Oct4* (as an epiblast-specific marker) in embryos of the indicated stages, determined by RT–PCR. Each colour-coded dot represents a group of four embryos. ND, not detected. Adjustments of individual colour channels were performed equally for all related images (**b–j**). Scale bars, 20 μm (**b–j**).

using PCR with reverse transcription (RT–PCR) confirmed that *Klf4* was downregulated during the progression from blastocyst to lumen. Meanwhile *Otx2* was induced in the rosette and *Oct6* in the lumen stage and in some rosettes (Fig. 1k). The epiblast therefore progresses to a rosette-specific state expressing the naive markers KLF4, NANOG and ESRRB with OTX2 before converting to the OTX2$^+$OCT6$^+$ primed state.

WNT and MEK signals couple successive steps in morphogenesis to pluripotency progression. We modelled rosette formation and lumenogenesis using ESCs seeded in basement membrane extract (BME)[2] to study the roles of WNT and MEK signals. After 48 h, 75% of ESC aggregates formed rosettes displaying apical–basal polarity of the actin cytoskeleton (Fig. 2a,b). About one-quarter of the aggregates contained KLF4$^+$ cells, more than 90% contained OTX2$^+$ cells and 60% contained OCT6$^+$ cells (Fig. 2c–e; Extended Data Fig. 2a,b). We monitored lumenogenesis by live staining for exposed PODXL (Extended Data Fig. 2c,d). About 70% of aggregates displayed a lumen after 65 h, similar to the percentage that formed rosettes (Fig. 2f,g).

The presence of 2i maintained KLF4 while blocking rosette formation, OTX2 and OCT6 induction and lumenogenesis. The GSK3 inhibitor CHIR (which acts as a WNT agonist) and WNT3A protein induced similar effects. Conversely, inhibiting endogenous WNT signals via the WNT inhibitor IWP2 (inhibitor of WNT production 2) promoted rosette formation, OCT6 induction and lumenogenesis (Fig. 2a–g; Extended Data Fig. 2a,b). WNT3A protein was less potent than CHIR, which is probably due to its low stability[8]. WNT/β-catenin acts by relieving transcriptional repression by TCF7L1 (refs. [9,10]). Accordingly, *Tcf7l1*$^{-/-}$ ESC aggregates, which lack TCF7L1 repression, did not form rosettes, even after WNT inhibition by IWP2 (Fig. 2h). WNT inhibition induced OTX2, which was dependent on TCF7L1 (Fig. 2i). OTX2 is a critical driver of rosette formation[3]. Indeed, ESCs in which *Otx2* was knocked out (*Otx2*KO) formed fewer rosettes, while *Otx2* overexpression (*Otx2*OE) induced rosettes even in the presence of the WNT agonist

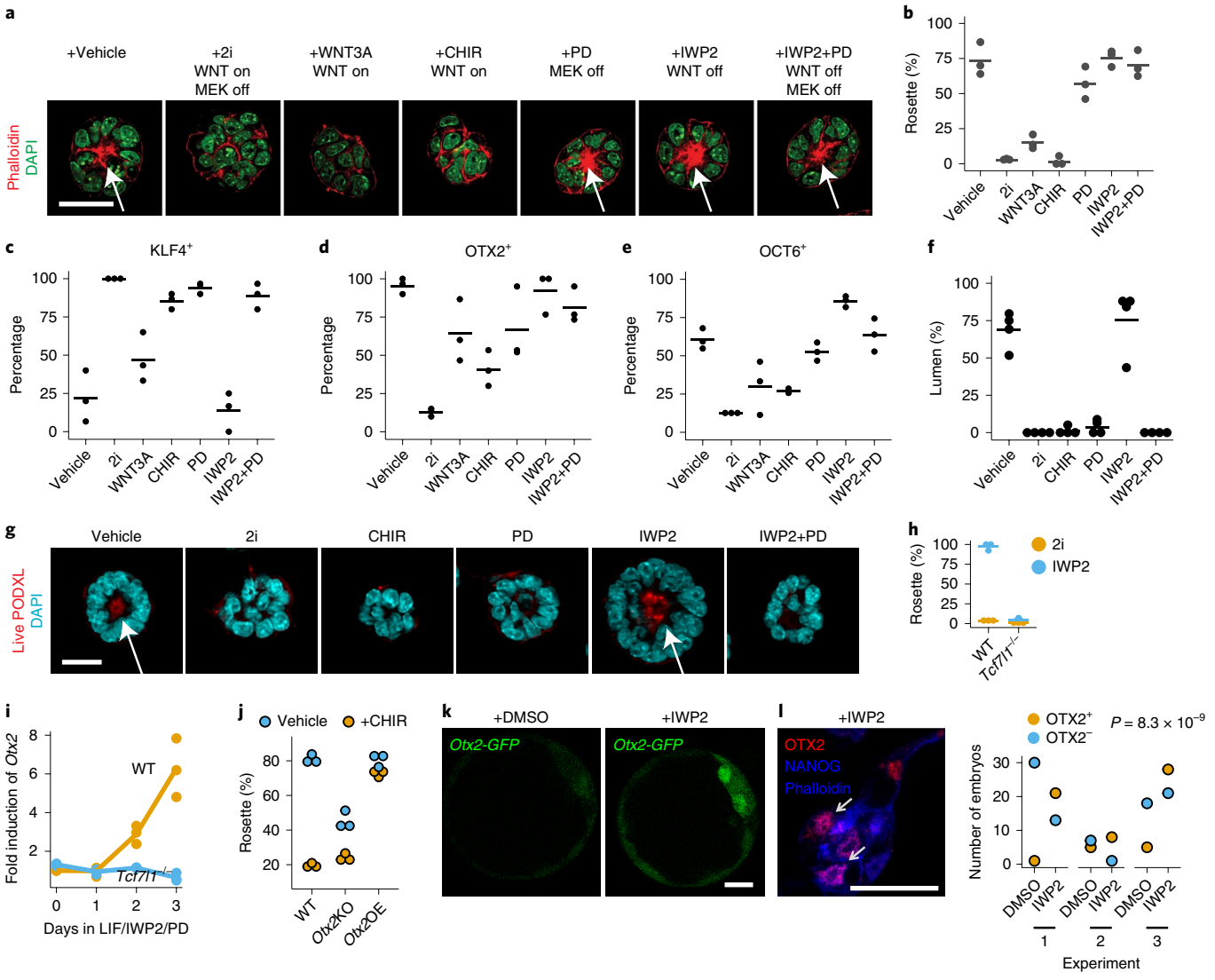


Fig. 2 | WNT and MEK signals couple successive steps in morphogenesis to pluripotency progression. a, Rosettes and aggregates 48 h after seeding ESCs in BME in the indicated conditions, stained for markers. The status of the WNT and MEK pathways as controlled by the treatments is indicated. Arrows indicate the rosette centres. **b**, The percentage of ESCs generating rosettes 48 h after seeding. $n = 3$ biological replicates. **c**–**e**, The percentage of ESC aggregates containing KLF4$^+$ (**c**), OTX2$^+$ (**d**) or OCT6$^+$ (**e**) cells 48 h after seeding. $n = 3$ biological replicates. **f**, The percentage of ESCs generating lumens 65 h after seeding. $n = 4$ biological replicates. **g**, ESC aggregates 65 h after seeding and live stained for PODXL, followed by DAPI. Arrows indicate lumens surrounded by PODXL. **h**, The percentage of wild-type (WT) or *Tcf7l1*$^{-/-}$ ESCs generating rosettes 48 h after seeding in the indicated conditions. $n = 3$ independent experiments. **i**, Induction of *Otx2* in *Tcf7l1*$^{-/-}$ ESCs compared to WT after transfer to LIF/IWP2/PD (RT–PCR, three technical replicates). **j**, Percentage of *Otx2*KO or *Otx2*OE ESCs generating rosettes 48 h after seeding. $n = 3$ independent experiments. **k**, Four-cell-stage *Otx2*–GFP embryos cultured for 75 h and imaged for GFP. A GFP$^+$ DMSO-treated (3 from 11) and representative IWP2-treated (10 from 11) embryos are shown. **l**, Four-cell-stage embryos cultured to the expanded blastocyst stage. Arrows indicate OTX2 in NANOG$^+$ epiblast of an IWP2-treated embryo. Three independent experiments with $n = 158$ embryos (66 DMSO, 92 IWP2) are summarized in the chart on the right. Two-sided Fisher's exact test. **m**–**o**, Embryos of the indicated stages stained for markers. Arrows indicate pERK$^+$ epiblast cells. Images representative of three blastocysts (**m**), none positive; nine rosettes (**n**), three displaying a pERK$^+$ cell; four lumen stages (**o**), with six, four, three and four pERK$^+$ cells. pERK was also observed in primitive endoderm (**m**). **p**, Blastocysts cultured for 77 h with DMSO or PD and stained for markers; 8 out of 16 and 1 out of 46 OCT4$^+$ embryos displayed a lumen (arrow). Dotted white line indicates the epiblast (**m**–**p**). Images represent three (**a**) or four (**g**) independent experiments. Scale bars, 20 µm (**a**,**g**,**k**–**p**). For **b**–**f** and **h**, bars represent the mean.

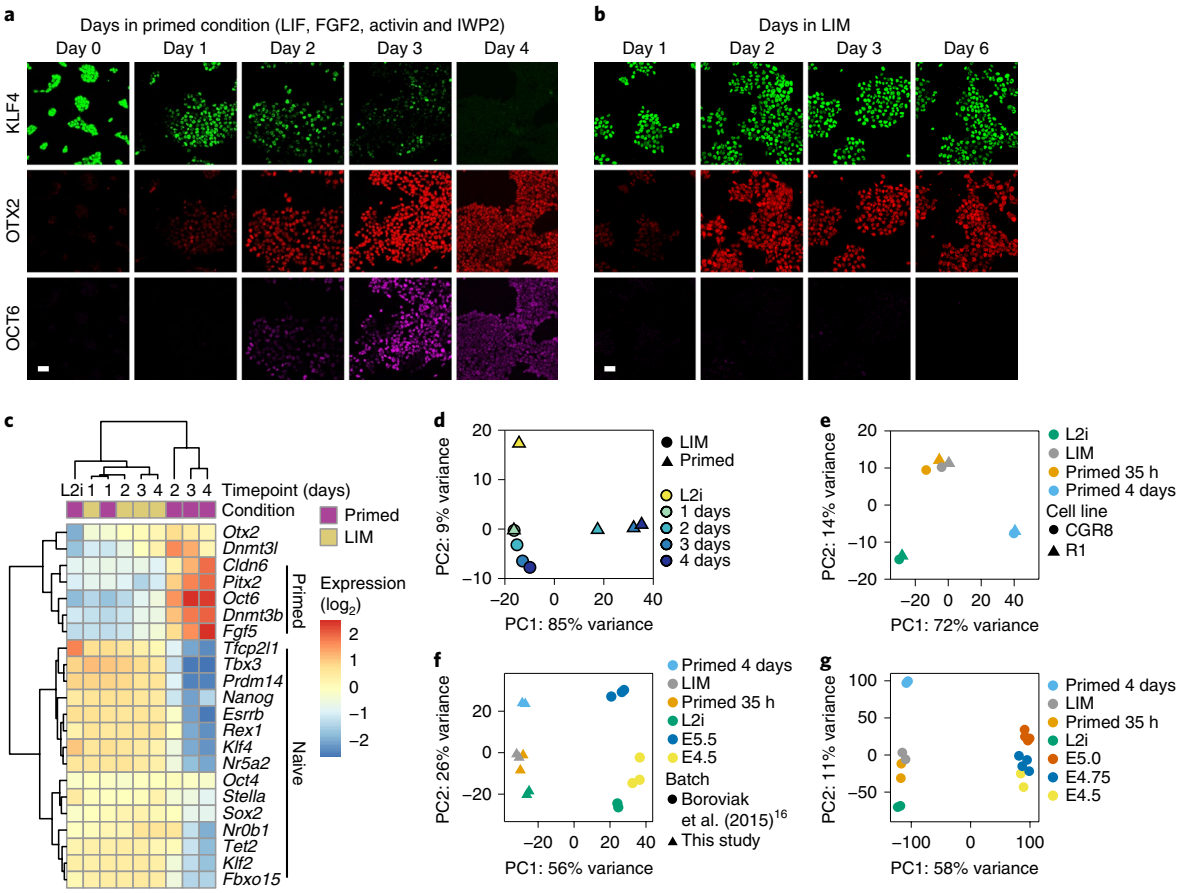


Fig. 3 | WNT controls the initial phase of the naive-primed transition. a,**b**, ESCs plated in primed (**a**) or LIM (**b**) conditions for the indicated number of days and immunostained for the indicated markers (3 independent experiments). **c**, RNA-seq heatmap from R1 ESCs plated in primed or LIM conditions for the indicated number of days. **d**, PCA of RNA-seq data from R1 ESCs plated in primed or LIM conditions for the indicated number of days. At day 1 the samples for both conditions overlap. One sample for each condition. **e**–**g**, PCA of RNA-seq data from R1 and CGR8 ESC lines maintained in the indicated conditions compared with L2i ESCs (**e**), E4.5 and E5.5 epiblast (**f**, from ref. [16]), or with E4.5, E4.75 and E5.5 epiblast (**g**, from ref. [3]). Each dot indicates a biological replicate. Scale bars, 40 µm (**a**,**b**).

CHIR (Fig. 2j). OTX2 is therefore a key effector repressed by the WNT pathway. The MEK inhibitor PD permitted rosette formation and OTX2 induction, especially in combination with IWP2, but most rosettes maintained KLF4 expression and few induced OCT6 and progressed to lumenogenesis, arresting in the KLF4$^+$OTX2$^+$ state (Fig. 2a–g; Extended Data Fig. 2a,b). Downregulation of the WNT/β-catenin–TCF7L1 pathway therefore induces OTX2 and rosette formation, whereas MEK signals drive the subsequent lumenogenesis and progression to primed pluripotency.

WNT signals are active in the blastocyst[6,11–13]. To investigate whether these prevented the inner cell mass (ICM) from inducing OTX2, we cultured four-cell-stage *Otx2*–GFP (green fluorescent protein) reporter embryos in the presence of IWP2. After 56 h, blastocysts formed in which the ICM displayed faint GFP signals that rapidly increased in the subsequent 6 h. In contrast, in the absence of IWP2, the ICM induced faint GFP in few embryos after 75 h (Fig. 2k). Immunostaining of four-cell-stage embryos cultured until the expanded blastocyst stage confirmed this result, whereas phalloidin staining did not reveal evidence of rosette morphogenesis (Fig. 2l). Thus, blastocyst WNT signals prevent premature progression of the ICM to OTX2$^+$ rosette pluripotency, while additional factors, for example, basal lamina deposition[2], are required for rosette morphogenesis.

MEK signals induce the phosphorylation of ERK, typically as a series of stochastic pulses at a frequency reflecting the strength of the signal[14]. The epiblast of late blastocysts and most rosettes did not display phosphorylated ERK (pERK) (Fig. 2m). However, some rosettes contained a single pERK$^+$ cell (Fig. 2n). These rosettes displayed OTX2 but not OCT6 expression (Extended Data Fig. 2e). In lumen-stage epiblasts, all cells were OCT6$^+$ and many displayed ERK phosphorylation (Fig. 2o; Extended Data Fig. 2f). Thus, pERK pulses increase in frequency during progression to the lumen stage. To investigate their function, we cultured blastocysts to the lumen stage. The embryos lost KLF4 expression 41 h after attachment to the substrate and displayed lumenogenesis after 77 h. However, MEK inhibition maintained the expression of KLF4 and NANOG in the epiblast, which generated a ball-shaped or disorganized mass lacking a lumen (Fig. 2p). These data suggest that pluripotency progression is coupled to embryonic morphogenesis first by WNT signals and, subsequently, in the rosette stage, by MEK signals (Fig. 1a).

WNT controls the initial phase of the naive-primed transition. We observed a transient appearance of rosette-like KLF4$^+$OTX2$^+$ OCT6$^-$ cells 0.5–1 day after ESCs that were grown in N2B27 medium supplemented with LIF and 2i (hereafter called L2i) were differentiated in N2B27 medium without supplements, or differentiated towards epiblast-like cells (EpiLCs), which are primed cells resembling egg cylinder epiblast[15] (Extended Data Fig. 3a,b). To focus on the roles of WNT and MEK, we maintained the

presence of LIF while inhibiting endogenous WNT signals using IWP2. We also activated MEK using FGF2 and supported the primed cells with activin. LIF slows but does not inhibit the naive-primed transition in this condition (hereafter called 'primed condition')[6]. OTX2 was induced after 1 day and coexpressed with KLF4 until conversion to the KLF4$^-$OTX2$^+$OCT6$^+$ primed state after 3 days (Fig. 3a). Importantly, ESCs transferred from L2i into conditions in which both WNT and MEK were inhibited (hereafter called LIM for LIF, IWP2 and MEK inhibitor) also induced OTX2, but arrested in the KLF4$^+$OTX2$^+$OCT6$^-$ rosette-like state (Fig. 3b). Since L2i and LIM differ only in WNT pathway activity (active versus repressed, respectively), this result indicates that WNT signals are required to prevent OTX2 induction and pluripotency progression.

RNA-sequencing (RNA-seq) showed that repression of naive and induction of primed genes mostly occurred after 2 days in primed conditions and appeared complete after 3–4 days, while after 4 days in LIM, pluripotency gene expression resembled that after 1–2 days in primed conditions (Fig. 3c; Supplementary Table 1). Principal component analysis (PCA) separated primed from other conditions alongside principal component 1 (PC1), and L2i from LIM alongside PC2 (Fig. 3d). During the first day, however, the transcriptomes for both LIM and primed conditions changed only alongside PC2, despite the opposite status of the MEK pathway. Extending this time frame, the transcriptomes of two ESC lines after 8 (R1) or 15 (CGR8) days in LIM grouped together with those after 35 h in primed conditions (Fig. 3e; Supplementary Table 2). Loss of WNT signals therefore drives part of the naive-primed transition, independent from MEK activity. Moreover, combined WNT and MEK inhibition arrests pluripotent cells in an intermediate naive-primed, rosette-like state.

A previous transcriptome analysis showed that naive pluripotency appears sometime within E3.75–4.5 and is absent from E5.5 epiblasts[16]. PCA of these data with ours captured the developmental progression in PC2 (Fig. 3f). ESCs from both studies mapped before E4.5 ICM alongside this developmental axis, while the 4-day-primed samples mapped alongside E5.5 epiblasts. Importantly, cells in LIM mapped together with 35-h primed cells, confirming their partway transition towards primed pluripotency. A comparative transcriptome analysis with peri-implantation embryos[3] again mapped the L2i ESCs before E4.5 ICM, while cells in LIM showed most similarity with E5.0 epiblasts (Fig. 3g). These data support our findings that loss of WNT signalling induces the naive-primed transition, while MEK inhibition arrests the cells only at a more advanced rosette-like state.

MEK and WNT inhibition maintains rosette characteristics. We derived stable cell lines from blastocysts in LIM. Outgrowths generated dispersed colonies that expanded long term (Extended Data Fig. 3c). After blastocyst injection, all three tested cell lines contributed to live chimeras, induced strong sex conversion and showed germline transmission (Extended Data Fig. 3d; Supplementary Table 3). Although characteristic for ESCs, contribution to blastocyst chimeras is found for a spectrum of pluripotent states, including E5.0 peri-implantation epiblast cells[17] and EpiSCs maintained in differentiation-suppressing conditions[18,19].

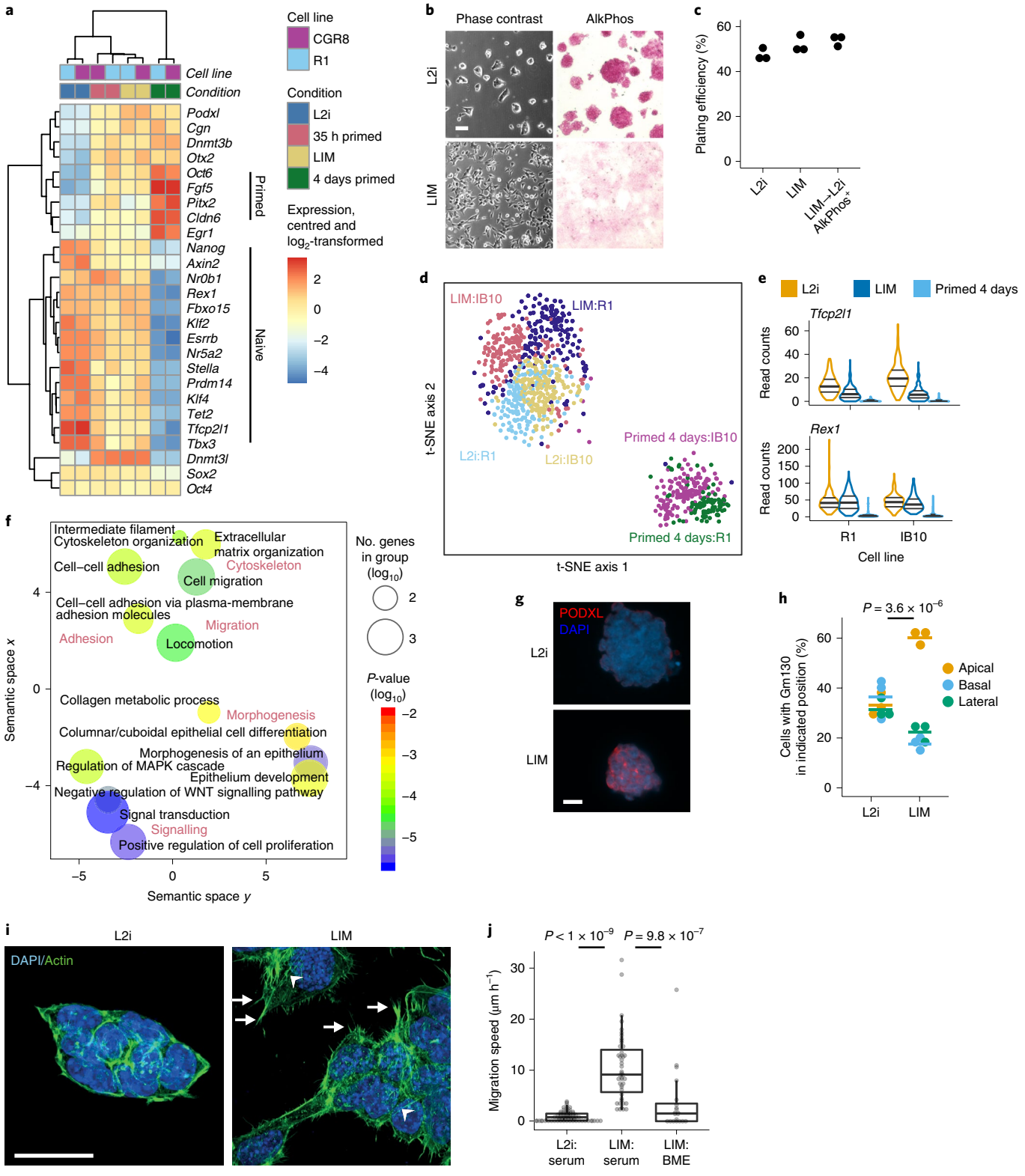
LIM supported the expansion of ESCs and blastocyst-derived cell lines for more than 28 passages on serum or BME coatings. Repression of the WNT target *Axin2* and the MEK target *Egr1* verified pathway inhibition (Extended Data Fig. 3e). The lines maintained expression of OCT4, SOX2, the naive markers KLF4, NANOG and ESRRB (at a slightly reduced level), and OTX2 while repressing OCT6 (Extended Data Fig. 4a). The cells displayed dispersed monolayer morphology, low alkaline phosphatase activity, tolerated clonal passaging, retained naive gene expression, LIF dependency and active X chromosomes, and proliferated slightly slower than ESCs (Fig. 4a–c; Extended Data Figs. 3e and 4b–d). Single-cell sequencing and t-distributed stochastic neighbour embedding (t-SNE) analysis revealed a KLF4$^+$OTX2$^+$OCT6$^-$ LIM population distinct from naive and primed cells (Fig. 4d; Extended Data Fig. 4e). While expression of the pluripotency marker TFCP2L1 was reduced, most pluripotency markers, such as REX1, showed similar expression level distributions in cells grown under LIM and L2i conditions (Fig. 4e). Of note, LIM cells repressed OTX2, regenerated alkaline-phosphatase-positive colonies and acquired ESC-like proliferation rates following transfer to L2i (Fig. 4c; Extended Data Fig. 4d,f), which indicates that WNT induced reversion to the ESC state.

Based on bulk transcriptome data, 531 and 196 genes were more than twofold differentially expressed between LIM and L2i cells (Supplementary Table 2). Gene Ontology (GO) analysis revealed processes involving cell adhesion, cell migration, epithelial cell differentiation and morphogenesis, and MEK signalling, which is consistent with the phenotypic changes observed in vitro and in the rosette (Fig. 4f; Supplementary Table 4). Motility and adhesion-related genes are induced early in the naive-primed transition[20,21], which possibly reflects morphogenetic processes such as polarization and lumenogenesis[2]. Indeed, two factors instrumental for rosette lumenogenesis were induced: PODXL, which induces cell membrane separation, and cingulin (CGN), which mediates its cell surface exposure[3] (Fig. 4a,g). Furthermore, LIM cells gained apical–basal polarity and displayed filopodia, stress fibres and a dispersed appearance, which all indicate motility (Fig. 4b,h,i; Extended Data Figs. 3c and 4g). Live imaging showed the cells migrating at approximately 10 μm h^{-1} on serum coatings (Fig. 4j; Extended Data Fig. 4h). E5.0–6.0 epiblast cells disperse after detaching from the basal lamina during mitosis[22], which suggests that the absence of lamina contacts induces motility. Accordingly, surfaces coated with the basal lamina extract BME reduced motility (Fig. 4j). Together,

Fig. 4 | MEK and WNT inhibition maintains rosette characteristics. a, RNA-seq heatmap from R1 and CGR8 ESCs cultured in the indicated conditions. LIM 8 days (R1) or 15 days (CGR8). **b**, Phase-contrast and alkaline-phosphatase (AlkPhos)-stained images of R1 ESCs maintained in L2i or LIM for three passages. Three independent experiments were performed, with similar results obtained. **c**, The plating efficiency of ESCs and LIM cells, and the formation of alkaline-phosphatase-positive colonies by LIM cells after plating in L2i. 3 independent experiments. **d**, t-SNE plot of single-cell RNA-seq data of the indicated cell lines and conditions. **e**, Distribution of the expression levels of the indicated genes in the single-cell RNA-seq data. Violin plots show the median (centre bold bar), the first and third quartiles on top of the expression level distribution. $n = 159$, 178, 102 (R1) and 184, 149 and 164 (IB10) cells for L2i, LIM and primed, respectively (**d** and **e**). **f**, GO analysis of genes differentially expressed between L2i and LIM cells (two samples each). The *P* values were calculated from minimum hypergeometric scores from a ranked gene list using GOrilla[58]. GO terms were aggregated and plotted according to their semantic relationships using REVIGO[59]. **g**, Immunostaining for PODXL of R1 ESCs maintained in L2i or LIM for six passages (three biological replicates). **h**, Apical–basal polarity in L2i and LIM cells was determined by Gm130 immunostaining for polarized location of the Golgi complex. Two-way analysis of variance for effect of the condition. $n = 3$ independent experiments; bar represents the mean. **i**, L2i and LIM cells stained with phalloidin to visualize the actin cytoskeleton. Filopodia (arrows) and stress fibres (arrowheads) are indicated (three independent experiments). **j**, Migration speed of L2i and LIM cells on serum-coated or BME-coated slides. The centre line indicates the median, the boxes the first and third quartiles, while the whiskers extend to the most extreme value no further than 1.5-times the inter-quartile range. $n = 80$, 47, 20 cells for L2i, serum and BME, respectively. Two-tailed Mann–Whitney test. Scale bars, 20 μm (**g**,**i**) or 40 μm (**b**).

our data argue that WNT inhibition induces pluripotency progression to the rosette-like state, while MEK inhibition arrests further progression and maintains this state. Therefore, we propose that WNT/MEK-inhibited cells are called rosette-like stem cells (RSCs).

OTX2 enables MEK-driven transition to primed pluripotency. We monitored the naive-primed transition using the *Rex1*::destabilized-GFP (RGd2) reporter, which is thought to track exit from naive pluripotency[21]. While RGd2 ESCs and RSCs displayed similar levels of the reporter, consistent with our single-cell RNA-seq data, RSCs downregulated the reporter within 24 h in the primed condition, more than a day earlier than ESCs (Fig. 5a; Extended Data Fig. 5a). Likewise, RSCs carrying *Tbx3*–RFP naive and *Oct6*–GFP reporters[23] switched more quickly to primed



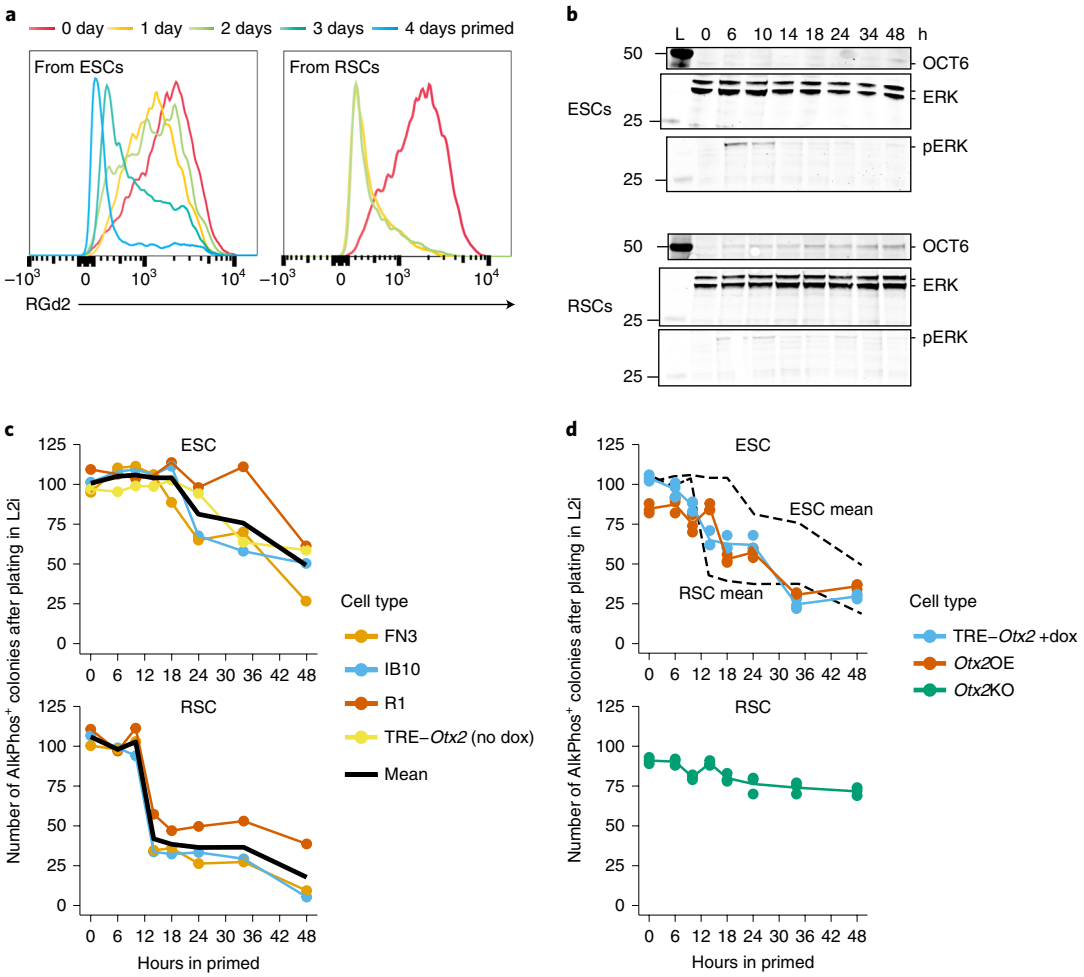


Fig. 5 | OTX2 enables MEK-driven transition to primed pluripotency. a, RGd2 ESCs and RSCs were plated in primed conditions and analysed daily for GFP by flow cytometry (three independent experiments). **b**, Western blots showing OCT6, ERK (same gel) and pERK (on different gel) in FN3 ESCs and RSCs after transfer to primed medium for the indicated durations (three biological replicates; full blots are available as source data). L, ladder. **c**,**d**, Number of alkaline-phosphatase-positive colonies generated in L2i by 200 ESCs (upper) or RSCs (lower) after incubation in primed medium for the indicated times. Cell lines used: FN3, IB10, R1 and TRE-*Otx2* (no doxycycline (dox) (**c**), and TRE-*Otx2* (+ dox), *Otx2*OE and *Otx2*KO (**d**). $n = 4$ (ESC) or $n = 3$ (RSC) biological (**c**) or $n = 4$ technical (**d**) replicates.

reporter expression than ESCs (Extended Data Fig. 5b). Moreover, RSCs induced OCT6 protein expression after 6 h in the primed condition, followed by a sudden drop in their ability to establish alkaline-phosphatase-positive colonies in L2i, which indicates their commitment to primed pluripotency. In contrast, ESCs induced OCT6 expression after 34 h, and their reversibility declined only slowly after 24 h (Fig. 5b,c; Extended Data Fig. 5c). Cell lines of diverse backgrounds, including non-permissive FVB/N (FN3), showed this difference. Therefore, RSCs are poised for conversion to primed pluripotency after release of MEK inhibition, whereas ESCs require a prior transition to the rosette-like state. Nonetheless, both ESCs and RSCs responded with similar pERK oscillations following transfer to primed conditions (Fig. 5b; Extended Data Fig. 5c). The WNT pathway—the only difference between L2i and LIM—must therefore modify the pERK response. OTX2 is induced following the loss of WNT activity (Fig. 2i) and acts early in the naive-primed transition[23–25]. Importantly, RSCs lacking OTX2 failed to commit to primed pluripotency and maintained their reversal efficiency even after 48 h in primed condition (Fig. 5d). Moreover, ESCs that over-expressed OTX2, either constitutively or by TRE-based doxycycline induction, showed a similarly rapid drop in reversibility as that of RSCs in the primed condition (Fig. 5d; Extended Data Fig. 5d). OTX2 therefore enables the progression to primed pluripotency in response to MEK signals.

A primed chromatin landscape in RSCs. L2i ESCs display DNA hypomethylation, which is associated with a low expression of DNA methyltransferases[26–30]. After transfer to primed conditions, 5-methylcytosine (5mC) levels rapidly increased from around 1% to around 4% after 3 days, while 5-hydroxymethylcytosine (5hmC) was low in naive and primed cells but elevated during the transition, which suggests that there is 5mC turnover (Fig. 6a). RSCs expressed *Dnmt3b* and *Dnmt3l*, and the protein DNMT3L was strongly induced in RSCs and embryonic rosettes, which suggests that there is a departure from the hypomethylated ground state (Figs. 4a and 6b; Extended Data Fig. 6a,b). Indeed, 5mC and 5hmC levels were similarly elevated in RSCs to that of cells halfway through the naive-primed transition (Fig. 6a).

L2i ESCs are further characterized by the presence of bivalent gene promoters[27,31], carrying both activating H3K4me3 and repressive H3K27me3 chromatin marks[32–34]. Comparative chromatin immunoprecipitation (ChIP) with sequencing (ChIP-seq) analysis of two RSC lines with L2i ESCs[27] and EpiLCs[35] showed that RSCs clustered with EpiLCs regarding H3K4me3, thus suggesting

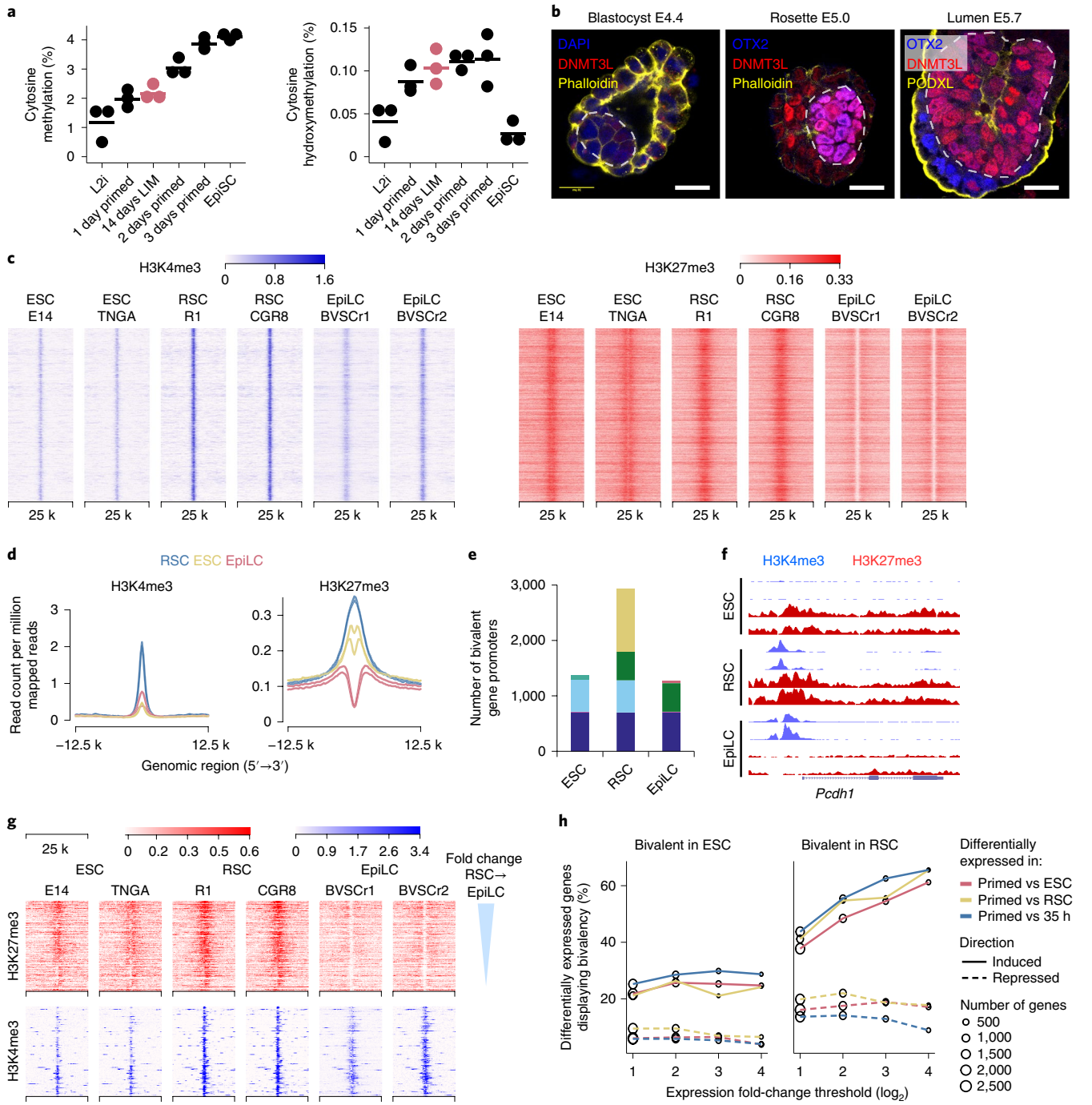


Fig. 6 | A primed chromatin landscape in RSCs. a, Levels of cytosine methylation and cytosine hydroxymethylation in ESCs (black) or RSCs (red) cultured in the indicated conditions ($n=3$ biological replicates, bars represent the mean). **b**, Blastocyst-stage ($n=4$), rosette-stage ($n=7$) and lumen-stage ($n=5$) embryos stained for DNMT3L, OTX2 and phalloidin or PODXL. The epiblast is encircled by a dotted white line. DNMT3L is also visible in the extraembryonic ectoderm (rosette-stage). **c**, H3K4me3 and H3K27me3 heatmaps of the bivalent cluster. **d**, Average intensity plots of H3K4me3 and H3K27me3 in bivalent regions. **e**, The number of promoters carrying bivalent marks in naive, rosette-like or primed cells. Groups of promoters are coloured depending on the cell type(s) in which they display bivalency. **f**, Screenshot from the UCSC genome browser showing H3K4me3 (blue) and H3K27me3 (red) peaks on the *Pcdh1* gene. **g**, H3K27me3 and H3K4me3 heatmaps centred over promoters of genes induced during the RSC-to-primed transition, ranked according to the fold-change in induction. **h**, The fraction of gene promoters carrying bivalent marks in ESCs (in L2i) or RSCs, plotted against expression level changes. Scale bar, 20 µm (**b**).

a shift towards the primed state, but revealed a distinct pattern of H3K27me3 (Extended Data Fig. 6c–e). *K*-means clustering of the ChIP-seq peaks revealed bivalent (Fig. 6c,d), H3K4me3-enriched and H3K27me3-enriched clusters (Extended Data Fig. 6f,g). RSCs gained H3K4me3 and H3K27me3 marks in the bivalent cluster compared to ESCs, while EpiLCs lost H3K27me3. Consequently, RSCs contained more than twice as many bivalent genes as ESCs or EpiLCs (Fig. 6c–e; Supplementary Table 5). Many primed genes, such as *Pcdh1*, *Oct6* and *Fgf5*, gained bivalency (Fig. 6f; Extended Data Fig. 6h). Moreover, genes that showed a stronger expression

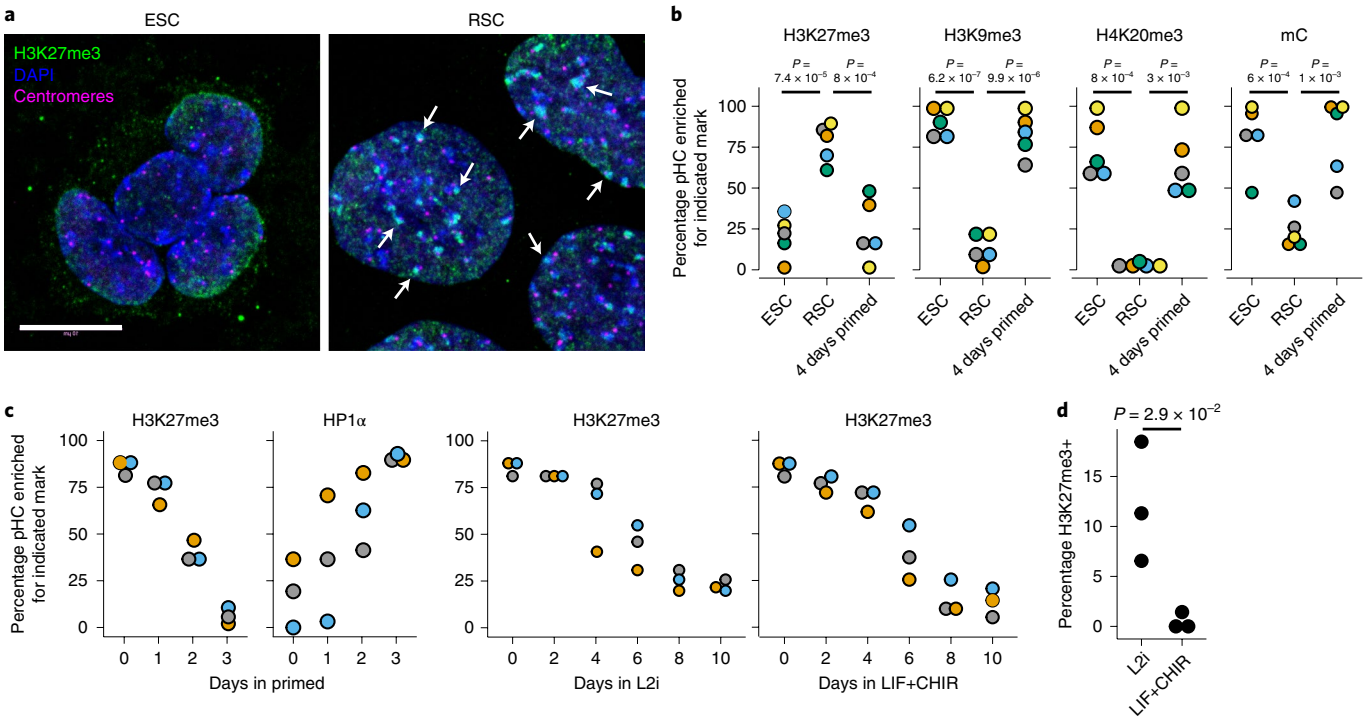


Fig. 7 | RSCs remodel pHC. a, Immunostaining for H3K27me3 (green) and the centromeres (magenta) in R1 ESCs and RSCs. Arrows indicate examples of pericentric H3K27me3 foci (light blue, overlap of H3K27me3 and DAPI). Images are representative of three experiments. **b**, The percentage of metaphases in indicated conditions in which the pericentric regions are enriched for the indicated mark. Each dot represents the average of 100 metaphases. $n=5$ biological replicates. **c**, RSCs were cultured in the indicated conditions and the percentage of cells displaying pHC foci positive for the indicated marks was determined. $n=3$ biological replicates. **d**, ESCs were cultured for 4 weeks in L2i or LIF+CHIR, and the fraction of cells displaying H3K27me3$^+$ pHC foci was determined. $n=3$ biological replicates. (**b** and **d**) Unpaired two-tailed *t*-test. Scale bars, 10 μm (**a**).

difference between primed cells and RSCs were more likely to display bivalency in RSCs, a correlation that was not observed with bivalent genes in ESCs or with downregulated or naive genes (Fig. 6g,h). Thus, the increase in bivalent genes in RSCs foreshadowed their induction following the transition to primed pluripotency, information that is absent from ESCs. The naive-to-rosette transition therefore sets up a chromatin landscape that is permissive for primed pluripotency, which is consistent with the rapid response of RSCs to MEK activation.

Rosette-specific formatting of pericentric heterochromatin. H3K27me3 is a feature of facultative heterochromatin, which evolved to regulate gene expression. However, immunostaining of RSCs revealed its accumulation on pericentric heterochromatin (pHC), which is 4,6-diamidino-2-phenylindole (DAPI)-dense heterochromatin surrounding the centromeres that is normally characterized by constitutive heterochromatin marks, including H3K9me3, H4K20me3, DNA methylation and HP1α[34] (Fig. 7a). Metaphase chromosome spreads showed enrichment of pericentric H3K27me3 in 60–90% of RSCs and only in a minority of ESCs or primed cells, while pericentric H3K9me3, H4K20me3 and 5-methylcytosine were depleted in most RSCs (Fig. 7b; Extended Data Fig. 7a–d), which indicates that RSCs erased constitutive heterochromatin marks and replaced them with facultative marks.

After transfer to primed conditions, RSCs lost KLF4 and erased most pericentric H3K27me3 within 2 days while simultaneously gaining pericentric HP1α, which indicates that there is a reversion to canonical constitutive heterochromatin following the transition to primed pluripotency (Fig. 7c; Extended Data Fig. 7e). Reverting the cells to ESCs by L2i also induced erasure of pericentric H3K27me3, but at a slower rate, and a subset of cells retained the mark, even after extensive culture (Fig. 7c,d; Extended Data Fig. 7f). Indeed, pericentric H3K27me3 was recently observed in a subset of L2i ESCs[36]. However, transfer to LIF+CHIR erased virtually all pericentric H3K27me3 (Fig. 7c,d), which suggests that MEK inhibition induces epigenetic heterogeneity in ESCs.

Importantly, while peri-implantation blastocysts displayed little H3K27me3 immunostaining, OTX2$^+$ embryos displayed nuclear accumulation of H3K27me3 in the epiblast, including staining of the pericentric regions. Post-implantation KLF4$^+$OTX2$^+$ embryos displayed further accumulation of H3K27me3 and intense pericentric staining in many epiblast cells, which was lost after lumenogenesis, and loss of KLF4 (Fig. 8a–c; Extended Data Fig. 8a,b). Conversely, rosettes lost pericentric H3K9me3 and displayed mosaic 5-methylcytosine staining (Fig. 8d,e). Thus, the rosette exchanges constitutive with facultative heterochromatin marks, which is reversed after lumenogenesis. Moreover, pericentric H3K27me3 appeared after the premature induction of rosette-stage pluripotency in blastocysts by IWP2, and persisted when arresting embryos in rosette pluripotency by MEK inhibition (Fig. 8f,g). The striking accumulation of pericentric H3K27me3 in the embryonic rosette and in RSCs supports the shared nature of their pluripotent state. Finally, *Otx2*KO ESCs repressed pericentric H3K27me3 accumulation in LIM, while *Otx2*OE ESCs increased it (Fig. 8h), thus confirming the central role of OTX2 in controlling rosette pluripotency.

Discussion

We identified an intermediate pluripotent state in the embryonic rosette and demonstrated that WNT controls the naive-rosette transition and MEK the rosette-primed transition. This sequential action allows the capture of embryonic (ESC)[37,38], rosette-like (RSC) and epiblast (EpiSC)[39,40] stem cells. RSCs differ from ESCs

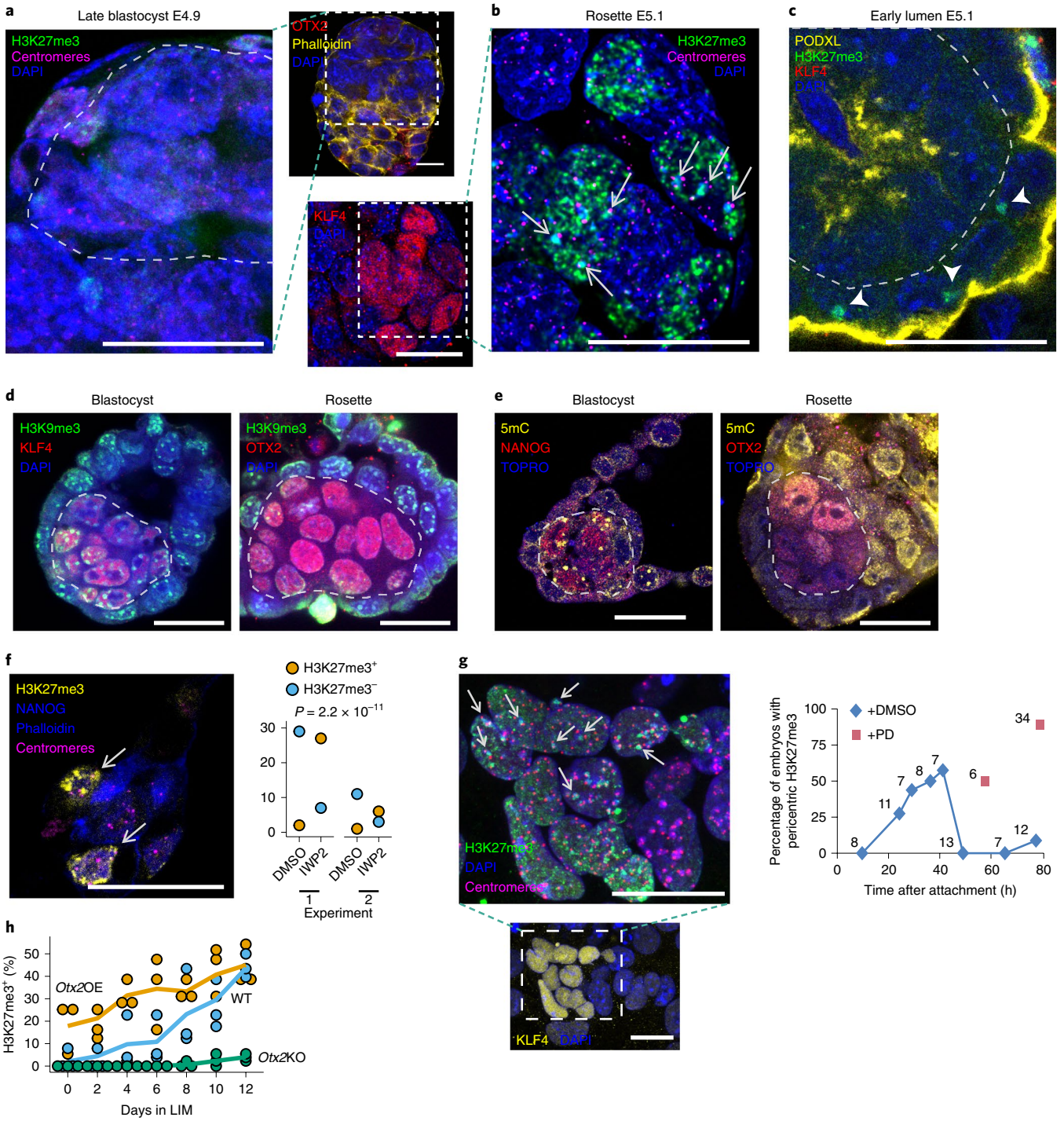


Fig. 8 | Rosette-specific formatting of pHC. a, Blastocyst (16 samples) stained for H3K27me3 (green), centromeres (magenta), OTX2 (red), phalloidin (yellow) and DAPI (blue) shows absence of OTX2 and H3K27me3 foci in the epiblast. **b**, Rosette (23 samples) stained for H3K27me3 (green), centromeres (magenta), KLF4 (red) and DAPI (blue). White arrows indicate some H3K27me3$^+$ pHC foci (light blue, overlap of H3K27me3 and DAPI). **c**, Early lumen-stage embryo (15 samples) stained for H3K27me3 (green), PODXL (yellow) and DAPI (blue). No enrichment of H3K27me3 at the pHC is observed. The outer primitive endoderm layer displays enrichment of H3K27me3 on the inactive X chromosomes (arrowheads). **d**, Blastocyst (8 samples) and rosette (7 samples) embryos stained for KLF4 or OTX2 (red), H3K9me3 (green) and DNA (blue). **e**, Blastocyst (15 samples) and rosette (18 samples) embryos stained for NANOG or OTX2 (red), 5mC (yellow) and DNA (blue). The epiblast is encircled by dotted white line (**a**,**c**–**e**). **f**, Four-cell-stage embryos were cultured to the expanded blastocyst stage in the indicated conditions and the presence of pericentric H3K27me3 (arrows) indicated in a representative IWP2-treated embryo. Two independent experiments, $n = 86$ embryos (43 for each condition); two-sided Fisher's exact test. **g**, Left: embryos cultured in the presence of a MEK inhibitor (PD) for 57 h after attachment and stained for KLF4 (yellow), H3K27me3 (green), centromeres (magenta) and DAPI (blue). Right: the number of embryos cultured for the indicated time, and the fraction that displayed H3K27me3-enriched pHC foci. Each time point indicates an independent experiment. **h**, *Otx2*OE, WT or *Otx2*KO ESCs were cultured in LIM for the indicated duration and the percentage of cells displaying H3K27me3$^+$ pHC foci quantified ($n = 3$ technical replicates). Scale bars, 20 µm (**a**–**g**).

Table 1 | Characteristics of mouse pluripotent stem cells

Characteristic	ESCs	RSCs	EpiSCs
Embryonic equivalent	E3.75–4.5 ICM[16]	Rosette	Pre-streak to early-streak epiblast[18]
Contribution to blastocyst chimeras	High efficiency	High efficiency	Low efficiency
Growth factor requirements	LIF + WNT[6]	LIF; inhibition of WNT and MEK signals	FGF2 + activin A[39,40]
Morphology	Dome-shaped colonies	Flat dispersed colonies	Flat colonies
Polarity	Unpolarized	Apical–basal polarity	Apical–basal polarity
Alkaline phosphatase	Strong	Faint	Faint
Clonogenicity	High	High	Low
Migratory	No	Yes	No
Transcriptome	Naive transcription factors	Naive transcription factors + OTX2	Primed transcription factors
X-inactivation status	XaXa	XaXa	XaXi
DNA methylation	Low	Intermediate	High
DNA hydroxymethylation	Low	High	Low
Promoter bivalency	Low	High in primed genes	Low
Main pHC mark	H3K9me3/5meC	H3K27me3	H3K9me3/5meC
Ability to efficiently revert to earlier state	Low	High	Low

XaXa, two active X chromosomes; XaXi, one inactive X chromosome.

by their bivalent marking of primed genes, facultative formatting of pHC, primed pluripotency progression in response to MEK activation and a similar transcriptome as cells halfway through the naive-primed transition, among other features (Table 1). We further showed that OTX2 acts downstream of WNT to establish rosette-like pluripotency, including the MEK responsiveness and pHC formatting. Finally, WNT and MEK signals control rosette formation and lumenogenesis, respectively, thereby coordinating pluripotency progression with embryonic morphogenesis (Fig. 1a).

Several studies argue that ESCs do not require WNT signals since they maintain naive gene expression and chimera-forming ability in their absence[10,41–43]. Our identification of rosette pluripotency—which shares those characteristics with ESCs—resolves this controversy: in the absence of WNT, ESCs convert into RSCs that share defining features with the embryonic rosette. Furthermore, our findings that WNT activation and MEK inhibition block sequential embryonic transitions—naive-rosette and rosette-primed, respectively—strongly argue that ESCs and RSCs represent distinct embryonic stages. Of note, the proposed attributes of the hypothetical 'formative' naive-primed intermediate, such as downregulation of naive gene expression and random X chromosome inactivation[5], are not observed in the rosette or RSCs.

Despite their primed chromatin landscape, WNT induces RSC-to-ESC reversion. This contrasts with the lack of reversion found in primed cells[21,44,45]. Epigenetic barriers such as DNA methylation and the repressive H3K9me3 histone mark enforce cell fate commitment[46–49]. The irreversibility of the primed state argues that MEK signals establish epigenetic barrier function in preparation for lineage specification. The erasure of H3K9me3 and 5meC and the accumulation of H3K27me3 in the rosette point towards unsuspected genome remodelling events, possibly clearing constitutive heterochromatin that would interfere with embryogenesis. In this light, it is suggestive that pericentric H3K27me3 maintains repression of the major satellites in ESCs when the H3K9me3 heterochromatin pathway is compromised[36,50]. H3K27me3 may act as a compensatory repressive mechanism in rosette-stage pluripotency until conversion to primed pluripotency. Further understanding of the ESC–RSC reversibility and RSC-primed transition may provide insight into the mechanisms of cell fate commitment and epigenetic barriers to cell reprogramming.

Recently, it was shown that L2i inhibits lumenogenesis in mouse embryos and maintains naive pluripotency, outlining a mechanism that coordinates the exit from naive pluripotency with amniotic cavity formation[3]. Our findings extend this seminal work by demonstrating that WNT signals link pluripotency progression to embryogenesis, not only by preventing rosette formation but also by preventing MEK signals from driving primed pluripotency progression. The rosette forms because of polarization of the ICM, which is induced by the deposition of a basal lamina by the primitive endoderm and the trophoblast[2]. The primitive endoderm differentiates from the ICM between E3.0 and E3.75 (refs. [51,52]). Interestingly, the mature primitive endoderm expresses the WNT antagonist DKK1 (refs. [53,54]) and may thereby signal its completion by repressing WNT signals, allowing *Otx2* induction, rosette formation and MEK-dependent pluripotency progression to proceed. Indeed, the pro-amniotic cavity is severely compromised in *Otx2* mutants[55], while mutants with constitutively activate β-catenin fail both to induce *Otx2* and form a pro-amniotic cavity[56,57]. Such a mechanism would constitute a checkpoint linking epiblast progression to maturation of the primitive endoderm by WNT signals.

Online content

Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at https://doi.org/10.1038/s41556-020-0508-x.

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Methods

Isolation of timed embryos. All animal experiments were conducted after approval by the Centrale Commissie Dierproeven (the Hague, the Netherlands) and by the Erasmus MC Animal Ethical Committee. Embryos were obtained from crosses between actin–GFP male[60] and superovulated C57Bl/6JolaHsd or FVB/N female mice. The male mice and newly weaned female mice were maintained under a shifted day–night rhythm (lights on at 1:00, lights off at 13:00). Superovulation was performed using an adjusted protocol: 5-week-old females received 0.1 ml of CARD HyperOva (Cosmo Bio) intraperitoneally at 8:00, and 48 h later, 5 units of 50 U ml^{-1} human chorionic gonadotropin. The superovulated females were housed with actin-GFP males after the human chorionic gonadotropin injection and checked for copulation plugs 3 and 6 h later. For isolation of peri-implantation and post-implantation embryos, isolated uteri were dissected under the lens of a fluorescence stereomicroscope to facilitate the detection of implanted GFP^{+} embryos. The embryos were then used in immunostaining, RNA isolation or culture protocols as described below.

The mouse *Otx2*–GFP strain, STOCK Tg(*Otx2*-EGFP)BD199Gsat/Mmnc, RRID:MMRRC_000239-UNC, was obtained from the Mutant Mouse Resource and Research Center (MMRRC) at Taconic/University at Albany, a NIH-funded strain repository, and was donated to the MMRRC by N. Heintz, The Rockefeller University. Embryos were obtained from mating *Otx2*–GFP males with superovulated females and isolation of the embryos at E1.5. Embryo culture was performed as described below.

Immunostaining and imaging of embryos. Embryos were fixed for 20 min in 4% paraformaldehyde (PFA), washed twice in PBS containing 0.05% Tween-20 (PBT) and permeabilized for 20 min in 0.3% Triton X-100 plus 0.1 M glycine in PBT. Samples were then washed three times in PBT and blocked for 1 h in blocking buffer consisting of 5% normal donkey serum plus 1% BSA in PBT. For 5mC and 5hmC staining, embryos were treated after permeabilization with 4 M HCl plus 0.1% Triton X-100 in water for 10 min, rinsed in PBT plus 20% blocking buffer and blocked as described above. The embryos were then incubated overnight at room temperature with primary antibodies in blocking buffer. The next day, embryos were washed three times in PBT, incubated for 2 h with secondary antibodies and again washed three times in PBT. If applicable, embryos were then incubated with 5 units per ml of phalloidin (labelled with Alexa-546 or Alexa-488) (ThermoFisher) in PBT for 20 min and washed in PBT. Embryos were then incubated with DAPI or, for 5mC or 5hmC staining, TO-PRO-3 in PBT for 10 min, rinsed in PBT and mounted on coverslips in Prolong Gold (ThermoFisher). The Prolong Gold was allowed to set for at least 1 day, after which embryos were imaged using a Leica SP5 confocal laser scanning microscope with 405-, 488-, 561-, 594- and 633-nm laser lines and ×40 or ×63 oil objectives. Images depicting pHC in embryos were deconvolved using Huygens Deconvolution software (Scientific Volume Imaging). Contrast settings were adjusted if necessary and equally over all relevant images using ImageJ. The antibodies used are listed in Supplementary Table 6.

Embryo culture. For culture of four-cell-stage embryos, embryos were isolated at E1.5 from superovulated females and cultured in KSOM medium (Millipore) until the four-cell stage, or in vitro fertilizations were performed as previously described[61] with modifications. In vitro fertilization culture medium, Mouse Vitro Fert (MVF; Cook Medical), was used for sperm incubation, in vitro fertilization and zygote culture. Straws with frozen sperm were thawed in a 37 °C water bath for 30 s. The sperm was pushed out of the straw into 500 µl of MVF and incubated for 11–15 min at 37 °C, 5% CO_2, 5% O_2. Meanwhile four to five superovulated 4-week-old female mice per fertilization drop were used as oocyte donors. After 4 h of co-incubation, the presumptive zygotes were washed three times in 50-µl MVF drops and those appearing normal were cultured overnight in a 50-µl MVF drop at 37 °C, 5% CO_2, 5% O_2. The next day, two-cell embryos were transferred into KSOM medium and cultured at 37 °C, 5% CO_2. Next, four-cell embryos were collected during a 10-h window and transferred into KSOM containing 2 µM IWP2 (Merck, 681671) or the same concentration of dimethylsulfoxide (DMSO) and cultured for 60 h. They were then fixed, immunostained and imaged as described above.

To culture blastocysts to the egg cylinder stage, morula-stage embryos (C57BL/6NRj, Janvier Quickblasto, or freshly isolated) or E4.0 blastocysts (see "Isolation of timed embryos") were cultured in KSOM medium until the expanded blastocyst stage. They were then transferred to 8-well iBidi plates containing IVC1 medium[62]. E4.0 blastocysts were cultured directly into iBidi plates containing IVC1. After attachment of the embryos, the medium was switched to IVC2 (ref. [62]), containing 1 µM PD as indicated. Embryos were then fixed, immunostained and imaged as described above.

Cell culture. Mouse ESCs were cultured on gelatine and FCS-coated plates in N2B27 medium supplemented with 1,000 U ml^{-1} LIF, 3 µM CHIR and 1 µM PD (L2i ESCs). Coating was performed by first briefly incubating the wells with 0.1% gelatin in PBS, then incubating them with FCS for 1 h at 37 °C followed by a rinse with PBS. N2B27 medium consisted of one volume DMEM/F12 combined with one volume Neurobasal medium supplemented with 0.5% N2 supplement, 1% B27 supplement, 0.033% BSA 7.5% solution, 50 µM β-mercaptoethanol, 2 mM Glutamax, 100 U ml^{-1} penicillin and 100 µg ml^{-1} streptomycin (all from Invitrogen). Cells were cultured at 37 °C and 5% CO_2 and passaged 1:3–1:10 every 2–3 days by triturating the colonies into a single cell suspension using 0.05% trypsin–EDTA. Trypsin was quenched using soybean trypsin inhibitor (Sigma). Epiblast stem cells (EpiSCs) were cultured on gelatine and FCS-coated plates in N2B27 medium supplemented with 20 ng ml^{-1} activin A, 12 ng ml^{-1} bFGF (both Peprotech) and 2 µM IWP2 to suppress spontaneous differentiation[18]. EpiSCs were passaged 1:4–1:10 every 3 days by triturating the colonies into small clumps using 0.5 mg ml^{-1} collagenase IV (Sigma). All cell lines in culture were tested for mycoplasma contamination every 2–3 months.

To transit from embryonic stem to MEK/WNT-inhibited or primed cells, trypsinized ESCs were seeded onto gelatine and FCS-coated plates in N2B27 medium supplemented with 1,000 U ml^{-1} LIF, 2 µM IWP2 and either 1 µM PD for MEK/WNT-inhibited cells or 20 ng ml^{-1} activin A and 12 ng ml^{-1} bFGF for primed cells. After 2 days, MEK/WNT-inhibited cells were passaged as ESCs, whereas primed cells were passaged as EpiSCs.

The following ESC lines were used (see Supplementary Table 6): R1 and CGR8 (both obtained from Stanford Transgenic Facility), IB10, SV7, FN3 and SV8 (all described in ref. [6]; SV8 is female and FN3 is from the non-permissive FVB/N background), RGd2 (ref. [21]), *Tcf7l1*$^{-/-}$ (ref. [9]), *Tbx3*–RFP;*Oct6*–GFP[23], *Otx2*KO and *Otx2*OE[25]. The following RSC lines were derived from blastocysts in this study (see below): 129LIMA2, 129LIMB1, 129LIMB3 (all male) and 129LIMA6 (female). For the generation of TRE–*Otx2* ESCs, a mouse *Otx2* complementary DNA (accession number BC017609) was amplified using primers incorporating an amino-terminal Kozak sequence and SalI and NotI restriction sites. The vector p2Lox.GFP[63] was digested with SalI and NotI to remove the GFP sequence, and the SalI/NotI-digested *Otx2* fragment ligated into its place to create p2Lox.*Otx2*. A total of 25 µg of p2Lox.*Otx2* was electroporated into A2Lox.Cre ESCs, followed by induction of the cells with 1 µg ml^{-1} doxycycline overnight to induce recombination, and the cells selected on 250 µg ml^{-1} of G418, as previously described[64].

IWP2 (Merck, 681671), CHIR (Axon Medchem), PD (Merck) were diluted from 2 mM (IWP2) or 10 mM stocks in DMSO and used at 2, 3 and 1 µM, respectively. WNT3A protein was used at 400 ng ml^{-1}. Recombinant mouse WNT3A protein was produced in *Drosophila* S2 cells grown in suspension culture (a gift from R. Nusse, Stanford University) and purified using Blue Sepharose affinity and gel filtration chromatography[65]. For this, the S2 cells were expanded in Schneider's *Drosophila* medium (Lonza) containing 10% fetal bovine serum and antibiotics, and media were collected when cell expansion reached a plateau. Up to 12 litres of conditioned medium was 0.45-µm filtered, adjusted to 1% Triton X-100 and then applied to a fast protein liquid chromatography column containing 200 ml Blue Sepharose 6 Fast Flow (GE Healthcare, 17094801). After washing with four volumes of washing buffer (150 mM KCl, 20 mM Tris-HCl, 1% CHAPS, pH 7.5), bound proteins were eluted with elution buffer (1.5 M KCl, 20 mM Tris-HCl, 1% CHAPS, pH 7.5). After analysis on a Coomassie gel, WNT3A-containing fractions were selected, combined and concentrated to 10 ml using Pierce Concentrators 20 K MWCO (Thermo Scientific, 89887A). The combined fractions were fractionated on a HiLoad 26/60 Superdex 200 gel filtration column (GE Healthcare) in PBS, 0.5 M NaCl, 1% CHAPS, pH 7.3. Fractions were analysed for purity and WNT activity by Coomassie gels and WNT activity reporter assays, respectively, and selected fractions were combined for use.

Derivation and blastocyst injection of RSCs. E3.5 blastocysts were obtained from crosses between 129S2/SvHsd males and C57Bl/6JolaHsd females (Envigo). Blastocysts were flushed from the uteri and placed intact in 96-wells plates coated with gelatine and FCS and filled with 100 µl N2B27 medium plus LIF. Late next day, when hatching and attachment of the embryos had occurred, an equal volume of medium containing 4 µM IWP2 and 2 µM PD was added. Embryo outgrowths were trypsinized 4 days later, the trypsin quenched with soybean trypsin inhibitor, and plated in 96-wells plates coated with gelatine and FCS in N2B27 containing LIF, 2 µM IWP2 and 1 µM PD (RSC medium). After 4 days, wells containing colonies were passaged and expanded further in RSC medium. RSCs were recovered by treatment with 0.05% trypsin–EDTA and 10–15 cells were injected into blastocysts collected from C57Bl/6 mice. Blastocysts (10–15 per mouse) were transferred into the uterus of a pseudo pregnant mouse. The percentage of coat colour chimerism of the chimeras was visually estimated. Chimeras were mated with C57Bl/6 mice to determine germ line transmission.

Xist fluorescent in situ hybridization (FISH). For Xist RNA FISH and immunofluorescence staining, cells were adsorbed to poly-lysinated coverslips. Cells were fixed for 10 min with 4% PFA in PBS at room temperature, washed with 70% ethanol, permeabilized for 4 min with 0.2% pepsin at 37 °C and post-fixed with 4% PFA in PBS for 5 min at room temperature. Coverslips were washed twice with PBS and dehydrated in a gradient of 70%, 90% and 100% ethanol. A nick-labelled DNA probe (digoxigenin) was dissolved in hybridization mixture (50% formamide, 2XSSC (1XSSC: 0.15 M NaCl, 0.015 M sodium citrate), 50 mM phosphate buffer (pH 7.0) and 10% dextran sulfate) and 100 ng µl^{-1} mouse Cot DNA to a final concentration of 1 ng µl^{-1}. Probe was denatured for 5 min, prehybridized for 45 min at 37 °C, and coverslips were incubated overnight in a humid chamber at 37 °C. After hybridization, coverslips were washed once

in 2XSSC, three times in 50% formamide–2XSSC, both at 37 °C, and twice in TST (0.1 M Tris, 0.15 M NaCl, 0.05% Tween 20) at room temperature. Blocking was done in BSA–TST for 30 min at room temperature. Detection was done via subsequent steps of incubation with anti-digoxigenin (Boehringer) and two fluorescein isothiocyanate (FITC)-labelled antibodies for Xist RNA detection in blocking buffer for 30 min at room temperature. Coverslips were washed twice with TST between detection steps and once finally with TS (0.1 M Tris, 0.15 M NaCl). Dehydrated coverslips were mounted with ProLong Gold Antifade with DAPI (Molecular Probes).

Immunohistochemistry of cells. Cells were passaged onto gelatine- and FCS-coated glass slides 1 day before fixation. Cells were fixed with 4% PFA in PBS at room temperature for 10 min, washed with PBS containing 0.05% Tween 20 (PBST), permeabilized with 0.3% Triton X-100 plus 0.1 M glycine in PBST, washed with PBST, incubated in blocking buffer for 1 h (1% BSA, 5% donkey serum in PBST) and incubated overnight at room temperature in blocking buffer with the primary antibodies. The next day, slides were washed with PBST before incubation for 2 h with the secondary antibodies in blocking buffer. Nuclei were then stained with DAPI (Molecular Probes) in PBST for 5 min at room temperature, and slides mounted using Prolong Gold. Immunofluorescence confocal images were acquired using a Leica SP5 inverted laser-scanning confocal microscope. Contrast settings were adjusted if necessary and equally over all relevant images using ImageJ. The antibodies used are listed in Supplementary Table 6.

For alkaline phosphatase staining, cells were stained according to the manufacturer's protocol (Alkaline Phosphatase Detection kit SCR004, Millipore).

In vitro rosette and lumen assays. ESCs were triturated to a single cell suspension using 0.05% trypsin–EDTA (Life Technologies) and washed with PBS. Cells were resuspended in ice-cold BME (BME 2 RGF, AMSBIO, 3533-010-02) at a density of 5,000 cells per µl. Drops of 15 µl were plated on coverslips or in a prewarmed 8-chamber slide (Thermo Scientific, 154534) and incubated for 5 min at 37 °C until the BME solidified. The plate was then filled with prewarmed medium consisting of DMEM plus 15% FCS, 1× MEM nonessential amino acids, 50 µM β-mercaptoethanol, 100 U ml^{-1} penicillin and 100 µg ml^{-1} streptomycin (all from Life Technologies), containing supplements as indicated, and cultured at 37 °C and 5% CO_2. The cells and BME drops were then fixed with 4% PFA, prewarmed to 37 °C, for 30 min. For phalloidin staining, the fixed cells were rinsed with PBS, incubated with phalloidin–Alexa-488 (5 units per ml, Invitrogen) and DAPI in PBS for 1.5 h, and washed three times with PBS. For immunostaining, the fixed cells were permeabilized with 0.3% Triton/0.1 M glycine in PBS for 30 min at room temperature, washed with PBS containing 0.05% Tween 20 (PBST) for 20 min, incubated in blocking buffer for 1 h (1% BSA, 5% donkey serum in PBST) and incubated overnight at room temperature in blocking buffer with the primary antibodies. The next day, cells were washed three times for 30 min with PBST before incubation for 2 h with the secondary antibodies in blocking buffer. After three washes with PBST for 30 min, nuclei were stained with DAPI (Molecular Probes) for 30 min at room temperature and washed three times with PBS. For lumen assays, the live cultures were incubated for 1 h in a cell culture incubator with culture medium containing 1:500 anti-PODXL antibody (MAB1556, R&D Systems), followed by fixation and staining with secondary antibody as described above. The live staining ensured that only PODXL exposed at the cell exteriors was stained. The stained preparations were covered with Prolong Gold and imaged using a Zeiss Axioplan microscope or a Leica SP5 inverted laser-scanning confocal microscope. Structures were designated as rosette if they showed polarized, central localization of phalloidin or intracellular PODXL. Structures displaying central localization of PODXL after live staining were designated as lumen. The antibodies used are listed in Supplementary Table 6.

RNA-seq and data analysis. Total RNA was prepared using TriReagent (Sigma) according to the manufacturer's protocol. RNA-seq was performed at the Erasmus MC Center for Biomics (Rotterdam, the Netherlands). The RNA-seq library was prepared for analysis according to the Illumina TruSeq stranded mRNA protocol (www.illumina.com). Briefly, 200 ng of total RNA was purified using poly-T oligo-attached magnetic beads to result in poly-A containing mRNA. The poly-A tailed mRNA was fragmented, and cDNA was synthesized using SuperScript II and random primers in the presence of actinomycin D. cDNA fragments were end-repaired, purified with AMPure XP beads, and A-tailed using Klenow exo-enzyme in the presence of dATP. Paired-end adapters with dual index (Illumina) were ligated to the A-tailed cDNA fragments and purified using AMPure XP beads. The resulting adapter-modified cDNA fragments were enriched by PCR using Phusion polymerase as follows: 30 s at 98 °C, 15 cycles of (10 s at 98 °C, 30 s at 60 °C, 30 s at 72 °C), 5 min at 72 °C. PCR products were purified using AMPure XP beads and eluted in 30 µl of resuspension buffer. One microliter was loaded on an Agilent Technologies 2100 Bioanalyzer using a DNA 1000 assay to determine the library concentration and for quality checking. Cluster generation was then performed according to the Illumina TruSeq SR Rapid Cluster kit v2 (cBot) reagents preparation guide (www.illumina.com). Briefly, six RNA-seq libraries were pooled together to get a stock of 10 nM. One microlitre of the 10 nM stock was denatured with NaOH, diluted to 6 pM and hybridized onto the flow cell. The hybridized products were sequentially amplified, linearized and end-blocked according to the Illumina Single Read Multiplex Sequencing user guide. After hybridization of the sequencing primer, sequencing-by-synthesis was performed using the HiSeq 2500 with a single-read 50-cycle protocol followed by dual index sequencing.

Base calling and demultiplexing was performed using Illumina bcl2fastq v.2.19. Sequencing reads were aligned against GRCm38 (mm10) using HiSat2 v.2.0.4. Count data files were created using htseq-count v.0.6.0. Additional embryo and L2i ESC data[3,16] were downloaded from ArrayExpress E-MTAB-2958 and E-MTAB-5147. E-MTAB-2958 was aligned against GRCm38 and count data files created as described above, whereas count data files were already available from E-MTAB-5147.

Normalization and differential expression analysis were performed using R v.3.4.3 and the DESeq2 Bioconductor package v.1.18.1 with betaPrior set to TRUE, a False Data Rate cut-off alpha value of 0.1, and a log-fold change threshold lfcThreshold of 1. Heatmaps and PCA plots were generated from rlog-transformed data using the pheatmap package 1.0.8 and the DESeq2 function plotPCA, respectively, using the 500 genes displaying the highest row variance for PCA.

Significant differentially expressed genes were analysed for GO Biological Process terms enrichment using GOrilla[58], using a P value threshold of 0.001 and a background list of all genes that had an average read count of at least 1 over all samples. GO analysis was further processed using REVIGO[59].

Single-cell sequencing. Single-cell mRNA sequencing was performed according to an adapted version of the SORT-seq protocol[66]. In brief, single cells were FACS-sorted into 384-well plates containing 384 primers and mineral oil (Sigma). After sorting, plates were snap-frozen on dry ice and stored at −80 °C. For amplification, cells were heat-lysed at 65 °C followed by cDNA synthesis using the CEL-Seq2 protocol[67] and robotic liquid-handling platforms. After second-strand cDNA synthesis, the barcoded material was pooled into libraries of 384 cells and amplified using in vitro transcription. Following amplification, the rest of the CEL-seq2 protocol was followed for preparation of the amplified cDNA library using TruSeq small RNA primers (Illumina). The DNA library was paired-end sequenced on an Illumina Nextseq 500, high output, with a 1 × 75 bp Illumina kit (read 1: 26 cycles, index read: 6 cycles, read 2: 60 cycles).

During sequencing, read 1 was assigned 26 bp and was used for identification of the Illumina library barcode, cell barcode and unique molecular identifier. Read 2 was assigned 60 bp and used to map to the reference transcriptome of mm10 with Burrows–Wheeler Aligner. Data demultiplexing and statistical UMI poisson counting correction was done as previously described[68]. Mapping and generation of count tables was performed using the MapAndGo script (https://github.com/anna-alemany/transcriptomics/tree/master/mapandgo). Single-cell transcriptomics analysis was done using the RaceID3 algorithm, following an adapted version of the RaceID manual (https://cran.r-project.org/web/packages/RaceID/vignettes/RaceID.html). In total, 1,152 cells were sequenced. After removing cells with fewer than 8,000 UMIs and only keeping genes that were detected with at least 3 UMIs in at least 2 cells, 938 cells and 11,070 genes were left for further analysis. During clustering, genes contributing to batch effects (*Rn45s*, *Kcnq1ot1*, *Malat1*, A630089N07Rik, *Rpl13*, *Rpl39* and *Rps7*) were excluded from clustering, but were kept in the dataset for downstream visualization and analysis. Clustering was done using hclust. Differential gene expression analysis was done as previously described[66] using an adapted version of the DESeq2 algorithm[69].

Flow cytometry. For flow cytometry experiments, single-cell suspensions were made using trypsin–EDTA (5 min in 37 °C) or Enzyme-Free Cell Dissociation Buffer (Gibco) for 30 min at 37 °C and washed with 5% FCS in PBS. Hoechst 33258 (1 µg ml^{-1}) was used for live/dead cell assessment. Cells were analysed for GFP reporter gene expression using a FACS Fortessa (BD Biosciences), and data were analysed using FlowJo.

Live imaging. R1 L2i and MEK/WNT-inhibited cells were plated at clonal density on serum-coated or BME-coated plates and incubated overnight at 37 °C. The cells were tracked with CytoMate and pictures taken every minute. For the analysis, a stack of 12 pictures (taken at 5-min intervals, representing 1 h of tracking) was imported to ImageJ. Velocity was calculated, and the movement dotplots created using the 'Manual Tracking' plugin.

Real time RT–PCR analysis. Total RNA was prepared using a RNeasy mini or micro kit (Qiagen) according to the manufacturer's instructions. To extract RNA from embryos, four embryos from the same litter were combined, while other littermates were used for immunofluorescence to establish the presence of a rosette or lumen. Reverse transcription was performed using Superscript II (Invitrogen) or, for embryo RNA, a RevertAid RT Reverse Transcription kit (ThermoFisher, K1691). Quantitative PCR was carried out on a BioRad CFX machine using a SYBR-green-based master mix. Relative quantification was done using *Gapdh* as a reference gene or, for the analysis of embryo RNA, *Oct4* as a specific epiblast gene. All PCR assays were carried out in triplicate and the mean cycle threshold (CT) value was used for quantification. Primer sequences were designed such that they spanned splice junctions whenever possible, and sequences are provided in Supplementary Table 6.

Mass spectrometry of 5mC and 5hmC nucleosides. Mass spectrometry of 5mC and 5hmC was performed as previously described[70]. In short, for each sample, 2 µg of DNA (isolated from ~1×10^6 cells) was degraded into individual nucleosides using DNA Degradase Plus (Zymo Research). The individual nucleosides were measured using a high-performance liquid chromatography–tandem mass spectrometry (HPLC–MS/MS) system consisting of an Acquity UPLC (Waters) containing a Waters Atlantis Hilic column (2.1 mm × 100 mm, 3 µm) connected to a Micromass Quattro Premier XE (Waters). Quantification was performed using area-based linear regression curves derived from calibration standards containing internal standard solutions. The 5mC and 5hmC levels were calculated as a concentration percentage ratio of the percentage of 5-methyl-2′-deoxycytidine/2′-deoxyguanosine (%mdC/dG) and the percentage of 5-hydroxymethyl-2′-deoxycytidine/2′-deoxyguanosine (%hmdC/dG), respectively.

ChIP-seq analysis. R1 and CGR8 ESCs were cultured on gelatine and FCS-coated plates in N2B27 supplemented with LIF, 2 µM IWP2 and 1 µM PD for 8 days. Confluent cultures were fixed in PBS with 1% formaldehyde (Sigma) for 8 min and quenched using 100 mM fresh glycine for 15 min at room temperature. Fixed cell preparations were collected by scraping, then washed and sonicated in 1% SDS, 30 mM Tris pH 8.0, 150 mM NaCl using a Diagenode Bioruptor UCD-300 to a size range with a median of 200 bp. A total of 1 million cell equivalent chromatin was diluted seven times to 15 mM Tris pH 8.0, 1 mM EDTA, 150 mM NaCl, 1% Triton X-100 with fresh protease inhibitor cocktail (Roche) and 0.5 µg H3K27me3 or 1 µg H3K4me3 antibody (Diagenode) and incubated overnight at 4 °C with rotation. Protein A/G magnetic beads were washed in the same dilution buffer with 0.15% SDS and 0.1% BSA, added to the chromatin–antibody mix and rotated for 60 min at 4 °C. Beads were washed with 400 µl buffer for 5 min at 4 °C using (1) first a 150 mM NaCl wash buffer, (2) then twice a 500 mM NaCl buffer and (3) finally a 10 mM Tris buffer. DNA was eluted from the beads using 150 mM NaCl, 30 mM Tris, 1% SDS, 0.1 µg µl^{-1} Proteinase K buffer for 30 min at 55 °C followed by 2 h shaking at 65 °C. The beads were removed, after which the DNA was purified using a Qiaquick MinElute PCR purification kit (Qiagen) according to the manufacturer's protocol, and eluted in 20 µl of buffer EB.

Illumina library preparation was performed using a Kapa Hyper Prep kit using 5 ng of DNA isolated via ChIP. For end repair and A-tailing, double-stranded DNA was incubated with end repair and A-tailing buffer and enzyme and incubated first for 30 min at 20 °C and then for 30 min at 65 °C. Adapters were ligated by adding 30 µl ligation buffer, 10 µl Kapa DNA ligase, 5 µl 600 nM adaptor in a total volume of 110 µl and incubated for 15 min at 15 °C. Post-ligation clean-up was performed using Agencourt AMPure XP reagent, and products were eluted in 20 µl elution buffer. Libraries were amplified by adding 25 µl 2× KAPA HiFi Hotstart Ready-mix and 5 µl 10× Library Amplification Primer Mix followed by 10 PCR cycles. Samples were purified using a QIAquick MinElute PCR purification kit. Libraries were size- selected to ~300 bp using Agencourt Ampure XP beads. Correct size selection was confirmed using an Agilent BioAnalyzer. Sequencing was performed using an Illumina NextSeq500 with 42-bp read lengths. Sequencing of the input showed no specific signal.

Additional L2i[27] and EpiLC[35] H3K4me3 and H3K27me3 ChIP-seq data were downloaded from NCBI's Gene Expression Omnibus (GEO). The data were mapped against the mm9 reference genome using bowtie v.2.0.2. Duplicate and low-confidence mapping reads (mapq < 15) were removed. After sequencing and mapping of the ChIP-seq libraries, we normalized for the total number of mapped sequencing reads to allow quantitative comparisons between samples by generating reads per million mapped reads (RPM) values. The RPM values allow samples to be compared regardless of differences in sequencing depth. Peak calling was performed using SICER v.1.1 with window size 200, gap size 200 (for H3K4me3) or 600 (for H3K27me3) at an *E*-value of 0.1. Bivalency was demarcated using bedtools v.2.20.1 via the intersection of H3K4me3 and H3K27me3 replicate consistent peak-calls. Bedtools was used to collect read counts within peaks. DESeq2 v.1.16.1 was used for rlog transformation and normalization of the read counts. The data were clustered and visualized using R v.3.1.1, ngsplot v.2.61, bam2bw v.1.21 and the UCSC genome browser.

Preparation and immunostaining of chromosome spreads. Chromosome spreads were prepared as previously described[71]. Briefly, cells were treated with 0.1 µg ml^{-1} colcemid (Gibco) for 3 h. Cells arrested in mitotic prometaphase were collected by mitotic shake-off and spun down in culture medium containing colcemid for 10 min at 200 × *g*. Supernatant was removed to leave 1 ml of culture medium and cells were resuspended. An equal volume of a hypotonic buffer (pH 8.2) containing 30 mM Tris-HCl, 17 mM trisodium citrate dihydrate, 50 mM sucrose and 5 mM EDTA was added. After a 7 min of incubation, cells were spun down for 7 min at 200 × *g* and supernatant removed. Cells were resuspended in 100 mM sucrose (pH 8.2) to obtain a cell concentration of 5–15 × 10^6 cells per ml. From this suspension, 10 µl was applied to a slide dipped in a solution of 1% PFA and 0.15% Triton X-100 (pH 9.2). After drying for 1.5–2 h in a humid chamber, slides were washed with 0.08% Photo-Flo (Kodak Industrie), air dried and stored at −20 °C until further processing.

Immunofluorescence was performed as previously described[72]. The antibodies used are listed in Supplementary Table 6. Results from five independent experiments were evaluated by two observers, blinded to the culture condition and the type of modification detected. At least 100 metaphases were randomly selected and categorized per slide. The pHC region was identified by the DAPI-dense area between the centromeres and scored for absence or enrichment of the marker.

Statistics and reproducibility. No statistical methods were used to predetermine the sample sizes. The number and type of replicates and any statistical tests used are indicated in the figure legends, and all replicates successfully reproduced the presented findings. Most key experiments were repeated with cell lines and embryos from different genetic backgrounds (that is, from 129, C57Bl6 and FVB strains). Investigators were blinded to the experimental groups for analysis of in vitro rosette assays (Fig. 2j) and for analysis of chromosome spreads (Fig. 7b). Normality of data was tested using Kolmogorov–Smirnov test. Statistics were calculated using Graphpad Prism 8.3.0 or as indicated.

Reporting Summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability

Figures 3 and 4 have associated RNA-seq and single-cell RNA-seq data deposited in the GEO and are accessible through GEO series accession number GSE145727. Figure 6 has associated ChIP-seq data deposited in GEO with the accession number GSE112234 . Previously published RNA-seq data that were re-analysed here are available from ArrayExpress under accession codes E-MTAB-2958 and E-MTAB-5147. Previously published ChIP-seq data that were re-analysed here are available from the GEO under accession codes GSE23943 and GSE60204. Source data for Figs. 1, 2 and 4–8 and Extended Data Figs. 1, 3–6 and 8 are presented with the paper. All other data supporting the findings of this study are available from the corresponding author upon reasonable request.

Code availability

No custom computer code was used in this study.

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Author contributions

A.N., I.E., E.v.G., L.V., D.K., J.L., J.S., M.M. and D.t.B. designed and performed experiments and analysed data. A.N., I.E., D.K., J.L., J.S., R.A.M.D., G.v.M., M.M., M.D., E.B.B., H.M. and D.t.B. prepared the manuscript. J.S., D.t.B., R.A.M.D., G.v.M. and H.M. designed, performed and analysed the ChIP-seq experiments. D.t.B., J.S., A.T.d.D., R.W.W.B. and W.F.J.v.I. performed and analysed the RNA-seq experiments. E.v.G., D.t.B. and N.C.R. performed and analysed the single-cell sequencing experiments. A.N., C.E., E.v.G. and E.B.B. prepared and analysed the chromosome spreads. A.M. assisted with the mouse experiments and Y.G. performed additional experiments. H.M. and J.H.J. designed, performed and analysed nucleoside mass spectrometry analyses. D.t.B. conceived the study and coordinated the work.

Competing interests

The authors declare no competing interests.

Additional information

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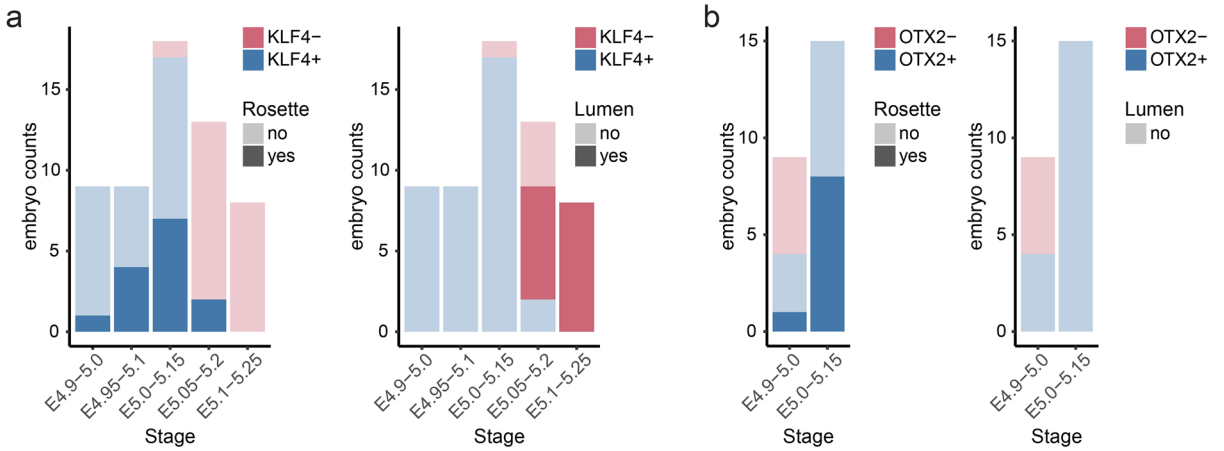
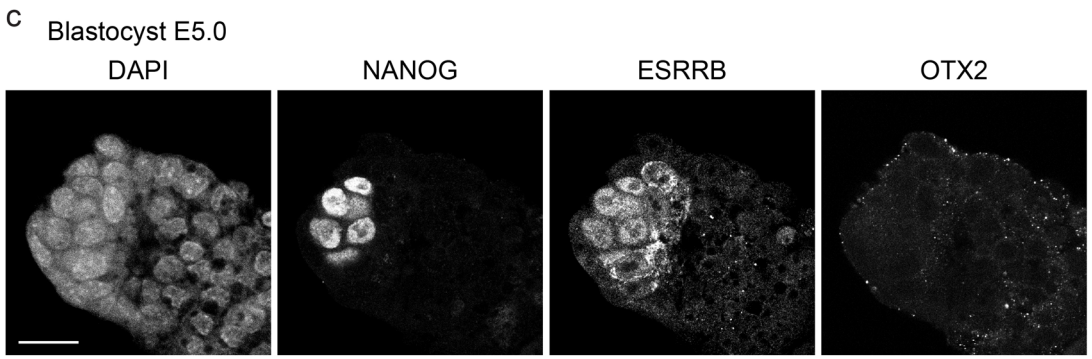
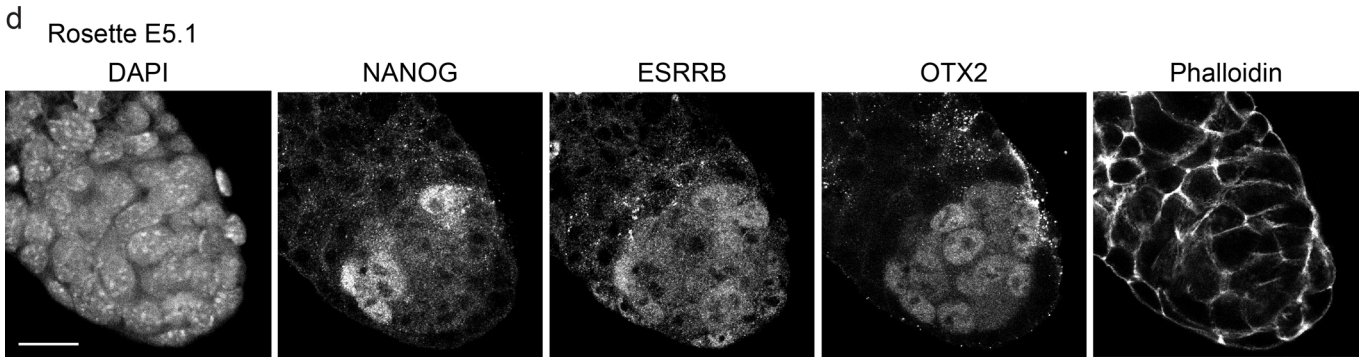
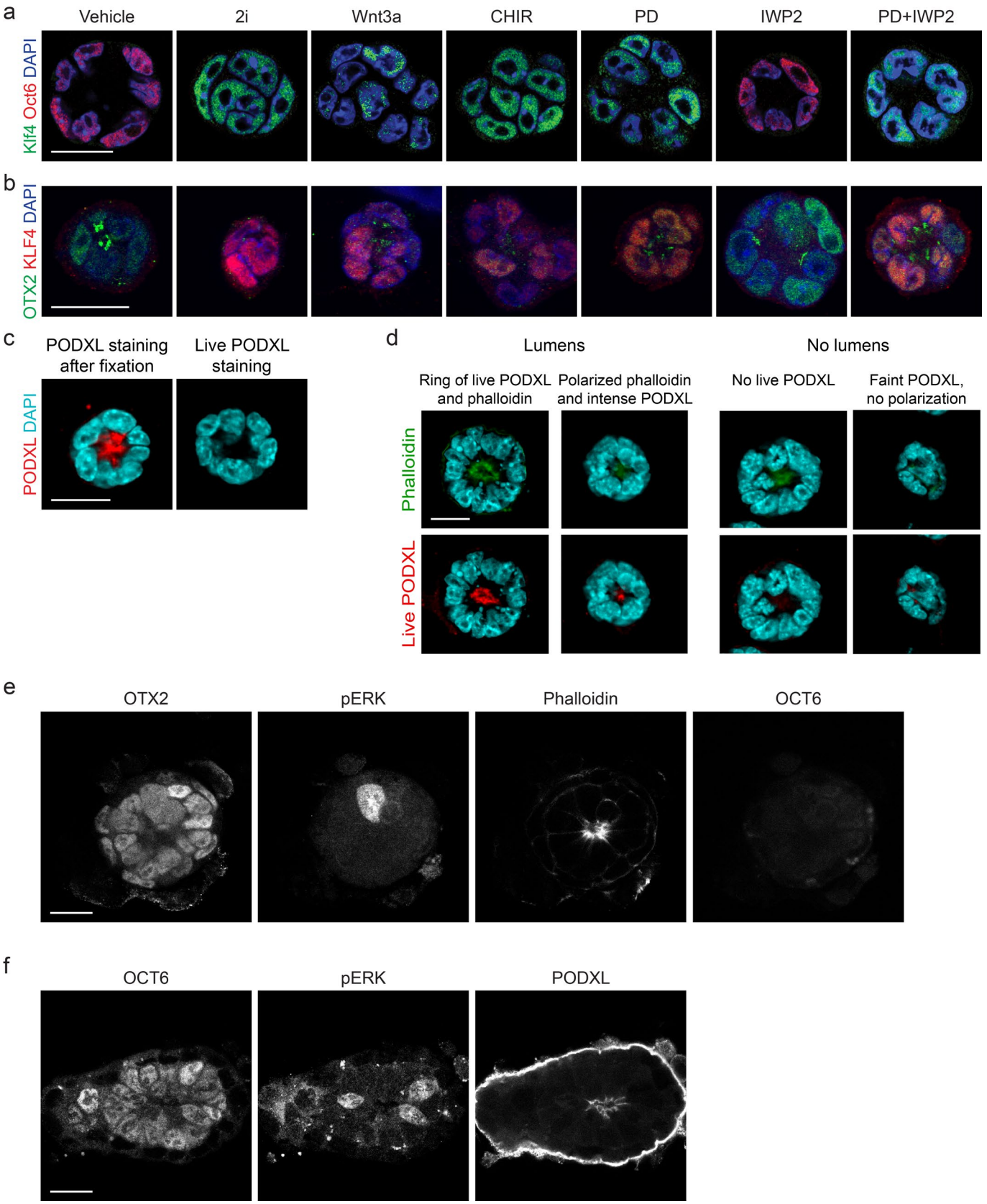




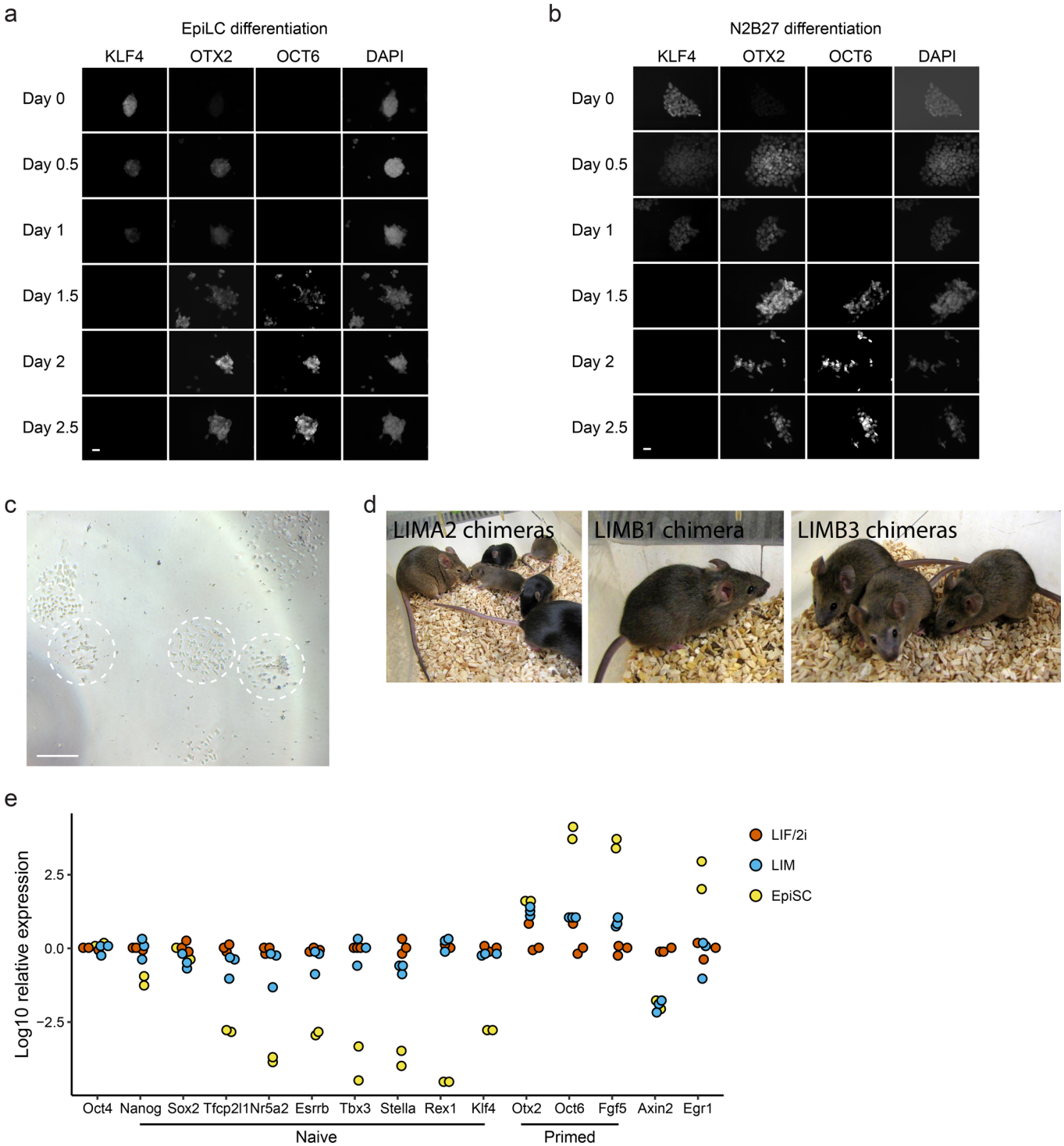


Extended Data Fig. 1 | Naive and primed markers in the embryonic rosette. a,**b,** Counts of embryos of the indicated stages displaying a rosette or lumen and KLF4 (**a**) or OTX2 (**b**) expression status. **a**, and (**b**) show different litters. **c**, Individual colour channels for Fig. 1h (representative of 5 experiments). **d**, Individual colour channels for Fig. 1i (representative of 8 experiments). Scale bars 20 μm.

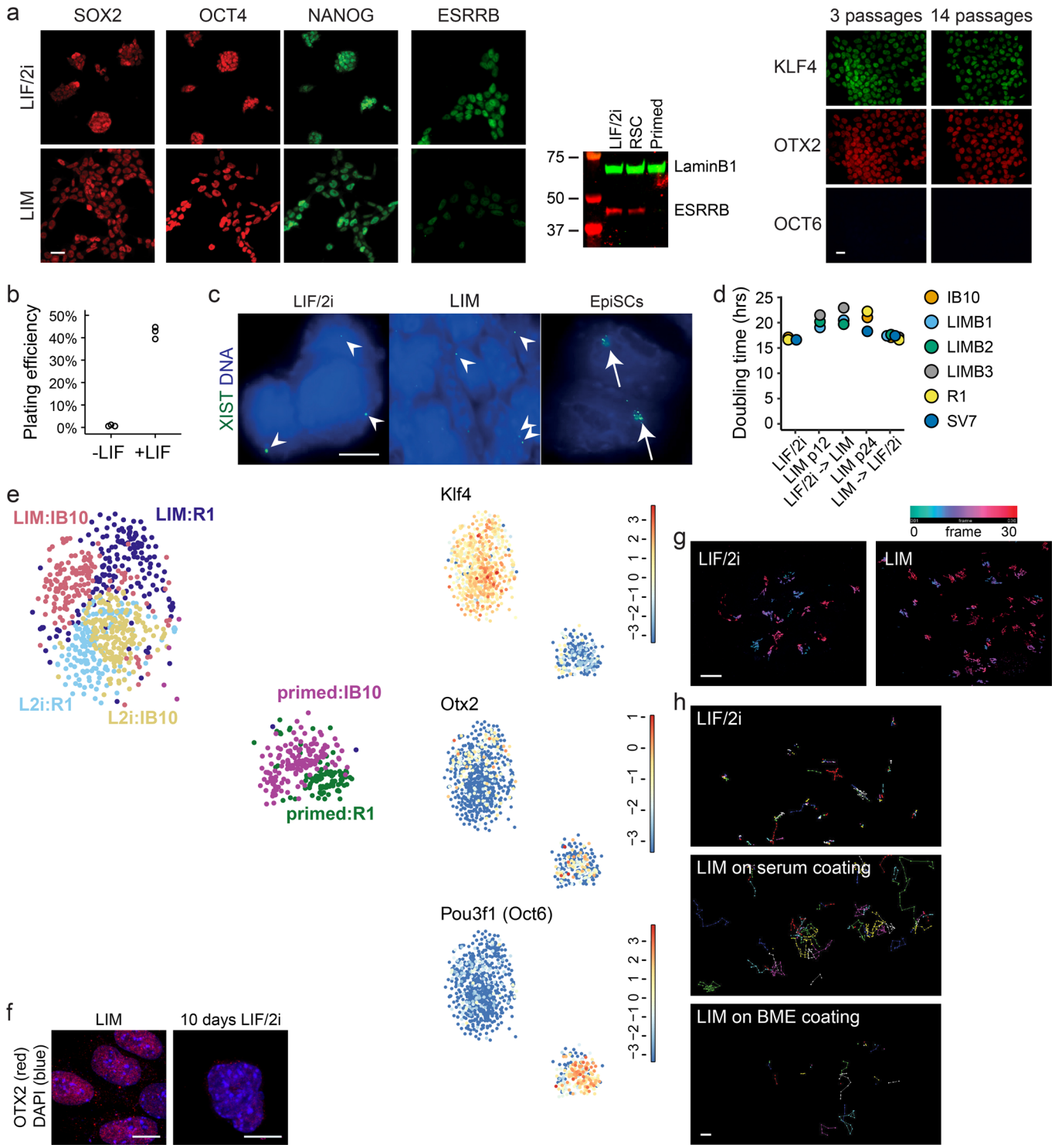


Extended Data Fig. 2 | See next page for caption.

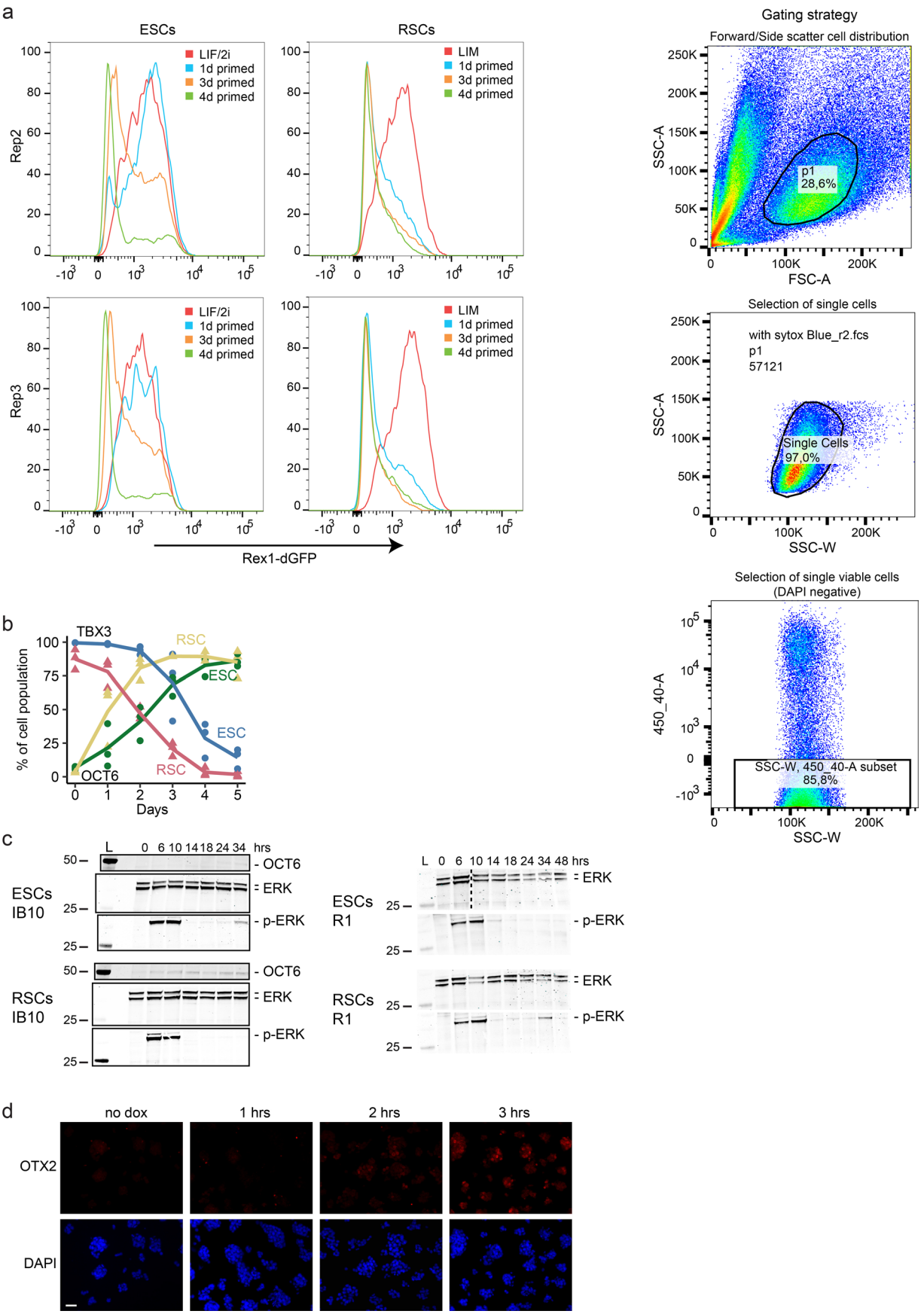
Extended Data Fig. 2 | WNT and MEK signals couple successive steps in morphogenesis to pluripotency progression. (**a**, **b**) Representative images of rosettes and aggregates generated by ESCs 48 hrs after seeding in BME in the indicated conditions and stained for the indicated markers. **a**, KLF4 (green) and OCT6 (red). **b**, OTX2 (green) and KLF4 (red). 3 independent experiments with similar results. **c**, Rosettes generated by ESCs 48 hrs after seeding in BME and stained for PODXL live or after fixation and permeabilization. Permeabilized samples showed PODXL staining, incorrectly suggesting presence of a lumen, while live staining revealed that the PODXL was not exposed at the cell surface and no lumen was actually present (illustrative examples, repeated > 5 times). **d**, Examples to illustrate criteria for calling ESC aggregates displaying a lumen, using live PODXL staining combined with phalloidin to reveal polarization (illustrative examples, repeated > 5 times). **e**, Individual colour channels for Fig. 2n (representative of 3 experiments). **f**, Individual colour channels for Fig. 2o (representative of 4 experiments). Scale bars 20 μm (a–f).



Extended Data Fig. 3 | WNT controls the initial phase of the naive-primed transition. (**a**, **b**) Immunostaining for the indicated markers of ESCs plated in EpiLC conditions (**a**) or N2B27 medium (**b**) for the indicated number of days (single experiments). **c**, Dispersed colonies (dashed white circles) obtained after passaging of a blastocyst grown out in the presence of IWP2, MEK inhibitor and LIF (LIM). 15 from 18 and 12 from 13 embryos yielded cell lines. **d**, Chimeras (brown), obtained after blastocyst injection of newly derived LIM cell lines (LIMA2, LIMB1 and LIMB3), with mate (black) and pups (only for LIMA2). Brown pups indicate that germline transmission occurred. **e**, RT-PCR analysis of ESCs maintained for 6 passages in L2i or MEK/WNT-inhibited conditions and of EpiSCs. n = 3 (LIF/2i and LIM) or n = 2 (EpiSC) biological replicates. Scale bars 20 µm (**a**,**b**), 100 µm (**c**).

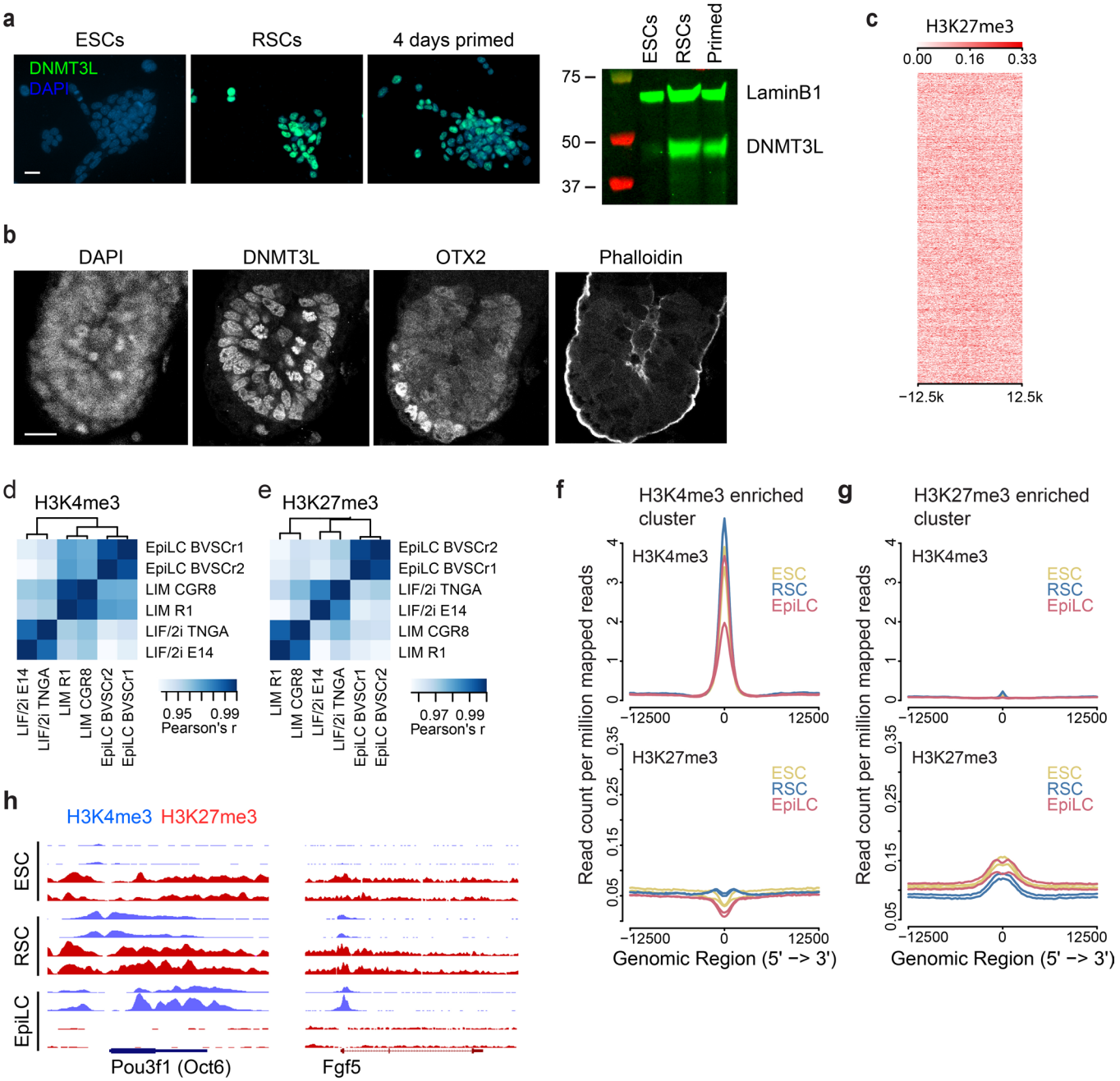


Extended Data Fig. 4 | MEK/WNT-inhibition maintains rosette characteristics. a, Immunostainings and western blot for the indicated markers of R1 ESCs maintained in L2i or LIM for 6 passages or as indicated (3 independent repeats with similar results; western blot 1 experiment, full blot in Source Data). **b**, Plating efficiencies of LIM cells in the presence and absence of LIF (n = 3 biological replicates, R1, FN3 and IB10 cells). **c**, Female SV8 L2i ESCs, LIM cells and EpiSCs stained for Xist by RNA-FISH. Puncta indicating Xist expression (arrowheads) are visible in both L2i and LIM cells, while only EpiSCs display Xist clouds indicating X-chromosome inactivation (arrows) (two independent experiments). **d**, Doubling times of the indicated cell lines in the indicated conditions. p12 and p24 indicate passage numbers of the cell lines. **e**, t-SNE plot of single cell RNA-Seq data, and log2 expression levels of the indicated genes plotted into the t-SNE plot. n = 159, 178, 102 (R1) and 184, 149 and 164 (IB10) cells for L2i, LIM and primed, respectively. **f**, RSCs were cultured for 10 days in the indicated conditions and immunostained for OTX2 (red). Representative images from 3 independent replicates. **g**, To determine apicobasal polarity, the Golgi apparatus was marked by Gm130 antibody staining, imaged by confocal microscopy and colorized according to z-stack level. Basal side of the cells correspond to low z-levels (green), and apical side to high levels (red). The data show apical Golgi localization in LIM cells, indicating their apicobasal polarity. Three experiments with similar results. **h**, Live tracking of individual cell locations during 24 hrs of L2i ESCs, LIM cells on serum-coating, and LIM cells on BME-coating. 80, 47 and 20 cells, respectively. Representative of 4 experiments. Scale bars 20 μm (a,g), 10 μm (c,f), 100 μm (h).

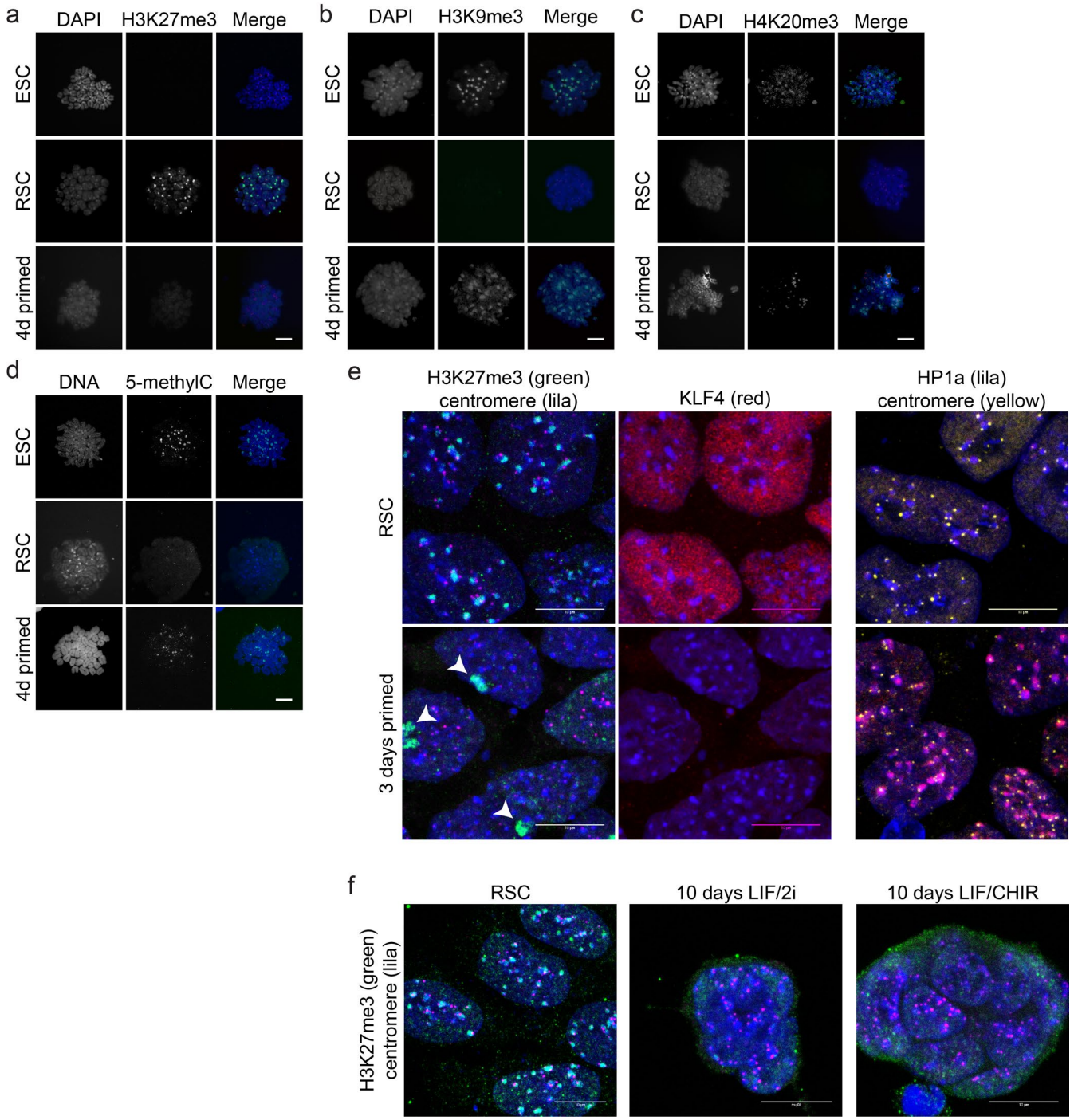


Extended Data Fig. 5 | See next page for caption.

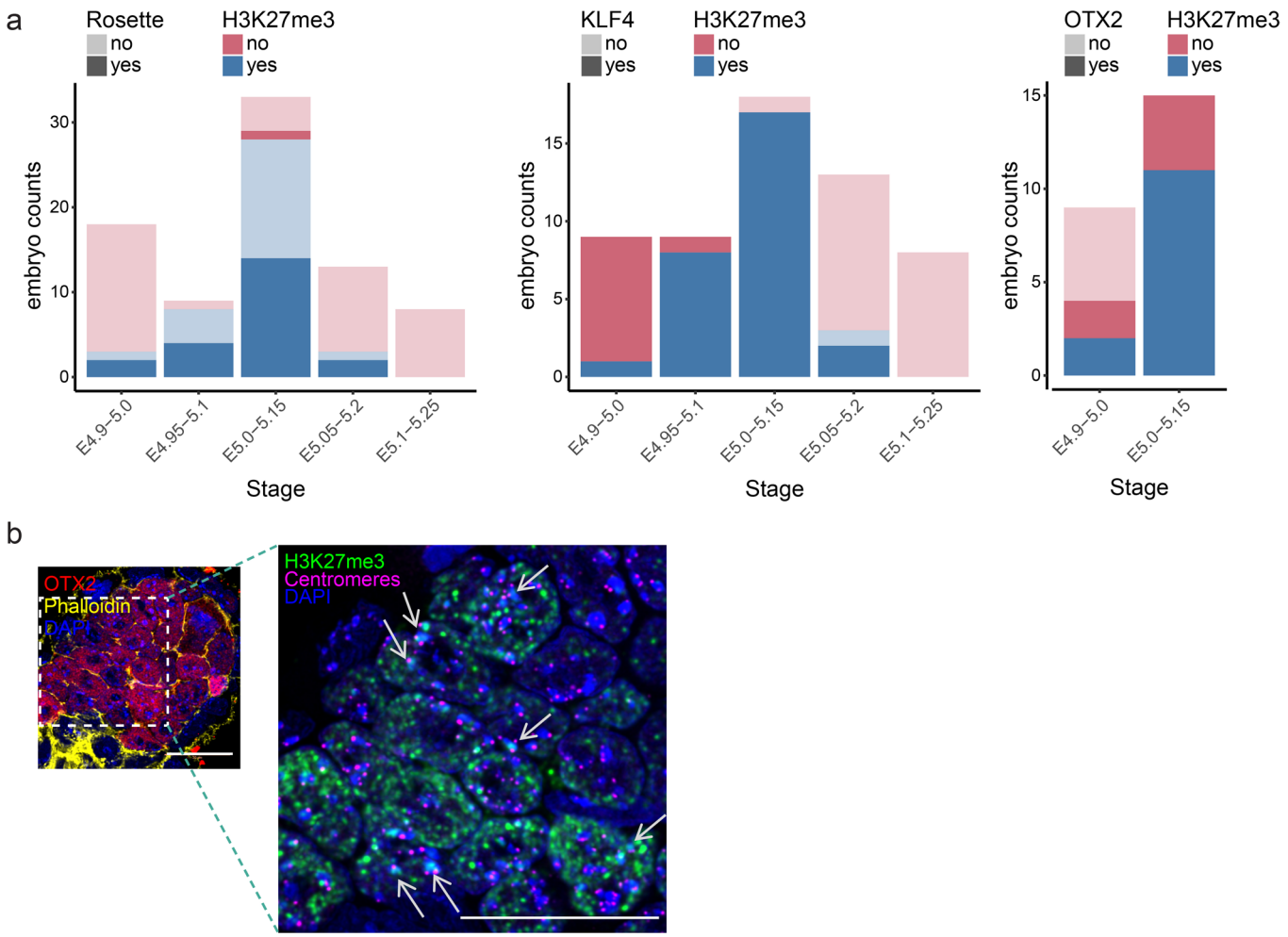
Extended Data Fig. 5 | Otx2 enables MEK-driven transition to primed pluripotency. a, RGd2 ESCs and RSCs were plated in primed conditions and analysed at several time points for GFP by flow cytometry. n = 3 independent replicates (3rd is shown in main Fig. 5a). **b**, ESCs and RSCs, carrying the Tbx3-RFP;Oct6-GFP reporters, were plated in primed conditions and analysed daily for expression of the reporters. Lines go through means, n = 3 independent experiments. **c**, Western blots showing OCT6 (only for IB10), ERK and phospho-ERK in IB10 and R1 ESCs and RSCs upon transfer to primed medium. Two (for OCT6) and three biological replicates. Full blots in Source Data. **d**, Immunostaining for OTX2 in TRE-Otx2 ESCs following induction by doxycycline for the indicated duration (single experiment). Scale bar 40 μm.



Extended Data Fig. 6 | A primed chromatin landscape in rosette-like stem cells. a, Immunofluorescence and western blot analysis for DNMT3L expression in the indicated cell types (3 independent repeats with similar results; western blot 1 experiment). **b**, Individual colour channels for Fig. 6b, right panel. Representative of 5 experiments. **c**, Representative input plot for H3K27me3 ChIP-Seq for the bivalent cluster. (**d**,**e**) H3K4me3 (d) or H3K27me3 (**e**) ChIP-Seq correlogram of ESCs, RSCs and EpiLCs. Two biological repeats for each condition. (**f**, **g**) Average intensity plots of H3K4me3 and H3K27me3 in the H3K4me3 (f) or H3K27me3 (g) enriched cluster. Two biological repeats for each condition. **h**, Screenshots from the UCSC genome browser showing H3K4me3 (blue) and H3K27me3 (red) peaks on the *Pou3f1* and *Fgf5* genes. Scale bars 20 μm (a,b).



Extended Data Fig. 7 | Rosette-like stem cells remodel pericentric heterochromatin. (**a–d**) Metaphase chromosome spreads of ESCs, RSCs and primed cells stained for the indicated markers. DAPI (blue), centromeres (red). **e**, Female RSCs (LIMA6) were cultured for 3 days in primed conditions and immunostained for the indicated markers. Arrowheads indicate H3K27me3-positive inactive X chromosomes demonstrating transition to primed pluripotency. **f**, RSCs were cultured for 10 days in the indicated conditions and immunostained for the indicated markers. Five (a–d) or three (e, f) biological replicates. Scale bars 20 μm (a-d), 10 μm (e, f).



Extended Data Fig. 8 | Rosette-specific formatting of pericentric heterochromatin. a, Counts of embryos of the indicated stages displaying pericentric H3K27me3 accumulation, and in which the presence of a rosette, KLF4 or OTX2 expression is detected. Embryos are the same as those of Extended Data Fig. 1a, b. **b**, Confocal microscopy images of E5.1 embryo stained for H3K27me3 (green), centromeres (magenta), OTX2 (red), and DAPI (blue). Some H3K27me3-positive pHC foci are indicated by white arrows (light blue, overlap of H3K27me3 and DAPI). Representative example from the embryos analysed in Extended Data Fig. 8a. Scale bars 20 µm.