

STEM CELLS AND REGENERATION

RESEARCH ARTICLE

H3K9 tri-methylation at *Nanog* times differentiation commitment and enables the acquisition of primitive endoderm fate

Agnès Dubois¹, Loris Vincenti¹, Almira Chervova¹, Maxim V. C. Greenberg^{2,3}, Sandrine Vandormael-Pournin¹, Déborah Bourc'h², Michel Cohen-Tannoudji¹ and Pablo Navarro^{1,*}

ABSTRACT

Mouse embryonic stem cells have an inherent propensity to explore gene regulatory states associated with either self-renewal or differentiation. This property depends on ERK, which downregulates pluripotency genes such as *Nanog*. Here, we aimed at identifying repressive histone modifications that would mark *Nanog* for inactivation in response to ERK activity. We found that the transcription factor ZFP57, which binds methylated DNA to nucleate heterochromatin, is recruited upstream of *Nanog*, within a region enriched for histone H3 lysine 9 tri-methylation (H3K9me3). Whereas before differentiation H3K9me3 at *Nanog* depends on ERK, in somatic cells it becomes independent of ERK. Moreover, the loss of H3K9me3 at *Nanog*, induced by deleting the region or by knocking out DNA methyltransferases or *Zfp57*, is associated with reduced heterogeneity of NANOG, delayed commitment into differentiation and impaired ability to acquire a primitive endoderm fate. Hence, a network axis centred on DNA methylation, ZFP57 and H3K9me3 links *Nanog* regulation to ERK activity for the timely establishment of new cell identities. We suggest that establishment of irreversible H3K9me3 at specific master regulators allows the acquisition of particular cell fates during differentiation.

KEY WORDS: ERK, H3K9me3, *Nanog*, ZFP57, Heterogeneity, Primitive endoderm, Mouse

INTRODUCTION

Mouse embryonic stem cells (ESCs) are derived from pre-implantation embryos and recapitulate numerous properties of the pluripotent inner cell mass of the blastocyst (Martello and Smith, 2014). *In vivo*, the culmination of pluripotency – the ability to give rise to all three somatic germ layers – takes place when the primitive endoderm – a source of extra-embryonic tissues – segregates from the epiblast, the founder of the embryo proper (Chazaud and Yamanaka, 2016). This segregation is strictly controlled by the transcription factor *Nanog*, which is required to form the epiblast

(Mitsui et al., 2003; Silva et al., 2009) and, additionally, stimulates FGF4 production. This extracellular signal is then transduced into neighbouring cells by ERK (MAPK1) activity, silencing *Nanog* and opening a window of opportunity to undergo commitment into primitive endoderm differentiation (Chazaud et al., 2006; Nichols et al., 2009; Yamanaka et al., 2010; Frankenberg et al., 2011; Saiz et al., 2016; Bessonnard et al., 2017). This binary cell-fate decision is characterised by substantial heterogeneity of NANOG expression, which creates the conditions required for epiblast and primitive endoderm specification (Chazaud et al., 2006). Subsequently, *Nanog* expression is downregulated in the epiblast, eliciting the establishment of somatic differentiation (Chambers et al., 2003). *In vitro*, ESCs also exhibit extensive *Nanog* heterogeneity, characterised by a subpopulation expressing no or extremely low levels of NANOG. Whereas NANOG-positive cells self-renew efficiently, NANOG-negative cells exhibit a propensity to differentiate even though they remain uncommitted and can spontaneously revert to the *Nanog*-expressing state (Chambers et al., 2007; Singh et al., 2007; Kalmar et al., 2009; Canham et al., 2010; Abranches et al., 2014). Notably, NANOG-negative cells spontaneously generated in culture or by homozygous targeted deletion show increased differentiation propensity towards both primitive endoderm and somatic fates (Chambers et al., 2007; Singh et al., 2007; Kalmar et al., 2009; Canham et al., 2010; Abranches et al., 2014).

NANOG heterogeneity has been proposed to result from two mechanisms: (1) from the architecture and the topology of the pluripotency network (Navarro et al., 2012; Karwacki-Neisius et al., 2013), and (2) from extrinsic cues, such as LIF/STAT3, WNT/GSK3b and FGF/ERK signalling (Kalmar et al., 2009; Wray et al., 2011; Marks et al., 2012; Abranches et al., 2014). For both, specific regulatory properties and their inherent stochastic nature have been suggested to play a role (Martinez Arias and Brickman, 2011). Nevertheless, ERK activity has emerged as a key trigger of *Nanog* heterogeneity, in line with its general role in eliciting exit from pluripotency (Kunath et al., 2007; Schröter et al., 2015). However, little is known about the chromatin-based aspects of *Nanog* heterogeneity. More specifically, it is unknown whether specific chromatin modifications (Jenuwein and Allis, 2001) contribute to stabilisation of the NANOG-negative state, which has been shown to be perpetuated for several cell divisions during self-renewal. Indeed, temporal analysis of NANOG fluctuations across cell generations has shown that the progeny of NANOG-negative cells is enriched in cells that maintain *Nanog* silencing, even though they can revert back and re-express NANOG (Hastreiter et al., 2018). This stability of the NANOG-negative state sets *Nanog* heterogeneity apart from other phenomena of gene expression heterogeneity, generally characterised by fast-switching dynamics resulting from intrinsic and extrinsic noise or encoded in regulatory networks themselves (Huang, 2009; Balazsi et al., 2011).

¹Department of Developmental and Stem Cell Biology, Institut Pasteur, Université Paris Cité, CNRS UMR3738, Epigenomics, Proliferation, and the Identity of Cells Unit, F-75015 Paris, France. ²Department of Genetics and Developmental Biology, Institut Curie, PSL Research University, INSERM, CNRS, 75005 Paris, France. ³Université Paris Cité, CNRS, Institut Jacques Monod, F-75013 Paris, France.

*Author for correspondence (pnavarro@pasteur.fr)

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In this study, we aimed at identifying histone modifications associated with gene repression that would be: (1) differentially enriched at the *Nanog* locus when it is active or inactive; (2) controlled by the signalling pathways associated with heterogeneity; and (3) heritable from mother to daughter cells (Gonzalez et al., 2021). We found histone H3 lysine 9 tri-methylation (H3K9me3; Jambhekar et al., 2019) enriched between the *Nanog* promoter and its -5 kb enhancer (Loh et al., 2006; Levasseur et al., 2008) to fulfil these criteria. We also found that H3K9me3 at NANOG depends on DNA methylation and on the binding of ZFP57, a transcription factor known to nucleate heterochromatin in ESCs (Quenneville et al., 2011; Zuo et al., 2012; Anvar et al., 2016; Riso et al., 2016; Li et al., 2008). Analysis of independent ESC mutants lacking H3K9me3 at *Nanog* revealed its role in promoting NANOG heterogeneity, commitment into differentiation and, most notably, effective differentiation along the primitive endoderm lineage. Moreover, our data also suggest that during differentiation H3K9me3 at *Nanog* becomes independent of ERK activity. Hence, we propose that the timely establishment of ERK-independent H3K9me3 at *Nanog* marks commitment into differentiation and impacts cell-fate acquisition in a lineage-dependent manner.

RESULTS

ERK-dependent, mitotically stable, H3K9me3 at the *Nanog* locus in ESCs

To explore the involvement of chromatin marks potentially distinguishing active and inactive *Nanog* states, we first performed a comparison of ESC populations exhibiting heterogeneity, cultured in the presence of fetal calf serum (FCS) and leukaemia inhibitory factor (LIF) (FCS+LIF) (Fig. 1A) and highly homogeneous populations obtained by double inhibition of ERK and GSK3b (2i+LIF; Fig. 1A). Although several euchromatic marks associated with active transcriptional states were found to be more enriched in 2i+LIF, as expected, we observed a single repressive mark, H3K9me3, to be enriched in FCS+LIF and lost in 2i+LIF (Fig. 1B). H3K9me3 was present at neither the promoter (P in Fig. 1B) nor the enhancer (E in Fig. 1B); rather, we detected H3K9me3 in the intervening region (IR in Fig. 1B). To characterise H3K9me3 further, we performed a high-resolution analysis of the *Nanog* locus (Fig. 1C, top), which confirmed the presence of a robust peak of H3K9me3 between the enhancer and the promoter in the FCS+LIF condition exclusively (Fig. 1C). Analysis of total H3 confirmed the specificity of H3K9me3 enrichment (Fig. 1C), which cannot be

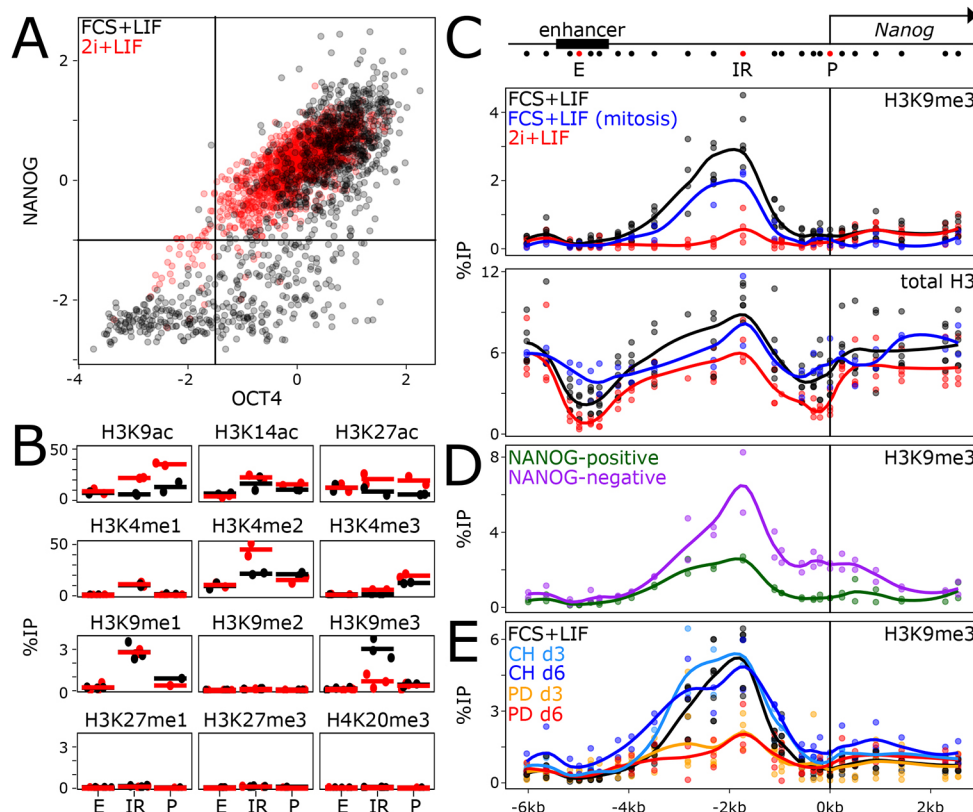


Fig. 1. Mitotically stable, ERK-dependent H3K9me3 at the *Nanog* locus. (A) Quantification of OCT4 (z-score; x-axis) and NANOG (z-score; y-axis) levels as assessed by immunostaining of ESCs cultured in FCS+LIF (black, $n=1125$) and 2i+LIF (red, $n=1445$). The difference in NANOG levels distribution was assessed with Kolmogorov–Smirnov test ($P<2.2e-16$). (B) ChIP-qPCR of several histone modifications, as indicated, at three positions of the *Nanog* locus (the *Nanog* promoter, P; the *Nanog* -5 kb enhancer, E; an intervening region, IR, located 1.7 kb of the *Nanog* transcription start site; see the red dots on the schematic in C) in ESCs cultured in FCS+LIF (black) or in 2i+LIF (red). Each dot represents the percentage of immunoprecipitation (%IP; y-axis) for independent replicates and the bar the corresponding mean. The loss of H3K9me3 observed at IR in 2i+LIF was assessed with unpaired one-tailed Student's t -test ($P=0.006135$). (C–E) Extended profile of H3K9me3 or total H3, as indicated, across the *Nanog* locus (x-axis represents genomic distances in kb with respect to the *Nanog* transcription start site, as schematised above), in ESCs cultured as indicated in each plot. Each dot represents the percentage of immunoprecipitation (%IP; y-axis) for independent replicates and the line the corresponding Loess regression. The differences in H3K9me3 enrichments at the IR region (from -3.5 kb to -0.9 kb) were analysed with unpaired one-tailed Student's t -test combining all individual primers ($P=2.316e-09$ for FCS+LIF versus 2i+LIF; $P=0.001528$ for NANOG negative versus positive; $P=1.367e-06$ for untreated versus PD₃). The increase in H3K9me3 at the promoter region between NANOG-positive versus -negative cells was analysed as for the IR but using primers within the -0.5 kb to 0 kb region ($P=2.447e-07$). d, day.

merely attributed to changes in nucleosome positioning or occupancy. Moreover, we observed good retention of H3K9me3 at *Nanog* in mitotic ESCs (Fig. 1C), in which drastic changes in nucleosomal densities could also be observed at the *Nanog* enhancer (Fig. 1C; Festuccia et al., 2019). Next, we used a previously described *Nanog*-GFP reporter (Chambers et al., 2007) to sort *Nanog*-positive and -negative ESCs by fluorescence-activated cell sorting (FACS). We observed that H3K9me3 was more prominent in *Nanog*-negative cells, with clear spreading towards the promoter (Fig. 1D). H3K9me3 was also found present, albeit at low levels, in *Nanog*-positive cells, obtained either by FACS (Fig. 1D) or by taking advantage of a puromycin selection cassette linked to the *Nanog*-GFP allele (Fig. S1A,B). Finally, we assessed the temporal ERK and GSK3b dependencies of H3K9me3 at *Nanog*. After 3 days of ERK inhibition with PD0325901, which induces high and homogeneous NANOG expression (Fig. S1C,D), H3K9me3 was significantly reduced (PD in Fig. 1E), whereas even after 6 days of GSK3b inhibition with CHIR99021, H3K9me3 levels remained globally unchanged (CH in Fig. 1E). Hence, the repressive H3K9me3 mark exhibits properties that indicate it may play a role in *Nanog* heterogeneity as it is readily dependent on ERK activity, lost in homogeneous NANOG populations, over-enriched in *Nanog*-negative cells, and maintained during mitosis.

H3K9me3 at *Nanog* impacts NANOG heterogeneity

To study the relevance of H3K9me3 at *Nanog*, we used a Crispr/Cas9 approach deleting ~1.8 kb between the enhancer and promoter (Fig. 2A, red box). Two clones ($\Delta K9.1$ and $\Delta K9.2$) were confirmed as homozygous deletions with a complete absence of H3K9me3 (Fig. 2A). *Nanog* mRNA levels were slightly upregulated in $\Delta K9$ cells (Fig. 2B), which presented a clear shift in NANOG expression, leading to a strong reduction of the proportion of cells expressing no or low NANOG (Fig. 2C), as confirmed at the mRNA level by single-molecule fluorescence in situ hybridisation (smFISH) (Fig. S1E,F). Nevertheless, the loss of heterogeneity was not as prominent as that achieved by ERK inhibition (Fig. S1D), indicating that ERK also inhibits *Nanog* transcription by other means. In line with this, *Nanog* expression further increased upon ERK inhibition in $\Delta K9$ clones (Fig. S1C). Moreover, whereas strong ectopic induction of NANOG leads to improved self-renewal (Chambers et al., 2007), the small upregulation of *Nanog* in $\Delta K9$ cells was associated with a marginal increase in self-renewal efficiency, as determined by clonal assays (Fig. 2D). However, $\Delta K9$ cells were more recalcitrant to efficient differentiation upon LIF withdrawal (Fig. 2D). Next, we performed RNA-sequencing (RNA-seq) analysis to compare wild-type and $\Delta K9$ cells (Table S1), which confirmed a small increase in *Nanog* expression [false discovery rate (FDR)<0.05; Fig. S2A]. We identified 235 and 402 genes that were up- or downregulated, respectively, in both $\Delta K9$ clones {FDR<0.05 and $\text{abs}[\log_2\text{fold change (FC)}]>0.3$; Fig. S2B}. Although differentially expressed genes exhibited small fold changes (Fig. S2C), consistent with the small increase in *Nanog* expression observed in $\Delta K9$ cells, they were nonetheless found enriched in the vicinity of NANOG-binding regions, compared with regions bound by OCT4 (POU5F1) and SOX2 but not NANOG (Festuccia et al., 2019; Heurtier et al., 2019; Fig. S2D). We conclude that deletion of the region harbouring H3K9me3 at the *Nanog* locus reduces the capacity of ESCs to explore the NANOG-negative state efficiently, leading to minimal gene expression changes and a measurable resistance to differentiation.

H3K9me3 at *Nanog* controls the timing of commitment into differentiation

We next aimed at characterising the status of H3K9me3 at *Nanog* in non-pluripotent cells. First, we established the H3K9me3 profiles over the *Nanog* locus in several cell types in which *Nanog* has been silenced during development (Fig. 3A). H3K9me3 was systematically found to be enriched between the *Nanog* enhancer and promoter, albeit at different levels. In trophectoderm stem cells (TSCs), the levels found at *Nanog* were lower than in FCS+LIF ESCs, except at the *Nanog* promoter where some spreading was detected. In extra-embryonic endoderm (XEN) cells, the levels of H3K9me3 were higher, exhibiting spreading towards the promoter. Finally, in mouse embryonic fibroblasts (MEFs), the level of H3K9me3 was particularly high, with prominent invasion of the

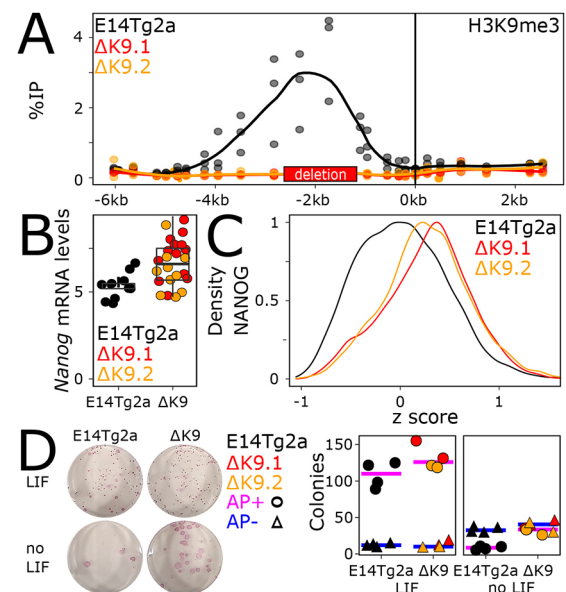


Fig. 2. The *Nanog* region harbouring H3K9me3 enables heterogeneous NANOG expression. (A) H3K9me3 profile across the *Nanog* locus, presented as in Fig. 1C, in wild-type ESCs (E14Tg2a, black) and in two mutant derivatives ($\Delta K9.1$ and $\Delta K9.2$, in red and orange, respectively), carrying a deletion of the region enriched for H3K9me3 (red box on the x-axis). The differences in H3K9me3 enrichments at the non-deleted parts of the IR region were analysed with unpaired one-tailed Student's *t*-test combining all individual primers and the two mutant clones ($P=0.001036$). (B) Expression of *Nanog* mRNA levels (normalised to *Tbp*) assessed by RT-qPCR in wild-type (E14Tg2a, black) and mutant (red and orange) clones. Each dot represents the level of *Nanog* mRNA for independent replicates and the boxplots the corresponding median (bar), 25-75% percentiles (box) and 1.5 \times the inter-quartile range (whiskers). The difference between E14Tg2a and $\Delta K9$ cells was assessed with unpaired one-tailed Student's *t*-test ($P=0.0002491$). (C) Histogram representing the density (y-axis) of NANOG expression levels [x-axis; $\log_2(\text{mean_intensity})$ corrected to the E14Tg2a median of each experiment, $n=2$] in wild-type (E14Tg2a, black; $n=8191$) or mutant ($\Delta K9.1$ and $\Delta K9.2$ in red and orange, respectively; $n=3835$, 3812) cells as assessed by immunostaining. The difference in NANOG distributions was analysed with a Kolmogorov-Smirnov test ($P<2.2e-16$ for each clone versus E14Tg2a). (D) Representative alkaline phosphatase staining of ESC colonies for the indicated cell lines and culture conditions. The plot shows the number of alkaline phosphatase (AP)-positive (circles) and -negative colonies (triangles) in wild-type (E14Tg2a, black) and $\Delta K9$ (red and orange) cells. Each dot represents an independent replicate and the bar the corresponding median for AP-positive (pink) or AP-negative (blue) colonies. The small difference in the number of AP-positive colonies between E14Tg2a and mutant cells cultured in FCS+LIF was not statistically significant (Mann-Whitney test, $P=0.1714$), whereas it was significant in the absence of LIF ($P=0.01429$).

Nanog promoter and gene body. Therefore, we conclude that, although H3K9me3 is found at *Nanog* in the three cell types analysed, its absolute levels and the degree of spreading towards the promoter are variable. This suggests that a certain level of developmental specificity impacts H3K9 methylation at *Nanog*. Moreover, and in contrast to ESCs, inhibition of ERK in MEFs did not abolish H3K9me3 at *Nanog*, which remained robustly enriched (Fig. 3A). This indicates that during differentiation, when *Nanog* silencing becomes irreversible, H3K9me3 at *Nanog* is liberated from its strict dependency on ERK.

We then aimed at assessing the dynamics of H3K9me3 at *Nanog* during ESC differentiation. We used a simple protocol starting from 2i+LIF (absence of H3K9me3) and based on the withdrawal of LIF and ERK/GSK3b inhibitors (Fig. S3A,B). We observed a step-wise increase of H3K9me3 (Fig. 3B): if it remained low during the first 48 h, it suddenly appeared after 3 days and increased at days 4 and 7, when low but clear signs of spreading to the promoter were also observed. In Δ K9 clones, however, H3K9me3 remained absent during differentiation (Fig. S3C). Somehow, unexpectedly, the appearance of H3K9me3 at day 3 did not correlate with a particularly strong reduction of *Nanog* expression (Fig. 3C). In fact, we observed *Nanog* downregulation taking place largely during the first 48 h, in the absence of high levels of H3K9me3. However, although *Nanog* expression continued to decrease during differentiation of wild-type cells, when H3K9me3 further increased and then spread to the *Nanog* promoter (Fig. 3B) Δ K9 cells displayed a stabilisation of low *Nanog* expression after the sharp decrease occurring during the first 2 days (Fig. 3C), despite efficient differentiation (Fig. S3A,B). Immunofluorescence analyses further indicated that the retention of low but measurable NANOG expression affected the vast majority of Δ K9 cells (Fig. S4). This different global behaviour of *Nanog* expression in Δ K9 clones, temporally correlated with the time at which H3K9me3 is first established and subsequently spreads to the promoter in wild-type cells, prompted us to determine whether commitment into differentiation – the moment at which cells cannot easily come back to an undifferentiated state – was altered in Δ K9 cells. For this, we seeded wild-type and Δ K9 cells at clonal density and after 2, 3 or 7 days of differentiation, we replaced the culture medium with 2i+LIF: only cells that have not yet irreversibly lost their capacity to self-renew will survive, proliferate and form undifferentiated, alkaline phosphatase-positive colonies (Kalkan et al., 2017; Fig. 3D,E). In wild-type cells, we observed a striking coincidence of the time of commitment, taking place between days 2 and 3, with the appearance of H3K9me3 at *Nanog*. In Δ K9 clones, however, we observed a significant number of alkaline phosphatase-positive colonies after 3 and even 7 days of differentiation, indicating a delay in commitment. Altogether, these analyses suggest that in ESCs cultured in 2i+LIF, H3K9me3 at *Nanog* is established during differentiation, when it marks the irreversible commitment into effective differentiation. However, it is not strictly required for differentiation per se and rather enables the appropriate timing of commitment.

Δ K9 cells exhibit delayed differentiation

Given the delay in differentiation commitment observed in Δ K9 clones, we monitored the expression of several genes reflecting the loss of pluripotency and the transition to a differentiated state (Kalkan et al., 2017; Fig. S3B). Whereas naïve pluripotency genes [*Esrrb*, *Klf4*, *Prdm14*, *Rex1* (*Zfp42*)] showed a less drastic downregulation, mimicking *Nanog* expression, differentiation markers (*Fgf5*, *Dnmt3b*, *Otx2*, *Wnt3*) showed slightly delayed

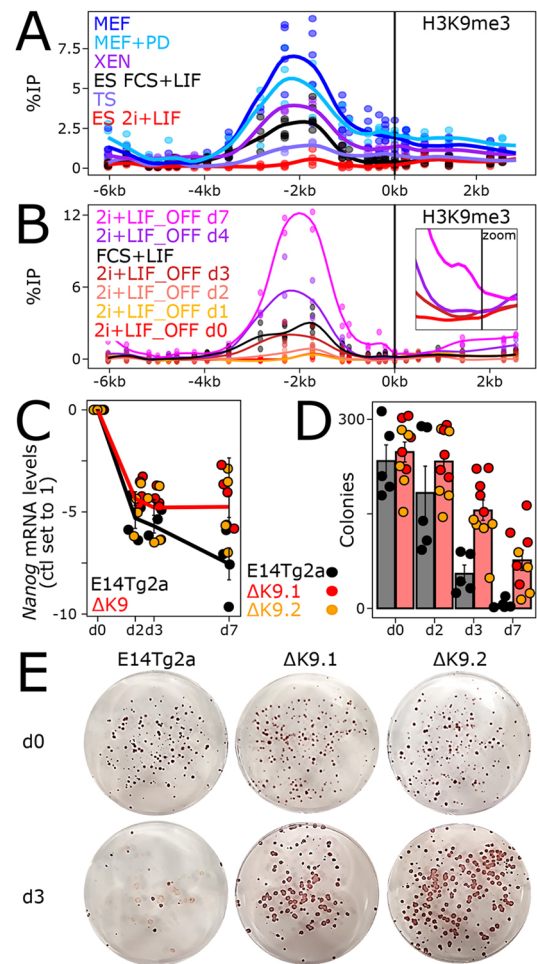


Fig. 3. H3K9me3 at *Nanog* times commitment into differentiation.

(A) H3K9me3 profile across the *Nanog* locus, presented as in Fig. 1C, in the indicated cell lines and conditions. The enrichment of H3K9me3 at either the IR (from -3.5 kb to -0.9 kb) or the promoter region (from -0.5 kb to 0 kb) in non-pluripotent cells was compared with 2i+LIF ESCs with unpaired one-tailed Student's *t*-test combining all individual profiles falling in each subregion ($P=0.002566$ and 2.307×10^{-6} for IR and promoter, respectively, for TSCs; $P=0.001053$ and 4.716×10^{-6} for XEN cells; $P=4.878 \times 10^{-8}$ and 3.79×10^{-9} for MEFs). (B) Identical profiles for ESCs undergoing differentiation (labelled 2i+LIF_OFF) for the indicated number of days. The same statistical analyses as in A showed that the enrichment at the IR and promoter region was statistically significant from day 3 onwards ($P=0.00226$, 0.0001353 , 0.0004633 for the IR, and $P=0.04536$, 0.0004678 , 0.002106 for the promoter, at days 3, 4 and 7, respectively). The inset focuses on the promoter region (-1 kb to 0.5 kb) to better appreciate H3K9me3 spreading from day 0 to day 3, 4 and 7. (C) *Nanog* log2 relative mRNA levels (day 0 set to 1) measured by RT-qPCR and normalised to *Tbp*, during ESC differentiation in the indicated cell lines. Each dot represents an independent replicate and the line the corresponding mean with s.e.m. The comparative analysis of the temporal reduction of *Nanog* between E14Tg2a and Δ K9 clones (unpaired one-tailed Student's *t*-test) showed that it was statistically significant only at day 7 ($P=0.08368$, 0.05471 and 0.01324 at days 2, 3 and 7, respectively). Moreover, the reduction observed for E14Tg2a between days 3 and 7 was also found to be significant ($P=0.03026$). (D) Number of alkaline phosphatase-positive colonies obtained after switching to 2i+LIF for wild-type (E14Tg2a, black) and Δ K9 (red and orange) cells seeded clonally and differentiated for the number of days indicated on the x-axis. Each dot represents an independent replicate and the histogram the corresponding mean and s.e.m. Differences in clonogenicity were assessed with a Mann-Whitney test ($P=0.2198$, 0.001332 , 0.002531 for days 2, 3 and 7, respectively). (E) Representative alkaline phosphatase staining of ESC colonies cultured in 2i+LIF after 0 (top) or 3 (bottom) days of differentiation for the indicated cell lines. d, day.

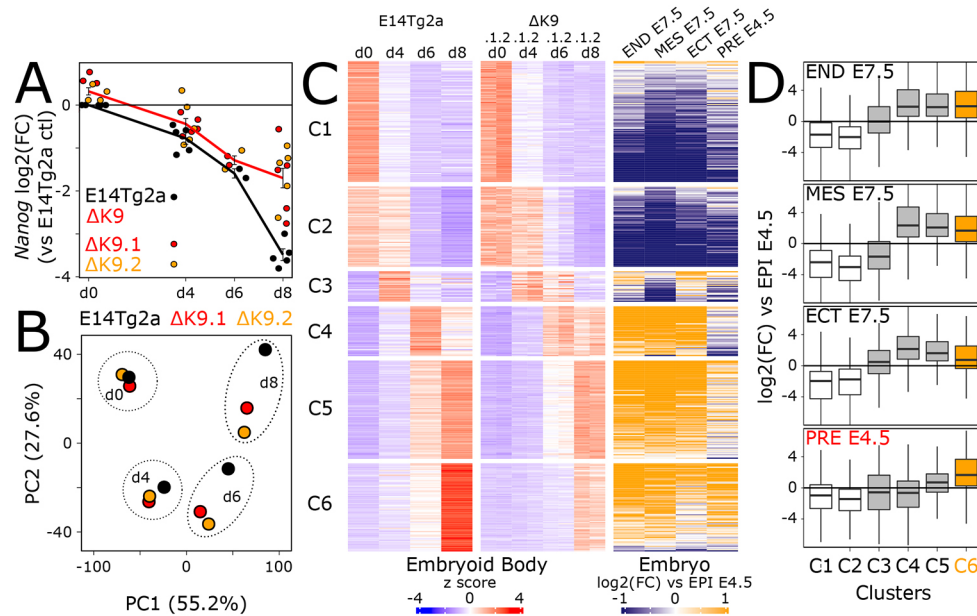


Fig. 4. The lack of H3K9me3 at *Nanog* leads to delayed differentiation. (A) Log2 *Nanog* mRNA levels measured by RT-qPCR and normalised to *Tbp* during EB differentiation in the indicated cell lines. Each dot represents an independent replicate and the line the corresponding mean with s.e.m. Expression differences between E14Tg2a and Δ K9 clones were evaluated at each day of differentiation with unpaired one-tailed Student's *t*-tests ($P=0.002143$, 0.0485 , 0.06709 , $7.367e-06$ for days 0, 4, 6 and 8, respectively). (B) PCA of 16,336 transcripts quantified by RNA-seq (TPM >1 in at least one sample) in wild-type (E14Tg2a, black; $n=3$) and Δ K9 (red and orange; $n=3$ for each) cells undergoing EB differentiation for the indicated days. (C) Heatmap of 4100 transcripts displaying gene expression changes during EB differentiation [FDR<0.05 and $\text{abs}(\log_2\text{FC})>1$ in at least one comparison to undifferentiated cells; $n=3$ for each cell line], clustered by k-means after z-score normalisation. Left: Relative gene expression (z-score) in wild-type (E14Tg2a) and Δ K9 cells during EB differentiation. Right: Corresponding $\log_2\text{FC}$ of each developmental stage indicated at the top versus E4.5 epiblast, as previously reported (Argelaguet et al., 2019). (D) Boxplot (median; 25–75% percentiles; $1.5\times$ the inter-quartile range) of the $\log_2\text{FC}$ shown in C for each cluster. Cluster C6 is highlighted in orange. The highest distribution of $\log_2\text{FC}$ for PRE E4.5 for C6 with respect to the other clusters was assessed with a Kolmogorov–Smirnov test ($P<1e-07$ for all comparisons). d, days; ECT, ectoderm; END, endoderm; EPI: epiblast; MES, mesoderm; PRE, primitive endoderm.

dynamics. Next, we differentiated wild-type and Δ K9 cells into embryoid bodies (EBs), a paradigm that recapitulates the establishment of multiple lineages. At the morphological level, we observed Δ K9 EBs to be often characterised by defective sealing at their periphery (Fig. S5A, top). Moreover, cellular outgrowths derived from Δ K9 EBs also exhibited less morphological typologies compared with those derived from wild-type EBs, suggesting altered multi-lineage differentiation (Fig. S5A, bottom). Gene expression analyses of EBs after days 4, 6 and 8 of differentiation confirmed the attenuated downregulation of *Nanog* expression (Fig. 4A). Moreover, principal component analysis (PCA) analysis of RNA-seq profiling (Table S1) highlighted a transcriptome-wide delay of both Δ K9 clones, starting at day 4 and progressively increasing through time (Fig. 4B). Gene ontology analysis of the top 1000 loadings of the PCA identified focal adhesion genes among the most enriched cellular components (GO:0005925; $P<10^{-5}$), in line with our morphological observations (Fig. S5A). Therefore, the lack of H3K9me3 at *Nanog* is strongly associated with delayed differentiation, as evaluated with three distinct differentiation paradigms (Fig. 2D, Fig. 4B, Fig. S3B).

The absence of H3K9me3 at *Nanog* leads to major defects in primitive endoderm differentiation

To explore further the molecular nature of the defects observed during EB differentiation, we first identified around 4000 genes displaying temporal gene expression changes in either differentiating wild-type or mutant cells compared with their undifferentiated controls [FDR<0.05, $\text{abs}(\log_2\text{FC})>1$; Table S1]. These genes were then clustered using k-means; this allowed us

to identify six groups of genes displaying different expression dynamics during EB differentiation (Fig. 4C, left; Table S1). Although all identified clusters underscored the delay into differentiation, one in particular, cluster C6, showed a blatant attenuation of gene upregulation at day 8. To characterise each cluster relative to known developmental trajectories, we plotted the fold change reported in a previous study (Argelaguet et al., 2019), in which embryonic endoderm, mesoderm and ectoderm cells of embryonic day (E) 7.5 embryos, as well as primitive endoderm cells of E4.5 embryos, were directly compared with the E4.5 epiblast (Fig. 4C, right). Whereas clusters C1 and C2 showed a marked downregulation upon epiblast differentiation into all lineages, clusters C3 to C6 displayed a clear upregulation in at least one E7.5 lineage. Notably, cluster C6, which shows the strongest Δ K9 versus wild-type differences, was the only one enriched for genes displaying a prominent upregulation in the primitive endoderm with respect to the epiblast (Fig. 4D). Analysis of individual primitive endoderm markers during EB differentiation confirmed the altered induction of these genes (Fig. S5B,C).

Finally, in light of these results, we wanted to ascertain whether the defective primitive endoderm signature identified in EBs implies a deficiency in the capacity of Δ K9 clones to engage in primitive endoderm differentiation. Thus, we challenged wild-type and Δ K9 ESCs with a primitive endoderm differentiation protocol (Anderson et al., 2017; Fig. S6). In wild-type cells, but not in Δ K9 clones, we observed the appearance of endoderm-like cell clusters from day 4 onwards. Moreover, Δ K9 clones showed increased cell death. Immunofluorescence of NANOG, GATA6, GATA4 and PDGFR α confirmed NANOG silencing in cells expressing

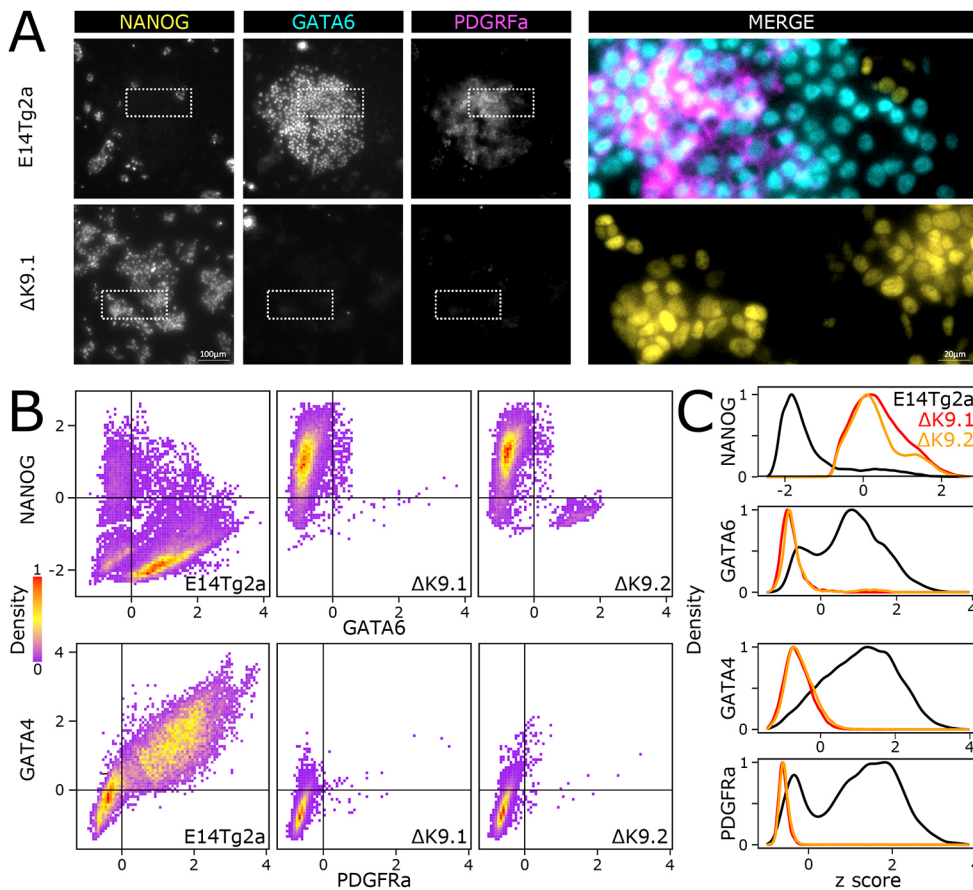


Fig. 5. Lack of H3K9me3 at *Nanog* abolishes primitive endoderm differentiation. (A) Representative immunostaining of ESCs (E14Tg2a, top; Δ K9.1, bottom) subject to a directed differentiation protocol into primitive endoderm. Boxed areas are shown at higher magnification, with signals merged, on the right. (B,C) Quantification of dual immunostaining for NANOG and GATA6 or GATA4 and PDGFR α in E14Tg2a ($n=19,507$, 20,641), Δ K9.1 ($n=11,348$, 19,476) and Δ K9.2 ($n=9713$, 19,742) cells differentiated as in A. $P<2.2e-16$ (Kolmogorov–Smirnov tests) for all E14Tg2a versus Δ K9 comparisons.

primitive endoderm markers, as expected (Fig. 5). In Δ K9 clones, however, NANOG expression was prominent with only the occasional appearance of cells expressing primitive endoderm markers. Therefore, Δ K9 cells are not able to efficiently undergo directed differentiation into primitive endoderm. We conclude that H3K9me3 is required to stably silence *Nanog* during differentiation and that failing to do so has different consequences depending on the differentiation lineage, with primitive endoderm being particularly sensitive.

DNA methylation and ZFP57 binding trigger H3K9me3 at *Nanog*

To investigate the mechanisms by which H3K9me3 is established at the *Nanog* locus, we used the Cistrome database (Zheng et al., 2019) to identify candidate factors binding the region displaying maximal levels of H3K9me3 at *Nanog*. Among the identified factors (Fig. S7A), ZFP57, KAP1 (TRIM28) and DNMT3a appeared particularly relevant. Indeed, ZFP57 has been shown to bind its cognate DNA motif (TGCCGC) only when the CpG is methylated (underlined) and then recruit KAP1, DNA methylases and H3K9 methylases to nucleate heterochromatin formation (Quenneville et al., 2011; Zuo et al., 2012; Anvar et al., 2016; Riso et al., 2016; Li et al., 2008). Two perfect ZFP57 motifs were identified at the exact region showing H3K9me3 in ESCs (Fig. 6A). Moreover, analysis of DNA methylation datasets (Domcke et al., 2015) showed that the two CpGs required for ZFP57 binding are methylated in FCS+LIF but not in 2i+LIF (Fig. 6A,B), as expected given the reduced expression of DNMTs upon ERK inhibition (Li et al., 2016; Spindel et al., 2021 preprint) and the global loss of DNA methylation in 2i+LIF-cultured ESCs (Leitch et al., 2013; Grabole et al., 2013).

Hence, we aimed at profiling ZFP57 binding by chromatin immunoprecipitation-quantitative PCR (ChIP-qPCR). We observed robust recruitment of ZFP57 at the H3K9me3-enriched region in ESCs cultured in FCS+LIF but not in 2i+LIF (Fig. 6C,D). Using previously described *Zfp57* knockout ESCs (ZKO; Riso et al., 2016) we were able to confirm the specificity of our assay (Fig. 6C,D). Moreover, upon knockout of all three DNMT genes [*Dnmt1/3a/3b*; triple knockout (TKO)] and the ensuing loss of DNA methylation (Fig. S7B,C), we also observed a complete loss of ZFP57 binding (Fig. 6C). As expected, in both TKO and ZKO cells H3K9me3 at *Nanog* was completely abrogated (Fig. 6D), establishing that it is triggered by a canonical mechanism dependent on DNA methylation and ZFP57 recruitment. Finally, we aimed to address whether the loss of H3K9me3 at *Nanog* triggered by either DNMT or *Zfp57* knockouts is accompanied by changes in NANOG heterogeneity and in the capacity of ESCs to differentiate into primitive endoderm derivatives, as shown in our Δ K9 mutant cells. In both knockouts, we observed a clear increase of NANOG expression before differentiation, depleting the NANOG-low compartment (Fig. 6E), and a strong attenuation of both NANOG downregulation and induction of GATA6 upon directed differentiation into primitive endoderm (Fig. 6F,G), reproducing the effects observed in Δ K9 ESCs. This reinforces the idea that H3K9me3 at *Nanog* is required to lock the *Nanog* silent state during differentiation, an event that is particularly important for proper primitive endoderm differentiation.

DISCUSSION

Gene expression heterogeneity has emerged as a main motor of lineage diversification during development, particularly in stem cell

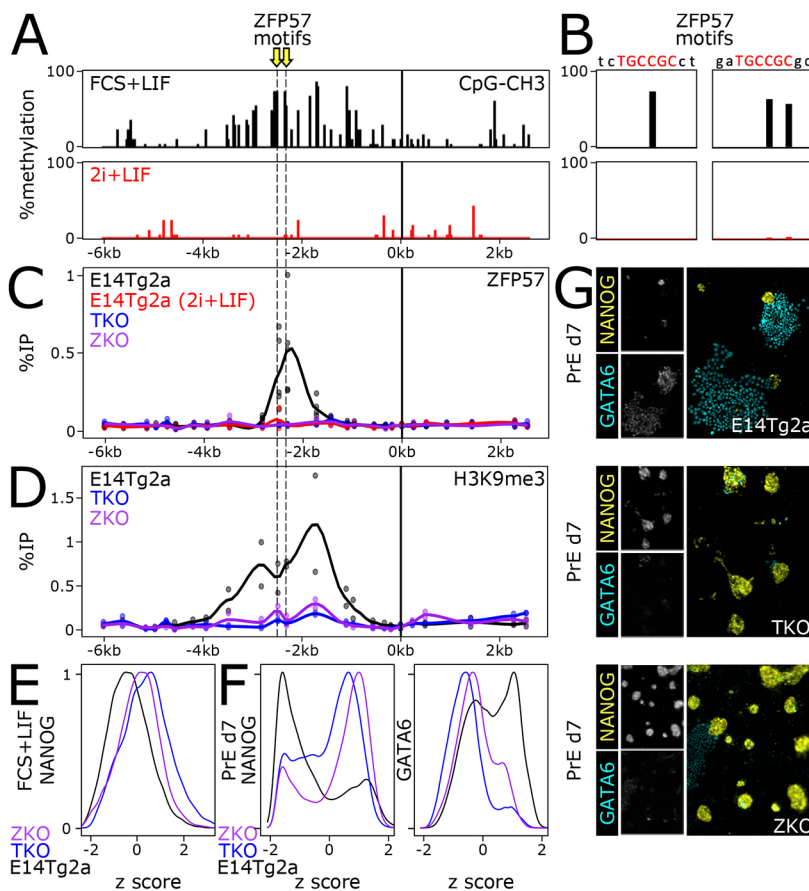


Fig. 6. DNA methylation and ZFP57 binding drive H3K9me3 enrichment at *Nanog*. (A) Analysis of DNA methylation at the *Nanog* locus, using available datasets in ESCs cultured in FCS+LIF or in 2i+LIF. The position of two ZFP57 motifs is indicated by yellow arrows. (B) Identical analysis but focusing at the two ZFP57 motifs. (C) Analysis of ZFP57 binding by ChIP-qPCR, presented as in Fig. 1C, in the indicated cell lines: E14Tg2a cultured in FCS+LIF (black) or in 2i+LIF (red), triple-negative DNMT knockout ESCs (TKO, blue) and *Zfp57* knockout ESCs (ZKO, purple). (D) Analysis of H3K9me3 at *Nanog* in E14Tg2a, TKO and ZKO ESCs cultured in FCS+LIF, presented as in C. (E) Analysis of NANOG expression in E14Tg2a, TKO and ZKO ESCs. The differential distribution in TKO and ZKO versus E14Tg2a was assessed with Kolmogorov–Smirnov tests ($P < 2.2 \times 10^{-16}$). (F) Analysis of NANOG and GATA6 expression after 7 days of directed differentiation of E14Tg2a, TKO and ZKO into primitive endoderm. Differences in distribution levels were assessed with Kolmogorov–Smirnov tests ($P < 2.2 \times 10^{-16}$ for both TKO and ZKO versus E14Tg2a). (G) Representative immunostaining of E14Tg2a, TKO and ZKO subjected to a directed differentiation protocol into primitive endoderm. Insets show individual signals, and main panels merged signals.

and progenitor populations: upon appropriate stimuli, these expression differences are translated into an effective decision-making process that culminates in commitment (Huang, 2009; Martinez Arias and Brickman, 2011; Balazsi et al., 2011). When these heterogeneities are dynamic, such as in the case of *Nanog* and ESCs, the fluctuating expression is eventually fixed. A likely model accounting for the transition from reversible to irreversible silencing involves epigenetic repression, in the sense of heritable chromatin states incompatible with transcription that do not depend on the original triggers (Berger et al., 2009). The acquisition of epigenetic repression, including H3K9me3 (Nicetto and Zaret, 2019), is indeed perceived as a general means to restrict developmental fate choices as cells differentiate (Yadav et al., 2018). Accordingly, we show that *Nanog* silencing during differentiation is accompanied by H3K9me3, which irreversibly locks its expression and thus contributes to commitment into differentiation. Two observations, however, indicate that additional complexity may characterise both *Nanog* silencing and its developmental implications.

Although epigenetic repression is often instated *de novo* during differentiation, for example at *Oct4* (Feldman et al., 2006), the regulation of *Nanog* appears to involve an intermediary state in which H3K9me3 is already established but not yet fixed. In undifferentiated cells, H3K9me3 is readily detected at *Nanog*, particularly in *Nanog*-negative cells. Moreover, it is transferred from mother to daughter cells during mitosis. However, it is strictly dependent upon ERK signalling, the main driver of NANOG heterogeneity. This dependency is likely mediated by the impact of ERK on DNA methylation (Leitch et al., 2013; Grabole et al., 2013; Li et al., 2016; Spindel et al., 2021 preprint), in this case of two

motifs of the ZFP57 transcription factor, which recruits proteins involved in H3K9 methylation (Li et al., 2008; Quenneville et al., 2011; Zuo et al., 2012; Anvar et al., 2016; Riso et al., 2016). By integrating ERK activity with mitotically stable H3K9 methylation, ZFP57 may confer to *Nanog* silencing the required stability to be inherited and, at the same time, sufficient flexibility to revert back to transcriptional activity. During differentiation (at least as judged by the analysis of embryonic fibroblasts), H3K9me3 becomes independent of ERK and, with respect to ERK, irreversible. Therefore, even if subjected to variations in ERK stimuli, *Nanog* will remain silent. Despite these considerations, we suggest that H3K9me3 at *Nanog* becomes epigenetic exclusively during differentiation, when it is no longer dependent on its original trigger. Hence, our data point to an integrated network of ERK signalling, DNA methylation and ZFP57 binding as the mechanism responsible for H3K9me3 enrichment at *Nanog*. This scenario, whereby DNA methylation plays a central role, is compatible with the kinetics of H3K9me3 loss upon ERK inhibition, which are suggestive of passive erasure of CpG methylation. It is also compatible with the incapacity of ESCs to efficiently maintain DNA methylation and H3K9me3 when transiently induced using epigenomic editing tools (Carlini et al., 2022). Whether the transition from reversible to irreversible H3K9me3 is mediated by direct mechanisms operating at the locus or on the H3K9me3-associated machinery, or linked to the general lack of strong epigenetic repression in ESCs (Carlini et al., 2022; Festuccia et al., 2017), remains unknown. Also, it remains possible that the global increase of H3K9me3 taking place during differentiation, which directly depends on the loss of activity of OCT4 and to a lesser extent of NANOG (Bernard et al., 2022), leads to an improvement

in the establishment of robust epigenetic repression. Nevertheless, the dependency of H3K9me3 on ERK shown here and previously on OCT4 (Bernard et al., 2022 preprint) is reminiscent of the dependency of other repressive marks, such as H3K27me3, on LIF signalling and NANOG activity (Heurtier et al., 2019), suggesting a general dependence of repressive chromatin marks on more dynamic regulators in undifferentiated ESCs. By displaying regulated dependencies towards signalling and/or transcription factor activity, repressive chromatin modifications may facilitate conditional heritability and excitability (before differentiation commitment) or, by contrast, fixed gene expression states to enable commitment (Festuccia et al., 2017).

Nanog is known to counteract differentiation when ectopically expressed at high levels (Chambers et al., 2007). Because the deletion of the region harbouring H3K9me3 leads to a minor increase of NANOG expression, it was not expected to block differentiation. After all, upon the collapse of the pluripotency network triggered by differentiation signals, *Nanog* would lose most of its activators and be downregulated, as we observed. However, the lack of H3K9me3 triggered by the deletion leads to a lack of complete *Nanog* silencing, providing an opportunity to evaluate the importance of fully repressing *Nanog* during differentiation. Similarly, as cells lacking the H3K9me3-enriched region at *Nanog* reduce their heterogeneity in a context in which *Nanog* can nevertheless be downregulated (in contrast to ectopic expression systems), the importance of NANOG heterogeneity in lineage priming can also be inferred from our experimental setup. In this regard, our observation that cells lacking the H3K9me3-enriched region can differentiate rules out a deterministic role for NANOG heterogeneity in the capacity to exit the undifferentiated state. Nevertheless, using multilineage protocols we observed delayed commitment and altered differentiation into all germ layers of cells lacking the H3K9me3-enriched region. Yet the highest consequences affect genes normally upregulated in the primitive endoderm, an observation that was fully confirmed by their incapacity to differentiate efficiently into primitive endoderm using a directed differentiation protocol. Although the existence of other forms of regulation mediated by the deleted region cannot be formally excluded, the fact that DNMT and *Zfp57* knockouts (which have a wild-type *Nanog* locus) phenocopy the loss of NANOG heterogeneity and the alteration of primitive endoderm differentiation suggests that H3K9me3 plays a major role. The differential impact observed for somatic versus primitive endoderm lineages in ESCs further underscores the relative and lineage-specific importance of *Nanog* in repressing differentiation. Furthermore, *Nanog* heterogeneity in ESCs has been proposed to either reflect the heterogeneity observed in early blastocysts, whereby cells of the inner mass can either express NANOG or GATA6, or the early downregulation of *Nanog* taking place around implantation to elicit somatic differentiation events of the epiblast (Chambers et al., 2007; Singh et al., 2007; Kalmar et al., 2009; Canham et al., 2010; Abranches et al., 2014). Indirectly, thus, our results could be interpreted as *Nanog* heterogeneity and its subsequent full silencing being functionally associated with epiblast versus primitive endoderm specification. In this regard, two observations are noteworthy. First, the repressive H3K27me3 mark has been shown to play a preponderant role in downregulating genes that prime ESCs for primitive endoderm differentiation (Illingworth et al., 2016). Second, *Nanog* sustains H3K27me3 in undifferentiated and early differentiating ESCs (Heurtier et al., 2019). In light of these findings and of our observation that the loss of H3K9me3 also alters the differentiation balance between somatic and primitive endoderm lineages, we suggest that a signalling and transcription factor

dialogue established through repressive histone methylation contributes to epiblast versus primitive endoderm specification and differentiation. In this model, ERK dynamically controls H3K9me3 at *Nanog*, which sustains H3K27me3 levels and keeps primitive endoderm genes in check, generating reversible and mosaic expression patterns associated with either epiblast or primitive endoderm fates.

Overall, our observations add to the notion that heterochromatin contributes to cell-fate restriction during differentiation processes. They also suggest that these events are more nuanced in their action than anticipated, given the regulation of its dependency on signalling cues in respect to its epigenetic potential and the differential impact that the ensuing stability of the repression may have in different lineages. Whether our findings can be extrapolated to other master regulators of pluripotency or to other developmental transitions represent important new avenues for future research.

MATERIALS AND METHODS

Cell culture

ESCs were cultured on gelatine in either FCS+LIF or 2i+LIF and passaged every 3–4 days. All 2i+LIF analyses were performed after at least three passages in 2i+LIF. ESCs were karyotyped and regularly tested *Mycoplasma*-free. MEFs were derived from F1 129sv/129sv E13.5 male embryos and cultured for no more than four passages. XEN cell lines (Artus et al., 2010) were routinely passaged every 4 days. TSC lines (Kunath et al., 2005) were cultured on mitomycin MEFs and passaged every 2–3 days. For N2B27 differentiation (2i_OFF), ESCs were seeded on poly-L-ornithine/laminin-coated, cell-culture-treated surfaces in 2i+LIF medium but omitting LIF and ERK/GSK3b inhibitors. The medium was changed daily. EBs were obtained by seeding cell aggregates onto non-cell-culture-treated dishes. Primitive endoderm differentiation was performed as previously described (Anderson et al., 2017) using activin A, CHIR99021 and LIF. The medium was changed daily. Commitment assays were performed with cells cultured in 2i+LIF and subject to 2i_OFF differentiation; at each differentiation time point, 2i+LIF was added back for seven additional days, after which cells were fixed and stained for alkaline phosphatase activity. Full details are available in supplementary Materials and Methods.

Generation of ESC knockout lines

To generate $\Delta K9$ clones, we used gRNAs (5'-CAGAGGAGGGCTTAA-GAGAT and 5'-CACTCTAACCAGCTTAAGT) cloned under the control of a U6 promoter in a vector conferring puromycin resistance (Heurtier et al., 2019). These vectors were co-transfected with a Cas9/mCherry expression vector (Addgene plasmid #64324) in E14Tg2a cells, selected with puromycin and FACS-sorted for mCherry fluorescence. Puromycin-resistant and mCherry-positive cells were seeded at clonal density and ~100 clones were picked 10 days later. Clones were screened by PCR with primers spanning the deletion (Table S2) by real-time qPCR using primers along the *Nanog* locus (Table S2), and by cloning and sequencing of PCR products (Table S2). Two karyotypically normal independent clones, $\Delta K9.1$ and $\Delta K9.2$, were selected for this study. *Dnmt1/3a/3b* TKO cells were generated in E14Tg2a using gRNAs that were previously described (Domcke et al., 2015). The three gRNAs targeting each DNMT gene were cloned in pX459 (Addgene plasmid #48139) and transfected into E14Tg2a cells. After puromycin selection, ~96 colonies were picked and screened by DNA digestion with the DNA methylation-sensitive restriction enzyme HpaII. DNA methylation mutant clones were further confirmed by Sanger sequencing (Table S2). For the TKO clone used in this study, complete loss of DNA methylation was confirmed by the luminometric methylation assay (LUMA; Karimi et al., 2006), which was performed exactly as described (Walter et al., 2016).

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Nanog-GFP cells (TNG; Chambers et al., 2007) were sorted using a MoFlo Astrios with the 'highest % of purity' parameter selected. After sorting, GFP-negative and GFP-positive sorted populations were re-processed, with

the same parameters used for sorting, to check the purity of each fraction (>95%).

Imaging analyses

To enable direct comparison of undifferentiated E14Tg2a with $\Delta K9$, TKO and ZKO cells, cells were individually incubated either with Rhodamine Red (E14Tg2a) or Deep Red ($\Delta K9$ or TKO or ZKO) dyes, collected, mixed at a 1:1 ratio, seeded and cultured for ~6 h. Cells were then fixed in 4% formaldehyde for 10 min and used for immunostaining. Differentiating cells were processed separately. Antibodies used for all immunostaining experiments are listed in Table S2. Imaging was performed with an inverted Nikon Eclipse X microscope equipped with $\times 20/0.45$ (WD 8.2–6.9) objective, LUMENCOR excitation diodes, Hamamatsu ORCA-Flash 4.0LT camera and NIS Elements 4.3 software. Quantifications were performed using CellProfiler (Carpenter et al., 2006). For smFISH analyses, cells were grown at low density in medium without phenol-Red, collected by trypsinisation, fixed, cytospun and stored in 70% ethanol at 4°C until use. The slides were dehydrated and hybridised for 24 h at 37°C with the *Nanog* mRNA probe (Stellaris Probe Designer version 4.2 on Biosearch Technologies). Image stacks (0.5 μ m gap) were acquired using a Nikon Eclipse X microscope equipped with $\times 63$ oil immersion objective (N.A1.4), LUMENCOR excitation diodes, Hamamatsu ORCA-Flash 4.0LT camera and NIS Elements 4.3 software. The analysis was performed using ImageJ and CellProfiler. Additional details are available in supplementary Materials and Methods.

Gene expression analyses

RNA extraction and DNase treatment were performed with the NucleoSpin RNA Mini kit (MACHEREY-NAGEL) according to the manufacturer's protocol. Reverse transcription was performed with 1 μ g of total RNAs with random hexamers following manufacturer's protocol (Roche). Real-time quantitative PCR was performed in a LightCycler 480 (Roche) using LightCycler 480 SYBR Green I Master (Roche) and normalised to *Tbp* (see Table S2 for primer sequences). Stranded, poly-A-selected RNA-seq libraries were prepared and sequenced (paired-end 150 bp reads; around 50 million each) by Novogene UK. Reads were aligned to the mm10 genome and transcripts per million (TPM) computed. All differential expression tests were run with DESeq2 (Love et al., 2004). For EB differentiation, we considered genes with absolute $\log_2FC > 1$ and $FDR < 0.05$ at any day of the differentiation versus undifferentiated cells for either E14Tg2a or $\Delta K9$ cells. k-means clustering was computed with R using the function k-means with options $k=6$, $nstart=50$, $iter.max=50$. Only differentially expressed genes as identified during EB differentiation were used (z-scored mean TPM). The number of clusters was chosen as the minimal value identifying at least one cluster with maximal expression at each day of differentiation, including day 0. Correlations with developmental gene expression were made by directly plotting the \log_2FC reported in a previous study using scNMT-seq (single-cell nucleosome, methylation and transcription sequencing) around gastrulation of mouse embryos (Argelaguet et al., 2019). Additional details are available in supplementary Materials and Methods.

Chromatin analyses

After trypsinisation, ESCs were cross-linked and nuclei isolated and sonicated using a Covaris M220. Fragmented chromatin (typically 200–600 bp) was used for ChIP using the antibodies listed in Table S2. ChIP and the corresponding Input samples were analysed by qPCR using the primers listed in Table S2. Additional information is available in supplementary Materials and Methods.

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Competing interests

The authors declare no competing or financial interests.

Author contributions

Conceptualization: A.D., P.N.; Methodology: A.D., M.C.-T.; Formal analysis: A.D., A.C., P.N.; Investigation: A.D., L.V., S.V.-P.; Resources: M.V.C.G., D.B.; Data curation: A.D., A.C., P.N.; Writing - original draft: P.N.; Writing - review & editing: A.D., M.C.-T., P.N.; Supervision: P.N.; Funding acquisition: P.N.

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Data availability

All RNA-seq datasets generated for this study have been deposited in Gene Expression Omnibus under accession number GSE178835.

Peer review history

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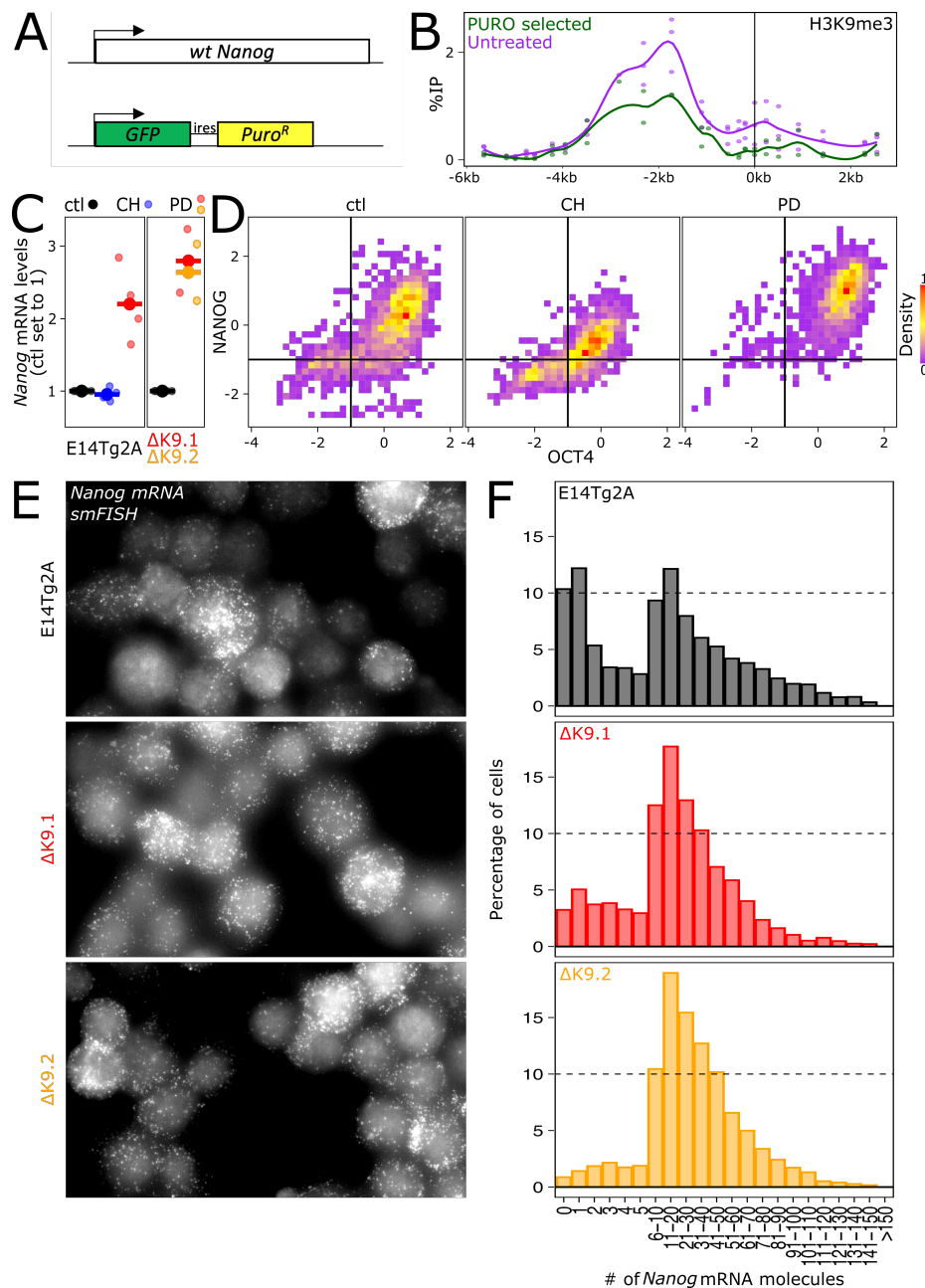


Fig. S1. Additional information on *Nanog* heterogeneity.

(A) Schematic representation of the two *Nanog* alleles in *Nanog*-GFP reporter cells (TNG; Chambers et al. 2007). Selection with Puromycin leads to a fully NANOG-GFP+ve homogeneous population (Chambers et al. 2007, Navarro et al. 2012). **(B)** H3K9me3 ChIP-qPCR analysed and presented as in Fig. 1C, D ($p = 0.02373$ and 8.404×10^{-6} for the IR and promoter region, respectively). **(C)** Relative mRNA levels (ctl set to 1) of *Nanog*, measured by RT-qPCR and normalised to *Tbp*, upon 3d of PD or CH treatment of FCS+LIF cultured cells. The increase observed upon PD treatment was tested with Student's t test ($p = 0.009019$ for E14Tg2a and 0.002947 for the two $\Delta K9$ clones combined). **(D)** Quantification (z score) of NANOG and OCT4 immuno-staining in untreated E14Tg2a cells (ctl; $n = 2053$) and after 3d of CH ($n = 1554$) or PD ($n = 1857$) treatment, as indicated. The changes in NANOG distribution in PD treated cells were assessed with a KS test ($p < 2.2 \times 10^{-16}$). **(E)** Representative image of smFISH assays using a *Nanog* mRNA probe in the indicated cell lines. **(F)** Quantification of the number of *Nanog* mRNA molecules per cell for each indicated cell line. The distributions were compared with a Chi-squared test ($p < 2.2 \times 10^{-16}$).

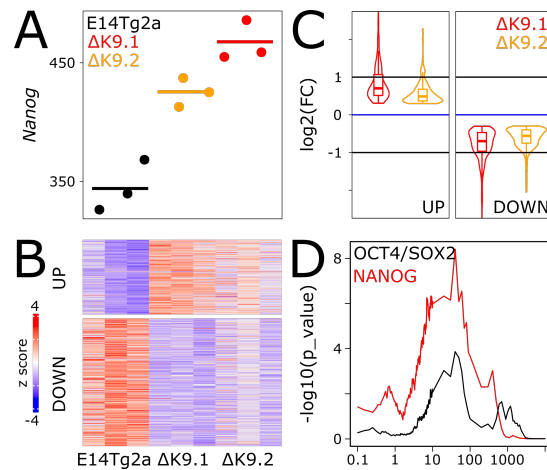


Fig. S2. Gene expression consequences in $\Delta K9$ cells. (A) Confirmation of *Nanog* upregulation in $\Delta K9$ cells by RNA-seq (Transcripts Per Million, TPM). (B) Z scored heatmap of genes identified as differentially expressed in $\Delta K9$ cells. (C) Violin-Boxplot of gene expression fold-changes measured in $\Delta K9.1$ and $\Delta K9.2$ clones compared to wild-type E14Tg2a cells. (D) Statistical association (Y-axis, $-\log_{10}(\text{Fisher exact test p-value})$) between the differentially expressed genes shown in (B) with the presence of NANOG (red) or OCT4/SOX2 (black) binding sites as a function of the distance (X-axis, kb).

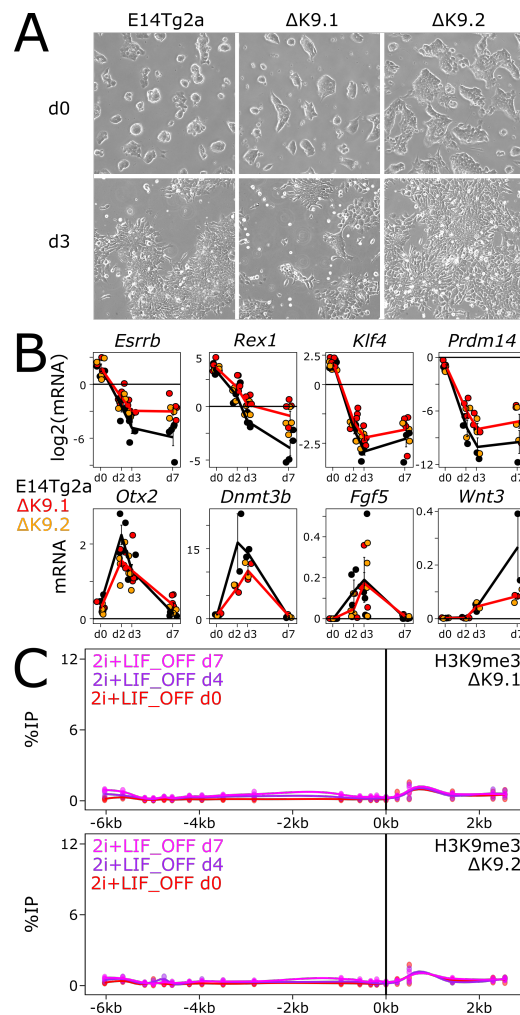


Fig. S3. Differentiation of wild-type and $\Delta K9$ cells upon 2i+LIF withdrawal.

(A) Representative bright-field photomicrographs of wild-type (E14Tg2a) and $\Delta K9$ cells cultured in 2i+LIF (top) and after 3 days of withdrawal (bottom). **(B)** RT-qPCR analysis of markers of naïve pluripotency (top) and differentiation (bottom) after 0, 2, 3 and 7 days of 2i+LIF withdrawal. *Tbp* was used for normalisation. Differences in naïve marker downregulation were assessed with a Student's t test at the day of strongest reduction in wild-types (d7 for *Esrrb* and *Rex1*; d3 for *Klf4* and *Prdm14*; $p = 0.02361, 0.007429, 0.01211, 0.08776$, respectively). For differentiation markers, d2 was used ($p = 0.0214, 0.1431, 0.1604$ for *Otx2*, *Dnmt3b* and *Fgf5*, respectively) except for *Wnt3* (d7 $p = 0.135$). **(C)** Analysis of H3K9me3 at *Nanog* in $\Delta K9.1$ and $\Delta K9.2$ cells analysed and presented as in Fig. 3B, showing a lack of enrichment throughout differentiation.

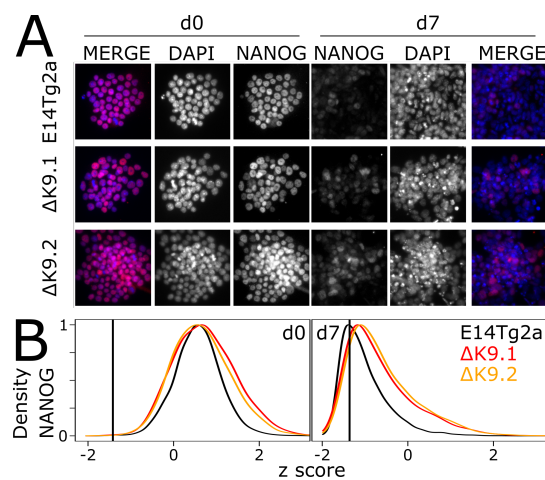


Fig. S4. General retention of low NANOG levels in differentiating Δ K9 cells.

(A) Representative illustration of undifferentiated (left) and differentiated (right; 7d of 2iLIF withdrawal) E14Tg2a, Δ K9.1 and Δ K9.2 cells stained for NANOG and DNA (DAPI).

(B) Quantification of NANOG levels in the conditions described in (A), presented as in Fig. 2C. The vertical line shows the median of differentiated E14Tg2a cells. Note that the vast majority of Δ K9 cells present higher NANOG levels than E14Tg2a cells after 7 days of differentiation (KS test $p < 2.2 \times 10^{-16}$).

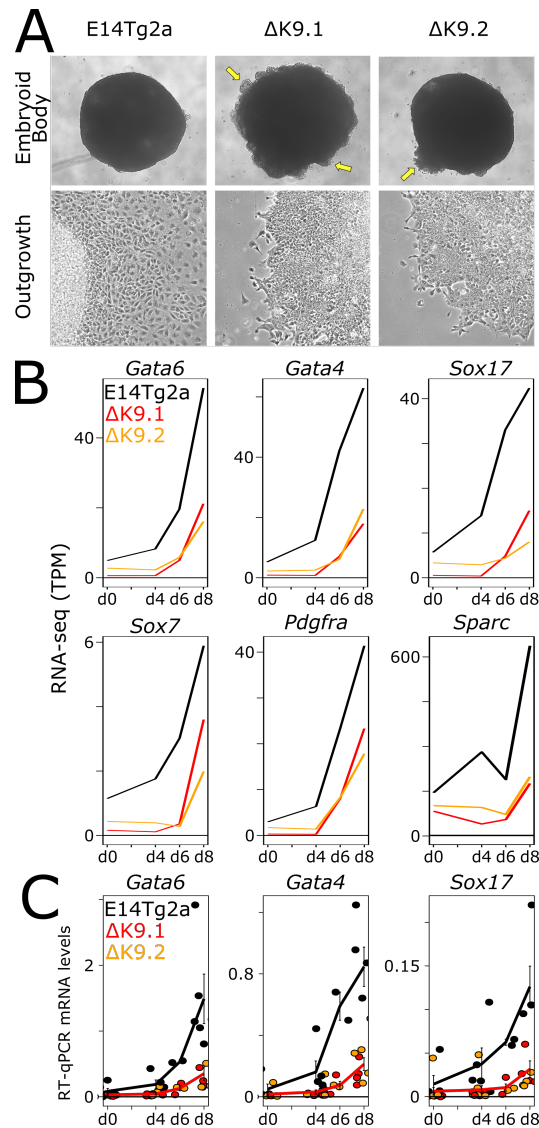


Fig. S5. Differentiation of wild-type and $\Delta K9$ cells into embryoid bodies.

(A) Representative bright-field photomicrographs of wild-type (E14Tg2a) and $\Delta K9$ embryoid bodies (top) and their derived cellular outgrowths (bottom). The arrows point to defective sealing of EBs periphery. **(B, C)** Analysis of endoderm markers after 0, 4, 6, 8 days of embryoid body differentiation, by RNA-seq (B) or by RT-qPCR (C). *Tbp* was used for normalisation.

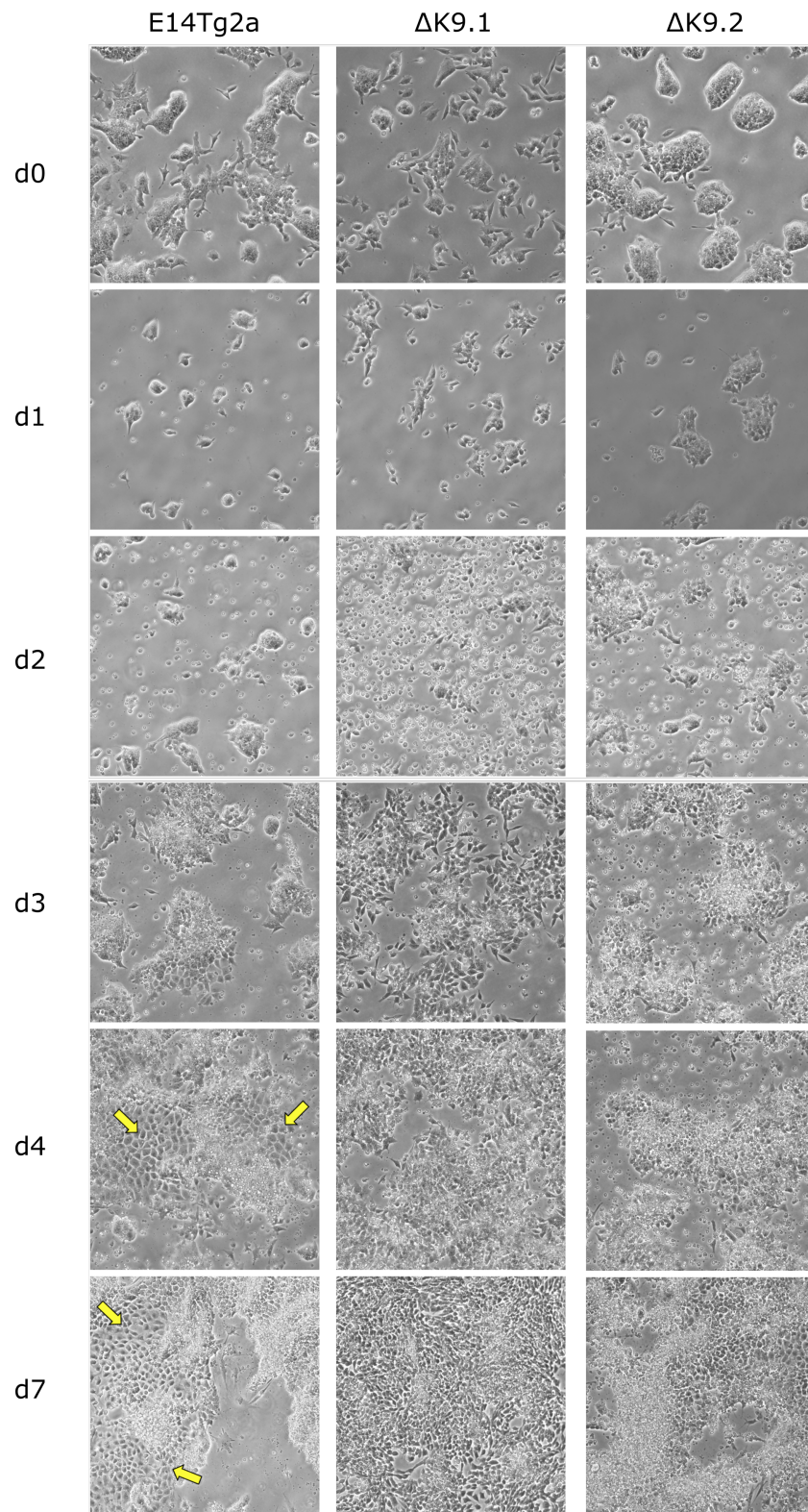


Fig. S6. Directed differentiation of wild-type and Δ K9 cells into primitive endoderm. Representative bright-field photomicrographs of wild-type (E14Tg2a) and Δ K9 cells during their treatment with a primitive endoderm differentiation protocol. The arrows point to clusters of primitive endoderm cells.

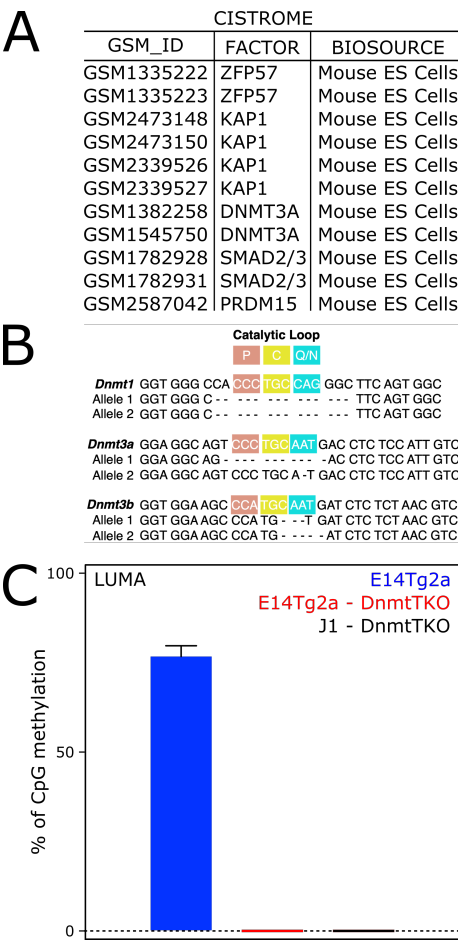


Fig. S7. Factors binding the IR region and generation of *Dnmts* TKO cells.

(A) The DNA sequence of the deleted region was used for a Cistrome analysis to identify candidate binding factors. The table shows those identified in ChIP-seq datasets performed in mouse ES cells. **(B)** Sanger sequencing tracks for each mutated allele of the three *Dnmt* genes. **(C)** LUMinometric Methylation Assay (LUMA) quantification of global CpG methylation levels in E14Tg2a WT and *Dnmt* TKO cells. DNA from a previously generated TKO line (J1 background; Tsumura et al., 2006) was used as a control. Data are shown as means \pm s.e.m for two independent replicates.

Table S1. RNA-seq counts and annotations.

[Click here to download Table S1](#)

Table S2. Primers and antibodies used in this study.

[Click here to download Table S2](#)

Supplementary Materials and Methods.

Cell Culture

General ES cell culture conditions.

ES cells (E14Tg2a; Δ K9 clones; TNG (Nanog-GFP reporter cells), a kind gift of Pr. Ian Chambers; TKO (triple knock-out DNMTs); ZKO (*Zfp57* knock-out), a kind gift of Basilia Acurzio & Andrea Riccio) were cultured (37°C, 7%CO₂) on 0.1% gelatine (SIGMA, Cat# G1890-100G) in DMEM, high glucose, GlutaMAX™ Supplement, pyruvate (Gibco, Cat# 31966-021), 10% FCS (Sigma, F7524), 100 μ M 2-mercaptoethanol (Gibco, Cat# 31350-010), 1 \times MEM non-essential amino acids (Gibco, Cat# 1140-035) and 10 ng/ml recombinant LIF (MILTENYI BIOTEC, Cat# 130-099-895). For TNG, when specified, puromycin was used at 2 μ g/mL. When indicated, cells were grown in serum-free 2i-containing medium (1 μ M PD0325901 and 3 μ M CHIR99021; Axon1408 & Axon1386 respectively): 0.5X DMEM/F12 (Gibco, Cat# 31331093), 0.5X Neurobasal (Gibco, Cat# 21103049), 0.5X N2 supplement 100X (Gibco, Cat# 17502048), 0.5X B27 supplement 50X (Gibco, Cat# 17504044), 10 μ g/ml Insulin (Sigma, Cat# I1882-100MG), 2 mM L-Glutamine (Invitrogen, Cat# 91139), 37.5 μ g/ml BSA (Sigma, Cat# A3311-10G), 100 μ M 2-mercaptoethanol (Gibco, Cat# 31350-010), 10 ng/ml recombinant LIF (MILTENYI BIOTEC, Cat# 130-099-895). Cells were seeded at 0.02M to 0.04M cells/cm², media changed every 2 days and cells passaged every 3-4 days. All 2i+LIF analyses were performed after at least 3-4 passages in 2i+LIF. All experiments inhibiting ERK or GSK3b were performed with 1 μ M PD0325901 and 3 μ M CHIR99021, respectively. Cells were karyotyped and regularly tested mycoplasma-free.

Culture conditions of MEFs, XEN and TS cells.

Mouse Embryonic Fibroblasts (MEFs) were derived from F1 129sv/129sv E13.5 male embryos and cultured (37°C, 7%CO₂), for no more than 4 passages, in DMEM, high glucose, GlutaMAX™ Supplement, pyruvate (Gibco, Cat# 31966-021), 10% FCS (Sigma, F7524), 100 μ M 2-mercaptoethanol (Gibco, Cat# 31350-010), 1 \times MEM non-essential amino acids (Gibco, Cat# 1140-035). When indicated, MEFs were treated for 3 days with 1 μ M PD0325901. Extra-Embryonic Endoderm (XEN) cell lines (Artus et al., 2010) were routinely cultured (37°C, 7%CO₂) in DMEM, high glucose, GlutaMAX™ Supplement, pyruvate (Gibco, Cat# 31966-021), 10% FCS (Sigma, F7524), 100 μ M 2-mercaptoethanol (Gibco, Cat# 31350-010), 1 \times MEM non-essential amino acids (Gibco, Cat# 1140-035) on 0.1% gelatine, without LIF, and passaged every 4 days. Trophectoderm Stem (TS) cell lines (F2 and F3) were initially isolated from F1 129/Sv Hprt-4 Pgk1a \times 129/Sv embryos using a published protocol (Tanaka et al., 1998) and cultured (37°C, 5%CO₂) in RPMI 1640 Medium, GlutaMAX™ Supplement (Gibco, Cat# 61870036) with 20% FCS (Sigma, F7524), 1 mM sodium pyruvate, 100 μ M β -mercaptoethanol (Gibco, Cat# 31350-010), 25 ng/ml hrFGF4 (235-F4-025, R&D) and 1 μ g/mL heparin (H3149-10KU, Sigma) on mitomycin-MEFs. Cells were passaged every 2-3 days. For TS analyses, MEFs were removed by adsorption onto gelatinized culture dishes for 1.5 hours.

Derivation of Δ K9 cells.

gRNAs (5'- CAGAGGAGGGCTTAAGAGAT and 5'- CACTCTAACCCAGCTTAAGT) were designed and cloned under the control of a U6 promoter in a vector conferring puromycin resistance, as described (Heurtier et al., 2019). 1 μ g of each gRNA-expressing vector and of a Cas9/mCherry expression vector (Addgene#64324) were lipofected in E14Tg2a cells according to manufacturer's instructions (Lipofectamine 2000; ThermoFisher). Lipofected cells were selected with Puromycin (1 μ g/ml) and FACS-sorted for mCherry fluorescence. Puromycin-resistant and mCherry-positive cells were seeded at clonal density and ~100 clones were picked 10 days later. Clones were screened by 2 independent PCR (LongAmp Taq PCR kit; BioLabs Cat# E5200S) with primers spanning the deletion (DelCTRL_F/R; N12c_F/N17b_R; Table S2), by real-time PCR using primers along the *Nanog* locus (Table S2), and by cloning and sequencing of PCR products (N12c/N17b). Two karyotypically normal independent clones, Δ K9.1 and Δ K9.2, were selected for this study.

Clonal assays.

600 cells were plated in single wells of 6-well plates coated overnight with poly-L-ornithine 0.01% (Sigma, Cat# P4957) at 37°C, washed twice with PBS 1X and coated 2 h with 1X laminin (Sigma, Cat# L2020). After 7 days, cells were fixed and stained using an alkaline phosphatase staining kit (Sigma Aldrich, Cat # 86R-1KT) according to the manufacturer's instructions or by fixation 20 min in PFA 4%, 2 washes with PBS 1X, incubation 15 min at RT in the dark in a staining solution (TrisMaleate 1M, MgCl₂ 1M, 100 mg/ml α -naphthyl-phosphate, 100 mg/ml Fast-Red TR), 2 washes with PBS 1X, 1 wash with milliQ-water. Colonies were counted under a stereo-microscope (NIKON-SMZ1500).

2i_{OFF} differentiation.

Cells were adapted from FCS+LIF to 2i+LIF for 3-4 passages, harvested and seeded at 10,000 cells/cm² on Poly-L-ornithine/Laminin coated cell culture treated surfaces, in 2i+LIF medium but omitting PD0325901, CHIR99021 and LIF. The medium was changed daily.

Embryoid Bodies differentiation.

Embryoid Bodies (EBs) imaged in Fig.S5A (top) were obtained with a hanging-drop protocol where a suspension of 0.1M cells/mL was distributed in 20 μ l drops (2000 cells/drop) onto inverted cover plates of several dishes filled with PBS 1X to avoid evaporation. The day after, the aggregates formed in each drop were pooled and seeded on non-cell culture treated dishes and cultured in 10%FCS-DMEM without LIF for 7 additional days. EB differentiation for outgrowths (Fig.S5A, bottom) and RNA-seq analysis (Fig.S5B) was performed by seeding 0.04M cells/cm² in 10%FCS-DMEM/LIF on 0.1% gelatinized cell culture treated dishes. After 3 days, the aggregates were harvested (2min trypsinization), transferred to non-cell culture treated dishes for 4 days in 10%FCS-DMEM medium without LIF and then replated at very low density on 0.1% gelatinized cell culture treated dishes for 2 and 4 additional days in 10% FCS-DMEM medium without LIF.

Primitive endoderm differentiation.

ES cells cultured in FCS+LIF were harvested and seeded onto 0.1% gelatinized single wells of μ -slide 4 well^{Ph+} ibiTreat (ibidi GmbH Ref#80446) at 37000 cells/cm² and differentiated as previously described (Anderson KGV et al., 2017). They were first cultured for 24H in endoderm base medium (EBM): RPMI 1640 Medium, GlutaMAXTM Supplement (Gibco, Cat# 61870036) supplemented with 2% B-27 minus insulin (Gibco, Cat# 15285074) and 100 μ M 2-mercaptoethanol (Gibco, Cat# 31350-010). Subsequently, Activin A (100 ng/ml; R&D, Cat# 338-AC-010), CHIR99021 (3 μ M) and LIF (10 ng/ml) were added (inductive PrE medium). The medium was changed every day.

Commitment assays.

Cells cultured in 2i+LIF were plated at clonal density (600 cells per well of 6-well plates) and subject to 2i_{OFF} differentiation, as described above. After each day, the medium was changed into 2i+LIF for 7 days, after which cells were fixed and stained for alkaline phosphatase activity as described.

Preparation of mitotic cells.

To obtain mitotic ES cells (>95% purity as assessed by DAPI staining and microscopy), we used a nocodazole shake-off approach, as described before (Festuccia et al., 2018).

Imaging analyses.

Bright field microscopy.

Cell culture pictures were taken on a Nikon Eclipse Ti-S inverted microscope equipped with: CFI S Plan Fluor ELWD $\times 20$ objective; 89 North PhotoFluor LM-75; Hamamatsu ORCA-Flash 4.0LT camera; NIS Elements 4.3 software.

Immunofluorescence of NANOG/OCT4 in E14Tg2a cells.

Cells were trypsinized, counted and fixed for 10 min in 4% formaldehyde (Sigma Cat#F8775) at room temperature and quenched immediately after with Glycine 125 mM for 5 min at room temperature. After one wash in PBS 1X, they were cytospun at 0.5M cells/200 μ l/spot (4 min – 300 rpm – low speed) on Superfrost+ slides. After washing the cells twice in PBS 1X, they were permeabilized with cold PBS 1X/0.5 % v/v Triton X-100 for 5 min, washed twice with cold PBS 1X, and blocked in PBS1X/1% Donkey Serum (DS) (Sigma, Cat# D9663) for 30 min on ice. Cells were incubated overnight at 4°C with primary antibodies (diluted in PBS1X/1% DS) within a humid chamber. After three washes in cold PBS1X, cells were incubated 1H at room temperature in the dark with secondary antibodies (diluted in PBS1X/1%DS), washed three times in PBS1X and nuclei counterstained with Vectashield antifade mounting medium with DAPI (Vectorlabs, Cat#H-1200). Imaging was performed with an inverted Nikon Eclipse X microscope equipped with: X20/0.45 (WD 8.2-6.9) objective; LUMENCOR excitation diodes; Hamamatsu ORCA-Flash 4.0LT camera; NIS Elements 4.3 software. Cell Profiler (Carpenter AE et al., 2006) was used for quantifications and ggplot2 (Wickham et al. 2016) for plotting in R.

c3/ Comparative immunofluorescence of NANOG (E14Tg2a vs mutant cells Δ K9, TKO or ZKO).

WT and each mutant cell line were trypsinized, counted and resuspended at 1M/ml in FCS free medium (DMEM-Glutamax/100 mM 2-mercaptoethanol/NEAA 1X) into sterile 1.5 mL Eppendorf tubes. Cells were then individually incubated either with 10 μ M Rhodamine Red dye (Invitrogen, Cat#CMTPX C34552) or 1 μ M Deep Red dye (Invitrogen, Cat#C34565) for 30 min at 37°C. The labeled cells were then collected by centrifugation, washed with PBS1X, resuspended in DMEM/10%FCS+LIF medium and mixed at a 1:1 ratio for WT^{Rhod} and mutant^{DeepRed} cells (usually \sim 0.4M each). The opposite labeling (WT^{DeepRed} & mutant^{Rhod}) was also performed with identical results. Mixed cells were seeded into a Poly-L-Ornithine/Laminin coated single well of a μ -slide 4 well^{Ph+} ibiTreat (ibidi GmbH Ref#80446) and incubated for \sim 6H at 37°C and 7% CO₂. Cells were then fixed directly into the well by freshly prepared PFA 4% (Fisher Scientific, Cat# 16431755) for 10 min at room temperature in the dark, washed twice in PBS1X for 10 min and used immediately for immunostaining or stored at +4°C for short time. Cells were washed in PBS-Tw0.1% (PBS1X/0.1 % Tween20 (Sigma, Cat#P9416)) and permeabilised with PBS1X/0.1% Triton X-100 (Sigma, Cat#T8787) for 10 min at room temperature. After three washes with PBS-Tw0.1%, cells were blocked with PBS-Tw0.1%/10% Donkey Serum (Sigma, Cat#D9663) for 30 min on ice in the dark and incubated overnight with the primary antibody (diluted in PBS-Tw0.1%/10% DS). Following three washes with PBS-Tw0.1%, 1H incubation with secondary antibodies at room temperature in the dark and 2 washes with PBS1X, nuclei were counterstained with 4',6-diamidino-2-phenylindole (DAPI; Sigma, Cat# D9542), washed in PBS1X and use immediately for imaging or stored at +4°C for short time. Imaging was performed with an inverted Nikon Eclipse X microscope equipped with: X20/0.45 (WD 8.2-6.9) objective; LUMENCOR excitation diodes; Hamamatsu ORCA-Flash 4.0LT camera; NIS Elements 4.3 software. Quantifications were performed using Cell Profiler (Carpenter AE et al., 2006). For each experiment, E14Tg2a and each mutant cells quantifications were attributed using the FlowJo software. For each experiment, the fluorescence intensity of each mutant cell was normalised to the median of the corresponding E14Tg2a intensities imaged on the same spot. The data was plotted using the ggplot2 package (Wickham, 2016) in R.

Immunofluorescence of NANOG and primitive endoderm markers.

Differentiated cells were processed into their respective well: one wash with PBS1X, fixation with freshly prepared PFA 4% for 10 min at room temperature, three washes of PBS1X, and used immediately for immunostaining or stored at +4°C for short time. Immunostaining was performed as in c3/ for NANOG in combination with primitive endoderm markers. Imaging was performed with an inverted Nikon Eclipse X microscope equipped with a x20 objective; LUMENCOR excitation diodes; Hamamatsu ORCA-Flash 4.0LT camera; NIS Elements 4.3 software. For each well, ~30 pictures were taken by randomly scanning the well. Quantifications were performed using Cell Profiler (Carpenter AE et al., 2006) and plotted using the ggplot2 package (Wickham et al. 2016) in R. Representative images were generated with ImageJ software using identical settings for

Single-molecule RNA Fluorescent In Situ Hybridisation (smFISH).

Cells were grown at low density, put in no RED medium for 24H before collecting them by regular trypsinization and resuspended in DMEM/10% FCS medium. Cells were fixed with 3.7% Formaldehyde (Sigma F8775) for 10 min at RT and the reaction was stopped by addition of 0.125M glycine (SIGMA G7126) for 5 min at RT. Cells were then centrifugated at 1000 rpm at 4°C, washed once in cold PBS/0.1% BSA, resuspended in cold PBS/0.1% BSA/1 mM EDTA at 1 million cells/mL and cytospun at 500 rpm (low speed) for 4 min on SuperFrost slides (Thermo J1800AMNT) – around 500K cells/spot in order to obtain around 40-60 nuclei/field (obj.x60) and no or few clumps. Slides were air dried for 1 min and stored in 70% EtOH at 4 °C for at least one night. The slides are dehydrated for 3 min in 100% EtOH and fully air dried. Each spot was hybridized for 24H at 37°C in a denaturing chamber (2X SSC/10% formamide) with the *Nanog* mRNA probe cocktail (2X SSC, 10% Formamide, BSA 1 µg/mL, 1 µL of *E.Coli* RNAs at 1 µg/mL, 1 µL of probe at 0.25 µM). The slides were washed for 1H at 37 °C in 2X SSC/10% Formamide then 2 times in 2X SSC for 10 min and mounted in Vectashield medium with DAPI (Vector-abcys H-1200). The *Nanog* mRNA probe was designed using Stellaris Probe Designer version 4.2 on Biosearch Technologies website with the maximum masking level (5) and was synthesized by the same company. Image stacks (0.5 µm gap) were acquired using a Nikon Eclipse X microscope equipped with: x63 oil immersion objective (N.A1.4); LUMENCOR excitation diodes; Hamamatsu ORCA-Flash 4.0LT camera; NIS Elements 4.3 software. The analysis was performed using ImageJ to identify smFISH spots using the Find Maxima function with adapted Noise tolerance and maximal projections of the stacks. The identified maxima were converted into grays scale images and, together with minimal projections of the DAPI staining, were imported into Cell Profiler (Carpenter et al., 2006) to count the number of smFISH spots per DAPI-segmented cell. The results were imported into R to compute the frequency of spot detection per cell and plot them with ggplot2 (Wickham, 2016).

Chromatin analyses.*Chromatin preparation.*

After trypsinisation, 10⁷ ES cells were crosslinked for 10 min in 3 ml DMEM/10% FCS/1% formaldehyde (Sigma Cat#F8775). Crosslinking was stopped with 125 mM glycine for 5 min at room temperature then 5min in ice. Cells were pelleted and washed with ice-cold PBS1X. Cells were resuspended in 1 ml of ice-cold swelling buffer (25 mM Hepes pH 7.95, 10 mM KCl, 10 mM EDTA) freshly supplemented with 1× protease inhibitor cocktail (PIC-Roche, Cat# 04 693 116 001) and 0.5% IGEPAL (Sigma, Cat#18896). After 20 min on ice, the suspension was passed 50 times in a dounce homogenizer. Cells were then centrifuged and resuspended in 1 ml of ice-cold D3 (0.1% SDS, 15 mM Tris pH 7.6, 1 mM EDTA) buffer, freshly supplemented with 1× PIC. Samples were sonicated using a Covaris M220 - Setpoint @6°C and 10 cycles with the following parameters for each cycle : 60 sec duration, peak

power of 67W, duty factor of 15% and cycles/burst of 500. A delay of 45 sec is added at the end of each cycle. Result of the average power is 10W. After centrifugation (15 min, 14000 rpm, 4 °C), the supernatant was stored at -80 °C until use. 20 µl were used to quantify the chromatin concentration and check DNA size (typically 200-600 bp).

Immunoprecipitation.

15 to 20 µg of chromatin were used for each ChIP after pre-clearing it for 1.5 hours rotating on-wheel at 4 °C in 1 ml of TSE150 (0.1% SDS, 1% Triton X-100, 2 mM EDTA, 20 mM Tris-HCl pH8, 150 mM NaCl) buffer containing 50 µl of protein G Sepharose beads (Active Motif, Cat#37499) 50% slurry, previously blocked with BSA (0.5 mg/ml ; Roche, Cat# 10711454001) and yeast tRNA (1 µg/ml ; Roche Cat# 10109495001). Immunoprecipitations were performed overnight rotating on-wheel at 4 °C in 500 µl of TSE150. 20 µl were set apart for input DNA extraction and precipitation. 50 µl of blocked protein G beads 50% slurry was added for 2 h rotating on-wheel at 4 °C. Beads were pelleted and washed for 5 min rotating on-wheel at room temperature with 1 ml of buffer in the following order: 2 × TSE150, 1 × TSE500 (as TSE150 but 500 mM NaCl), 1× washing buffer (10 mM Tris-HCl pH8, 0.25M LiCl, 0.5% IGEPAL, 0.5% Na-deoxycholate, 1 mM EDTA), and 2 × TE (10 mM Tris-HCl pH8, 1 mM EDTA). Elution was performed in 100 µl of elution buffer (1% SDS, 10 mM EDTA, 50 mM Tris-HCl pH 8) for 15 min at 65 °C after vigorous vortexing. Eluates were collected after centrifugation and beads rinsed in 150 µl of TE-1%SDS. After centrifugation, the supernatant was pooled with the corresponding first eluate. For both immunoprecipitated and input chromatin, the crosslinking was reversed overnight at 65 °C, followed by proteinase K treatment, phenol/chloroform extraction and ethanol precipitation.

qPCR analysis.

Input and IP samples were analysed by real-time quantitative PCR performed in duplicates in 384-well plates with a LightCycler 480 (Roche) using 4.6 µl of LightCycler 480 SYBR Green I Master (Roche, 04707516001), 5 µl of sample and 0.2 µl of each primer at 20 µM in a final reaction volume of 10 µl. Standard and melting curves were generated to verify the amplification efficiency (>85%) and the production of single DNA species. PCR primer sequences are listed in Table S2. The 2dCt method was used. All values were corrected to the input and plotted in R using ggplot2 (Wickham et al., 2016).

Antibodies. Antibodies used for all ChIP experiments are listed in Table S2.

Gene expression analyses.

RNA preparation, RT-qPCR and sequencing.

E14Tg2a and ΔK9 cells were differentiated in parallel as embryoid bodies and cells recovered at d0, d4, d6, d8. RNA extraction and DNase treatment of 3 independent assays were made with NucleoSpin RNA Mini kit (Macherey Nagel, Cat# 740955.50) according to the manufacturer's protocol. Reverse Transcription was performed with 1 µg of total RNAs with random hexamers following manufacturer's protocol (Roche 04379012001); qPCR was performed as described above (*Tbp* was used as a reporter; see Table S2 for primer sequences). Stranded, poly-A selected RNA-seq libraries were prepared and sequenced (paired-end 150bp reads; around 50 millions each) by Novogene Co Ltd.

Alignments and quantification.

Stranded paired end RNA-seq reads were aligned to the mm10 genome using STAR (Dobin et al., 2013) and quantified by RSEM (Li and Dewey, 2011), with additional options " --calc-pme --calc-ci --estimate-rspd --forward-prob 0.0 --paired-end". Transcripts per million (TPM) were computed after omitting outliers identified by Principal Component Analysis (30 or more standard deviations away from the mean for PC1 component).

Principal component analysis.

PCA was run in R (prcomp function; with option center=TRUE) using log2 transformed TPM after selecting genes with at least 1 TPM in at least one sample mean (n=3). The top 1000 PC1 loadings (55% of variance) were used for gene ontology analyses. Data visualisation was made in base R.

Differentially expressed genes.

For all differential expression tests RSEM estimated read counts per sample were rounded for use with DESeq2 (Love et al., 2014), which was run without independent filtering. For differentially expressed genes in undifferentiated ES cells, we selected those with an FDR < 0.05 in Δ K9.1 versus E14Tg2a and Δ K9.2 versus E14Tg2a and same fold-change direction with $\text{abs}(\log_2\text{FC}) > 0.3$. For embryoid body differentiation, we considered genes with absolute $\log_2(\text{FC}) > 1$ and FDR < 0.05 at any day of the differentiation versus undifferentiated cells for either E14Tg2a or Δ K9 cells. The corresponding heatmap was made in R with ComplexHeatmaps package (Gu et al., 2016).

Clustering of differentially expressed genes and developmental annotation.

K-means clustering was computed with R using the function kmeans with options k=6, nstart=50, iter.max=50. Only differentially expressed genes as identified during embryoid body differentiation were used (z scored mean TPM). The number of clusters was chosen as the minimal value identifying at least one cluster with maximal expression at each day of differentiation, including d0. Correlations to developmental gene expression were made by directly plotting the $\log_2(\text{FC})$ reported in a previous study using SCNMT-seq around gastrulation on mouse embryos (Argelaguet et al., 2019). The data was downloaded from ftp://ftp.ebi.ac.uk/pub/databases/scnmt_gastrulation. The heatmap was made in R with ComplexHeatmaps package (Gu et al., 2016) and the boxplot with ggplot2 (Wickham, 2016).

Enrichment analyses.

Gene Ontology analyses were performed using Enrichr (<https://maayanlab.cloud/Enrichr/>) and the top association was selected. To determine enrichments of each group of differentially expressed genes in proximity to TF binding sites for Nanog and Oct4/Sox2 (Festuccia et al., 2019), we calculated Fisher tests right tail p-values for the association between differentially expressed genes of each cluster within xbp of a binding site to a background of all genes clustered within xbp of a binding site, for x in [1, 1e+9] bp. Data visualization was made in base R.

Supplementary References.

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