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Targeted disruption of *DNMT1*, *DNMT3A* and *DNMT3B* in human embryonic stem cells

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Abstract

DNA methylation is a key epigenetic modification involved in regulating gene expression and maintaining genomic integrity. Here we inactivated all three catalytically active DNA methyltransferases in human embryonic stem cells (ESCs) using CRISPR/Cas9 genome editing to further investigate their roles and genomic targets. Disruption of *DNMT3A* or *DNMT3B* individually, as well as of both enzymes in tandem, creates viable, pluripotent cell lines with distinct effects on their DNA methylation landscape as assessed by whole-genome bisulfite sequencing. Surprisingly, in contrast to mouse, deletion of *DNMT1* resulted in rapid cell death in human ESCs. To overcome the immediate lethality, we generated a doxycycline (DOX) responsive tTA-*DNMT1** rescue line and readily obtained homozygous *DNMT1* mutant lines. However, DOX-mediated repression of the exogenous *DNMT1** initiates rapid, global loss of

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DNA methylation, followed by extensive cell death. Our data provide a comprehensive characterization of *DNMT* mutant ESCs, including single base genome-wide maps of their targets.

Introduction

DNA methylation is a heritable and reversible enzyme-mediated modification to DNA that is implicated in a diverse range of biological processes[1]. In mammals, there are three catalytic active DNA methyltransferases (DNMTs): DNMT1, which is largely responsible for the maintenance of DNA methylation over replication, and DNMT3A and DNMT3B, which generally perform *de novo* methylation of either unmethylated DNA or hemimethylated DNA to assist in maintenance[2]. Deletion of these enzymes in mice results in embryonic (*Dnmt1* and *3b*) or postnatal (*Dnmt3a*) lethality, confirming their essential roles in development[3,4]. In line with knockout mouse models, mutations affecting the human DNA methyltransferases are linked to several diseases, including acute myeloid leukemia (AML) and Immunodeficiency, centromere instability and facial anomalies (ICF) syndrome[5–8].

DNA methylation has been extensively mapped in somatic cells as well as ESCs in human and mouse[9–14]. Human ESC methylation patterns are most unique at hypomethylated regulatory elements that are enriched for binding of pluripotency-associated master regulators, such as OCT4, SOX2, and NANOG[14]. Additionally, both mouse and human ESCs display increased levels of non-CpG methylation that is not observed in most somatic lineages, consistent with the high activity of *de novo* methyltransferases in these cells[11,15–17].

While deletion of *Dnmt1* is lethal in all dividing somatic cells[3,18–21], mouse ESCs are viable despite global loss of DNA methylation. In fact, all three *Dnmts* can be removed from these cells without any deleterious effects in the undifferentiated state[22]. As such, mouse ESCs have become a powerful tool to study the role and function of *Dnmt* enzymes, which clarified some of their specific targets and provided many general insights into the biology of DNA methylation[23]. No comparable efforts have been reported for human pluripotent stem cells and loss of function studies have been limited to the depletion of *DNMT1* in the colon cancer cell line HCT116, which results in cell death[24,25] and therefore indicates a similar requirement for maintenance of DNA methylation patterns in human cells. DNMT3B was reported to cooperate with DNMT1 to maintain methylation in HCT116 cells[26,27] and its depletion results in altered timing of neuronal differentiation and maturation[28]. Recently, a human ESC model for ICF syndrome was reported by targeted disruption of *DNMT3B*[29].

The limited number of human studies can likely be attributed to long-standing difficulties of genetic manipulation in the human ESC system, rather than lack of interest. However, new classes of genome-editing tools, transcription activator-like effector nucleases (TALENs) and clustered regularly interspaced short palindromic repeats (CRISPR)/CRISPR-associated (Cas) systems have been applied for genomic editing in mammalian cells[30–34]. These techniques have substantially enhanced targeting efficiency into human ESCs, allowing previously difficult studies to be undertaken[33,35].

Here we report the inactivation of all three DNMT*s* in human ESCs and present a detailed analysis of the DNA methylation changes in *DNMT1*, *DNMT3A*, *DNMT3B* and tandem *DNMT3A/3B* double knockouts (*DNMT3A−/−3B−/−*). We further characterized the differentiation potential of the knockout lines using gene expression signatures, EB differentiation and teratoma formation. We also used directed differentiation towards the endoderm lineage to assess the role of DNMT3A in this transition. Most notably, we were unable to obtain *DNMT1* homozygous deletions without implementing a doxycycline-repressible rescue line, demonstrating that loss of DNMT1 is lethal. Taken together, our results highlight several unique aspects of DNA methylation biology in the context of human ESCs and provide controlled, tractable systems to dissect the function of DNMTs in precise detail.

Results

Disrupting the catalytic domain of all three *DNMT*s

Of the three catalytically active DNMTs, *DNMT3B* shows the highest level of expression in undifferentiated ESCs (Fig. 1a), but also the most variation when examined across 25 pluripotent lines (Fig. 1b). As ESCs differentiate, *DNMT1* and *DNMT3A* remain expressed at comparable levels, while *DNMT3B* is strongly downregulated and switches to predominant expression of a catalytically inactive isoform (isoform *3B1* NM_006892 to isoform *3B3* NM_175849; Fig. 1a, Supplementary Fig. 1a). While inactive, this isoform can likely still form complexes with catalytically competent DNMT3A and/or DNMT3B to contribute to DNA methylation activity[36].

To better understand the role of the three *DNMT*s in human pluripotent stem cells and over cellular differentiation, as well as to study their specific target spectrum at high-resolution across the genome, we created targeted deletions for *DNMT1*, *DNMT3A*, *DNMT3B* and both *DNMT3A* and *3B* in human ESCs. We selected the male line HUES64 for the following reasons: (i) it is on the NIH registry and generally available to researchers, (ii) it differentiates well into the three germ layers, (iii) it is karyotypically normal (Supplementary Fig. 1b) and grows well under standard culture conditions, and (iv) a substantial amount of publicly available transcriptional, epigenomic and transcription factor binding data have been generated for this line[9,37].

In order to avoid hypomorphic effects[24], we designed guide RNA G(N)19NGG sequences targeting the catalytic domains for *DNMT1*, *3A* and *3B* (Fig. 1c). After picking and expanding individual clones, we confirmed the targeted disruption of the catalytic domains by Sanger sequencing (Supplementary Fig. 1c, d). All experiments yielded high rates of mutations ranging from 53% to 66%, with homozygous deletion occurring in 3–6% of the clones for *DNMT3A* and *B* (Supplementary Fig. 1c). Double knockout cells were derived by targeting the already validated *DNMT3A−/−* clone (#139) with the *DNMT3B* gRNA and Cas9. To further validate the *DNMT3* knockouts, we performed qPCR (Fig. 1d) and western blotting (Fig. 1e). Notably, while we obtained homozygous knockouts for *DNMT3A* and *DNMT3B*, both singly and in combination, we were unable to obtain any *DNMT1−/−* clones, despite high (>50%) efficiency when deriving heterozygous lines.

DNMT3 knockout ESCs remain pluripotent

All of our *DNMT3* knockout clones grew well and were morphologically normal. We selected representative clones for each knockout (*DNMT3A*^{−/−}: #139; *DNMT3B*^{−/−}: #44; *DNMT3A*^{−/−}*3B*[−/−]: #21) and performed several assays to characterize these knockouts, both in terms of their molecular phenotype as well as of their differentiation potential (Fig. 2a). The clones were positive for the pluripotency-associated markers, *NANOG* and *TRA1-60* (Fig. 2b) and showed tri-lineage differentiation potential based on the TaqMan® hPSC Scorecard™ (Fig. 2c; Supplementary Fig. 2)[38]. We differentiated these clones into embryoid bodies and the resulting cells were stained for markers of the three germ layers (Fig. 2d). Finally, we injected each clone into the kidney capsule of three independent NOD-SCID mice to test for teratoma formation (Fig. 2e) and confirmed the presence of cell types representing the three germ layers in several sections per knockout (Fig. 2f). These results show that deletion of the *DNMT3* enzymes, either individually or in combination, has no obvious effect on morphology, viability and initial differentiation potential in human ESCs. This observation is consistent with prior studies of the respective knockouts in mouse ESCs, where both individual knockouts can differentiate, although the mouse double knockouts could only generate teratomas at early passage numbers[4,39].

Passaging results in progressive loss of DNA methylation

To characterize the effects of the *DNMT3* knockouts on the human ESC methylome, we harvested genomic DNA from our mutant lines at an early (2–7) and late (17–22) passage and generated WGBS libraries (Fig. 3a). We utilized the methylation status of 6.9 million CpGs that were covered at 5x in all eight samples for the initial comparative analysis. Hierarchical clustering of the WGBS data shows that WT, *DNMT3A*^{−/−} and *DNMT3B*^{−/−} cells cluster closely together, but remain clearly segregated nonetheless (Fig. 3b, left). Despite being derived from the *DNMT3A*^{−/−} clone, and therefore beginning in a DNMT3A-depleted background, the double knockout is clearly distinguishable, highlighting that complete *de novo* methyltransferase ablation dominates effects on global DNA methylation levels over those initiated by the original single knockout. To further assess the effects of the knockouts and of passage, we performed principal component analysis (Fig. 3b, right). The first two principal components explain 96% of the variation between the samples and separate them into two main groups: the first consists of the wild-type human ESCs, the *DNMT3A* and *DNMT3B* single knockouts, while the second includes the early and late passage double knockouts.

We then analyzed the WGBS data from the later passage samples, which captured 14.8 million CpGs covered at 5x coverage in all four samples, to see if specific genomic features were particularly sensitive to the loss of *de novo* DNA methyltransferase activity. We divided all CpGs into three classes according to their methylation values—high ($\geq$0.8), intermediate (>0.2 and <0.8) or low ($\leq$0.2)—and examined the relative proportion falling into each category (Fig. 3c). The fraction of intermediate and low methylated regions in the knockouts gradually increases over passage, especially when comparing *DNMT3A*^{−/−}*3B*[−/−] versus WT, which is similar to equivalent mouse knockout models (Supplementary Fig. 3a). Most genomic features show a mild decrease in DNA methylation in the single and a more

substantial change in the double knockouts, with the exception of satellites, which appear particularly sensitive to the loss of *DNMT3B* (Fig. 3c, Supplementary Fig. 3b–d).

We next divided the genome into 1kb tiles and compared the distribution of CpG methylation values within each tile across the samples (Fig. 3d). We defined differentially methylated regions (DMRs) as those tiles that were significantly different (weighted t-test, q value < 0.05) between the wild-type and any of the knockouts and had an average methylation difference greater than 0.4, ignoring repetitive tiles. These DMRs can be divided into four classes: (i) DNMT3A specific targets, (ii) DNMT3B specific targets, (iii) shared targets that require both DNMT3A and DNMT3B, (iv) redundant targets that require either DNMT3A or 3B and are only lost on *DNMT3A$^{-/-}$3B$^{-/-}$* cells. As suggested by the CpG level analysis above, almost all DMRs (~96%) are redundantly targeted by DNMT3A and DNMT3B and only lose methylation when both are ablated. For each class, we calculated the enrichment of different genomic features. Redundant targets are notably broad in scope and are enriched for intergenic regions, CpG island shores, intermediate and low CpG-density promoters, and introns (Fig. 3e). In contrast, the other three classes are largely enriched for CpG islands (Supplementary Table 1). These enrichments suggest that *DNMT3A* and *DNMT3B* appear to act redundantly in regions of low CpG density, but may be more selectively targeted to regions of high CpG density.

The individual knockouts show a mild global reduction, an effect that becomes slightly more pronounced in the double knockouts (Fig. 3f). In contrast to this limited global decrease, several loci, including *DPPA3* (*STELLA*), show complete loss of methylation in the *DNMT3A$^{-/-}$3B$^{-/-}$* cells (Fig. 3g), and form the set of redundant targets identified above. Another interesting pattern is seen at the *NANOG* promoter, which seems to be sporadically targeted by both *de novo* DNMTs, leading to low level methylation in the upstream region that is lost only in the *DNMT3A$^{-/-}$3B$^{-/-}$* cells (Supplementary Fig. 3c). *NANOG* is highly expressed, but the low level methylation in wild-type cells may reflect its known transcriptional heterogeneity within the ESC population.

We continued passaging of the *DNMT3A$^{-/-}$3B$^{-/-}$* cells for 22 passages, and profiled methylation over time using RRBS. In the absence of the *de novo* activity, CpG methylation is lost gradually due to the imperfect maintenance activity of *DNMT1* (44.5% to 35.4%; Fig. 3h). Using an estimated doubling time of 28.8 hours, we estimate the fidelity of *DNMT1* to be between 99.7% and 99.97%, with some noteworthy differences across genomic features (Supplementary Fig. 3b). In particular, features that are highly methylated and/or require continued repression by DNA methylation, such as transposable elements and satellite repeats[40], seem to exhibit slightly superior maintenance than other genomic features.

Lastly, we investigated the global level of non-CpG methylation, which is known to be present in mouse[15] and human[11,17] ESCs. Consistent with our previous knockdown based experiments[17], we find that DNMT3B depletion has a stronger effect than loss of DNMT3A (Fig. 3i, left). This could, however, simply be related to the comparatively high levels of *DNMT3B* in undifferentiated ESCs (Fig. 1a and b). In the *DNMT3A$^{-/-}$3B$^{-/-}$* cells, non-CpG methylation is rapidly reduced to background levels (Fig. 3i, right), confirming that continuous *de novo* DNA methyltransferase activity is required for non-CpG methylation in

dividing cells, while CpG methylation can still be largely propagated by DNMT1, with a slow, gradual decline due to imperfect maintenance over successive cell divisions (Fig. 3h).

Shared and unique targets of *DNMT3A* and *DNMT3B*

As noted above, we found a strong enrichment of CpG islands in the set of redundant DNMT3A/3B targets (Fig. 3e). To better understand this observation and gain a global perspective on the effect of DNA methylation on these targeted CpG islands and their neighboring regions (Supplementary Fig. 4a), we generated composite plots (Fig. 4a) and aggregate heatmaps (Supplementary Fig. 4b) for mean methylation levels across 25,489 CpG islands that were experimentally verified to be hypomethylated in somatic tissues[41]. We see a clear distinction in behavior between CpG islands that are hypomethylated in the knockouts, and the background set of CpG islands that are hypomethylated in both wild-type and knockout lines. These CpG islands are presumably specific targets of *de novo* methylation activity, since they become hypomethylated in the knockouts and then exhibit canonical CpG island methylation levels. In the *DNMT3A⁻/⁻3B⁻/⁻* cells, we see not only the focal loss of methylation seen at the center of the island, but also a general decrease in methylation at CpG island shores. These results are consistent with the enrichment results and suggest that DNMT3A and DNMT3B are specifically targeted to a subset of CpG islands.

Using our sequencing based data allows us to investigate the DNA methylation state of neighboring CpGs and classifying them into concordant or discordant patterns (Fig. 4b). We can then calculate the proportion of discordant reads (PDR) as the number of discordant reads divided by the total number of reads covering a CpG and plot the global distribution of this measure against CpG methylation[42]. In contrast to the single knockouts, the *DNMT3A⁻/⁻3B⁻/⁻* cells show a global increase in PDR (Fig. 4c), which suggests that DNMT3A and DNMT3B have similar mechanisms of action, and are likely to act redundantly to counteract global methylation loss due to imperfect DNMT1 fidelity. As expected, regions of low methylation in wild-type cells have low PDR, and these regions remain hypomethylated with low PDR in the knockouts (Fig. 4d, top panels). Regions of intermediate methylation show decreased methylation and lower PDR in knockout cells, with especially striking hypomethylation in the *DNMT3A⁻/⁻3B⁻/⁻* cells (Fig. 4d, middle panels). One exception are imprinting control regions that are known to be variable in ESCs but generally maintain stable, intermediate levels even in the *DNMT3A⁻/⁻3B⁻/⁻* cells (Supplementary Fig. 4c). Finally, regions of high methylation have low PDR in wild-type cells and the single KOs. These regions lose methylation specifically in the *DNMT3A⁻/⁻3B⁻/⁻* cells, with a corresponding increase in PDR (Fig. 4d, bottom panels).

To further assess the functional significance of the DMRs found above, we calculated the degree of overlap of the DMRs in the three knockout lines with published transcription factor binding sites[43] and compared this overlap to that seen with a background set of all 1kb tiles with an average methylation of at least 0.4 in wild-type cells (Fig. 4e). Both *DNMT3A⁻/⁻* and *DNMT3B⁻/⁻* DMRs show enrichment for the PRC2 components EZH2 and SUZ12 (Fig. 4e). The *DNMT3A⁻/⁻3B⁻/⁻* DMRs shows strong enrichment for many sequence-specific transcription factors, including CTCF, which is known to be impacted

with DNA methylation[44]. To see whether these targets are associated with any specific biological processes we took the set of 30,220 hypomethylated DMRs from the *DNMT3A*^{−/−}*3B*[−/−] cells and used GREAT[45] to associate them with genes and then calculate enrichments for functional ontologies. As expected, these regions are highly correlated with a large spectrum of developmental processes including patterning, regionalization, growth factor stimulation and corresponding pathways (Supplementary Fig. 4d and Supplementary Table 2). The enriched categories suggest that the DMRs we identify are generally hypermethylated in wild-type pluripotent cells by DNMT3A and DNMT3B, and normally lose methylation in a lineage and tissue specific manner as cells differentiate.

DNMT3A deletion prevents gain of DNA methylation

To further investigate the knockout effects during differentiation, we derived early endodermal cells from the *DNMT3A*^{−/−} cells (Fig. 5a; Supplementary Fig. 5). We then generated WGBS data and performed hierarchical clustering analysis, which included undifferentiated and sorted endoderm cells (dEN) of both WT and *DNMT3A*^{−/−} cells (Fig. 5b). These can be clearly segregated and undifferentiated cells in each case cluster more closely with their differentiated endoderm derivatives. We identified non-repetitive 1kb tiles that have significantly different DNA methylation levels between wild-type ESCs and dEN cells to determine their comparable DNA methylation state in undifferentiated and differentiated *DNMT3A*^{−/−} cells (Fig. 5c). We identified 575 DMRs that gain DNA methylation in wildtype dEN and are enriched for promoters and gene bodies (Fig. 5d). Strikingly, most of these DMRs remain hypomethylated in the *DNMT3A*^{−/−} derived dEN cells, suggesting that DNMT3A is required for their *de novo* methylation. For instance, we found that previously reported regions upstream of *FOXA2* and downstream *DBX1*, respectively, were no longer methylated in the knockout cells (Fig. 5e). We also differentiated the cells further towards the hepatoblast stage and stained the population with antibodies for FOXA2 and HNF4A. The loss of *DNMT3A* had no immediately detectable negative impact on the downstream differentiation in this lineage (Fig. 5f). In summary, we detect a unique molecular role for DNMT3A during endoderm differentiation. Although we have not yet determined the functional relevance or downstream consequences of the targeted *de novo* methylation, our system provides a new, powerful platform to dissect these effects in this and other lineages.

DNMT1 is essential for human ESCs

As described above, we were unable to obtain homozygous mutant lines for *DNMT1*, raising the possibility that human ESCs grown in standard FGF conditions require DNA methylation for viability. This hypothesis was further supported by the failure to create straightforward homozygous mutants using an alternative TALEN based strategy (Supplementary Fig. 6a) and our inability to obtain any viable lines with efficient *DNMT1* knockdown despite having tested a total of nine different shRNAs (Supplementary Fig. 6b). Notably, the same TALEN based strategy readily generated *DNMT3A and 3B* homozygous lines, but no viable *DNMT1* homozygous lines emerged, consistent with the CRISPR experiments.

We therefore designed a rescue strategy that would enable the generation of homozygous mutant lines without the immediate lethality of DNMT1 loss (Fig. 6a). We first introduced two separate lentiviruses containing the tTA and TRE-*DNMT1** into the wild-type HUES64 line, creating a doxycycline (DOX) responsive (TET/OFF) *DNMT1** cell line that we could subsequently use for targeting (Fig. 6b and Supplementary Fig. 6c). We excluded the KOZAK sequence (GCCACC) upstream of the start codon to prevent overexpression to levels that might cause adverse effects prior to targeting and created five point mutations within the target sequence and the Protospacer Adjacent Motif (PAM) sequence that have no effect on the protein sequence (Fig. 6a and Supplementary Fig. 6d) to avoid the gRNA from disrupting the exogenous *DNMT1**.

Both qPCR and Western blot analysis confirm tight control of expression for the exogenous *DNMT1**. In the absence of DOX, we observed an approximately threefold increase of overall DNMT1 levels that can be fully suppressed within 24 hours after supplementation with DOX (Fig. 6c). We next used the CRISPR/Cas9 approach to knock out the endogenous *DNMT1* while maintaining expression of the exogenous *DNMT1**. We could now readily obtain homozygous mutant lines in two independent rounds of targeting (Supplementary Fig. 6e, f), suggesting that our rescue strategy maintains a sufficient level of DNMT1 activity needed for survival. We selected two representative clones (#111, #122) for more detailed characterization and also confirmed by qPCR that *DNMT1* targeting does not affect the expression of *DNMT3A* and *DNMT3B* (Supplementary Fig. 6g).

Loss of *DNMT1* causes global demethylation and cell death

DNMT1 homozygous knockout ESCs proliferated normally and could be maintained over several passages as long as the exogenous *DNMT1** was expressed. The cells displayed a normal morphology and stained positive for NANOG and TRA-1-60 (Fig. 7a). Addition of DOX caused rapid downregulation of *DNMT1** (Fig. 7b) and cells began to die within a few days (Fig. 7c and Supplementary Fig. 7a). During this window we observed increased DNA damage and arrest in G1 (Supplementary Fig. 7b, c and Supplementary Table 3), consistent with previous studies that reported DNMT1 depletion in human epidermal progenitors and primary human fibroblasts (IMR90) results in G1 arrest[20,46]. Using a catalytic inactive version of DNMT1 (C1226W)[47], which failed to provide a rescue, we confirmed that this effect is due to the catalytic activity of DNMT1 (Supplementary Fig. 7d–7g).

Despite G1 arrest and cell death, our inducible system enabled us to assess the effect on global DNA methylation after acute loss of *DNMT1* activity, which highlighted a striking decrease over a one-week period (Supplementary Fig. 7h). We also confirmed that the catalytic inactive DNMT1[C1226W] could not rescue the cells and prevent the global loss of DNA methylation (Supplementary Fig. 7i). We then generated RRBS libraries to perform higher resolution analysis (Fig. 7d). Consistent with global, replication-dependent depletion, we find that all genomic features are affected similarly (Fig. 7e). We fitted an exponential model of the methylation decay of individual 1kb tiles and determined that it took 2.31 days for global DNA methylation to be reduced to half of the original levels (Fig. 7f). As expected, non-CpG methylation, which is the exclusive purview of *de novo* methyltransferases, was not affected by the acute loss of *DNMT1* (Fig. 7g).

Discussion

DNA methylation is essential for normal development. Over the past three decades, major advances in understanding its role have included the deletion of the responsible enzymes in the mouse model as well as the more recent emergence of comprehensive mapping tools. Here, we have applied both to the human ESC model system to study their function, providing an unprecedented view of the respective genomic targets (shared and unique) of these enzymes.

All of the *DNMT3* knockouts, including the *DNMT3A^−/−^3B^−/−^* cells, were able to differentiate into the three germ layers. Overall CpG methylation was only mildly affected in the single knockouts and even the *DNMT3A^−/−^3B^−/−^* cells retained around 65% methylation after 17 passages (wildtype HUES64 83%). This observation is consistent with previous mouse studies that suggested a dependence on the DNA methylation levels rather than the enzymes in the teratoma assay[39]. Our continued passing suggests an average fidelity for DNMT1 of around 99.8%, which is higher than that estimated for mouse ES cells[2]. Further experimentation is required to explore the underlying mechanism, however one can hypothesize that this simply reflects cell-type specific effects that have been noted for DNA methylation turnover rates[48]. We have recently observed increased DNA methylation discordance in chronic lymphocytic leukemia (CLL), which may be related to increased proliferation[42]. In this context, it is worth noting that mouse and human ESCs proliferate at very different rates (13.5±2.5 and 36±12 hours, respectively)[49,50].

Our *DNMT3* knockout human ESC lines provide a powerful model to study cell state-driven epigenetic effects during the differentiation into specific cell types. It is likely that phenotypes arise only post-differentiation, when *de novo* activity is required and/or compensation through DNMT3B is no longer available. *Dnmt3a* knockout mice develop to term, but die within a few weeks after birth[4], suggesting some essential, though still poorly understood, developmental function.

It is well established that mouse and human ESC represent different pluripotent states, which may explain the sensitivity of the human ESCs to loss of DNA methylation. All somatic cells require DNA methylation and the more mature epiblast state modeled by current human ESCs may represent the first developmental period where maintenance becomes essential. In fact, although no data were shown, a similar effect has been suggested for mouse epiblast cells[51]. In the future, it will be interesting to test the effect of *DNMT1* deletion under recently described human "naïve" conditions to dissect the transition to DNMT1-dependent pluripotency[52,53,54].

Our study provides a comprehensive, high-resolution view of the largely conserved roles that the three catalytically active DNMTs play in human cells. The mutant lines represent powerful tools to study their function in various differentiation models and will therefore greatly accelerate our understanding of DNA methylation in human development and disease.

Online Methods

Cell culture

Human embryonic stem cells were expanded on murine embryonic fibroblasts CF1 (Global Stem) in KO-DMEM (Life Technologies) containing 20% Knockout Serum Replace (Life Technologies) and FGF2 (10 ng/ml) (Millipore). Cells were passaged by enzymatic dissociation using Collagenase IV (1 mg/ml) (Life Technologies). For feeder free culture condition, plates were coated with Geltrex (Life Technologies) and the culture medium was mTeSR1 (Stemcell Technologies). HUES64 was derived in the Melton/Eggan labs[56] and maintained in our laboratory since then using the above conditions.

Electroporation

Human ESCs were harvested using 1mg/mL Collagenase Type IV (invitrogen) first and then dissociated into single cells using TrypLE™ (invitrogen). 1×10^7 cells resuspended in buffered saline (PBS) were electroporated with 15μg Cas9 and 15μg gRNA plasmids (Gene Pulser Xcell System, Bio-Rad: 250 V, 500μF, 0.4 cm cuvettes). Cas9 plasmid and gRNA backbone have been described previously[35]. Fluorescence-activated cell sorting (FACS) was carried out 48 hours posttransfection based on fluorescent-marker expression. Sorted cells were replated as single cells at very low density on irradiated CF1 feeders in hESC medium supplemented with Rho Kinase (ROCK)-inhibitor (Stemgent, Y-27632) for the first 24 hours. Individual colonies were picked and expanded on feeder free/mTeSR condition. Genomic DNA was then purified. Mutations were validated by PCR and DNA sequencing.

Constructs

GIPZ *DNMT1* shRNA: Thermo Scientific, RHS4531-NM_001130823

TRIPZ DOX inducible *DNMT1* shRNA: Thermo Scientific, RHS4740-EG1786

plenti-ef1a-*tTA*-2a-*mCherry: tTA*-2a-*mCherry* was inserted into lenti backbone (Addgene,17448). *tTA* was cloned from Addgene plasmid 26803[57]. *mCherry* was cloned from Addgene plasmid 23243[58].

plenti-ef1a-*puro*^r-TRE-*DNMT1**: *DNMT1* was clone from Addgene plasmid 36939[59]. After creating mutations, *DNMT1** was cloned into plenti-ef1a-*puro*^r-TRE backbone[60] which was a gift from Dr Lei Xiao's lab.

plenti-ef1α-*DNMT1*^WT-IRES-nlsEGFP: *DNMT1* was clone from Addgene plasmid 36939[59].

plenti-ef1α-*DNMT1*^C1226W-IRES-nlsEGFP: *DNMT1* point mutation C1226W was created on the *DNMT1*^WT.

Primers

DNMT3A U:

Forward: GACTCCATCACGGTGGGCATGG;

Reverse: TGTCCCTCTTGTCACTAACGCC.

DNMT3A D:

> Forward: TCCACTGTGAATGATAAGCTG;

> Reverse: GGAAACCAAATACCCTTTCCA.

DNMT3B U:

> Forward: GAGTCCATTGCTGTTGGAACCG;

> Reverse: ATGTCCCTCTTGTCGCCAACCT.

DNMT3B D:

> Forward: GACTGCTTGGAATACAATAGGA;

> Reverse: AAAGCCAAAGATCCTTTCGAG.

DNMT1 P:

> Forward: GGTTTCCTTCCTCAGCTACTGCGA;

> Reverse: CACTGATAGCCCATGCGGACCA.

ACTIN:

> Forward: TTTGAGACCTTCAACACCCCAGCC;

> Reverse: AATGTCACGCACGATTTCCCGC.

Sequence information for TALENs targeting strategy

For *DNMT1* knockout:

> tgtgggccatcgagatgtGGGACCCTGCGGCCCAggcgttccggctgaacaa

For *DNMT3A* knockout:

> tcttctttgagttctaccGCCTCCTGCATGATGCgcggcccaaggagggaga

Capital letters are the spacer region. Lower case letters are the TALEN binding sites.

Virus production and infection on human ESCs

For virus production, virus was produced by co-transfecting lenti plasmid with delta 8.91 and VSVG on 293 cell by using FuGENE® HD (Promega). 24 hours after transfection, medium was changes to mTeSR1 for virus collection. Supernatant of virus was collected twice 48h and 72h after transfection separately. Virus was stored in 4°C. Before infection, virus was filtered through 0.45μm filter.

For virus infection on human ESCs, cells were plated as small clumps on geltrex pre-coated plate. 24 hour after plating, cells were infected with virus supernatant for 3 hours in incubator. Medium was changed back to mTeSR1. Repeat the 3 hours infection the next day. In this case, cells were maintained well in pluripotent state.

Isolation of genomic DNA from 96-well plate

Cells in each well were about 90% confluent. Washed twice with PBS and then lysed with 50μl/well of lysis buffer (10mM Tris PH7.5, 10mM EDTA, 10mM NaCl, 0.5% Sarcosyl or SDS, 1mg/ml Proteinase K) at 37°C overnight. Genomic DNA was precipitated by adding 100μl/well of cold NaCl/EtOH (1.5% vol 5M NaCl in 100% cold EtOH) at room temperature (RT) for 2–4 hrs. Genomic DNA precipitate was washed 3 times with 70% EtOH. Each time discarding the alcohol by inversion. Air-dry DNA for 15–20 min at RT. DNA was re-suspended in 30–100 μl of DNase-free, RNase-free ddH$_2$O.

Antibodies

For immunostaining:

OCT4: BD Transduction Laboratories, 611202

NANOG: R&D, AF1997

TRA-1-60: Stemgent 090068

ANNEXIN V: Abcam, ab54775

FOXA2: R&D, AF2400

HNF4A: Abcam, Ab41898

DAPI: Hoechst 33342, Invitrogen, H3570.

γ-H2A.X (phospho S139): Abcam, ab2893.

For western blotting:

DNMT3A: Cell signaling, 2160S

DNMT3B: santa cruz biotechnology, sc-20704

DNMT1: Activemotif, 39204

ACTIN: Abcam, ab8226

Immunostaining

Human ESCs were passaged on Geltrex coated plates and cell were cultured in mTeSR1 medium. Then the cells were fixed in 4% paraformaldehyde (PFA) for 15 minutes at room temperature. Permeabilization and blocking was performed in 5% bovine serum albumin (BSA) and 1% Triton X-100 in PBS for 30minutes. Cells were stain with primary antibody in 4°C overnight. The secondary antibody was applied for 2 hours at room temperature. Nucleuses were stained with Hoechst 33342 (Life Technologies). Images were obtained using Olympus IX71 microscope and MetaMorph Advanced software.

RNA extraction

RNA was isolated from hESCs using TRIzol (Invitrogen, 15596-026), further purification was carried out with RNeasy columns (QIAGEN, 74104) and DNase treatment.

Embryoid body (EB) formation

Human ESCs were expanded in $10cm^2$ dishes on irradiated CF-1 MEFs (Global Stem). Cells were harvested using 1mg/ml Collagenase IV (Life Technologies) once they were confluent. After three times washes with $1 \times$ phosphate-buffered saline (PBS), cell clumps were plated into 6-well low attachment dishes (Corning) in 20% Knockout Serum Replacement (KOSR) (Life Technologies), 200mM Glutamax (Life Technologies), MEM Non-Essental Amino Acid, in KO DMEM (Life Technologies). Medium was changed every 2–3 days.

Directed differentiation into the three germ layers

hESCs were plated as clumps on 6-well plates coated with Geltrex (Life Technologies) in mTeSR1 culture medium (Stem Cell Technologies). On day 3, directed differentiation was induced and medium was changed every day.

Ectoderm differentiation medium contains 2μM A83-01 TGFb inhibitor (Tocris), 2μM PNU-74654 WNT3A inhibitor (Tocris), and 2μM Dorsomorphin BMP inhibitor (Tocris) in 15% KOSR (Life Technologies), MEM Non-Essential Amino (Life Technologies), and 55μM 2-mercaptoethanol.

Mesoderm differentiation medium contains 100ng/ml Activin A (Life Technologies), 10 ng/ml bFGF (Millipore), 100ng/ml BMP4 (Life Technologies), 100ng/ml VEGF 100ng/ml (Life Technologies) in 0.5% FBS (Hyclone), MEM Non-Essential Amino Acid (Life Technologies), 55μM 2-mercaptoethanol and Glutamax (Life Technologies) in DMEM/F12 medium (Life Technologies) for the first 24 hrs of differentiation. For the rest 4 days, the medium contains 10 ng/ml bFGF (Millipore), 100ng/ml BMP4 (Life Technologies), 100ng/ml VEGF 100ng/ml (Life Technologies) in 0.5% FBS (Hyclone), MEM Non-Essential Amino Acid (Life Technologies), 55μM 2-mercaptoethanol and glutamax (Life Technologies) in DMEM/F12 medium (Life Technologies).

Endoderm differentiation medium contains 100ng/ml Activin A (Life Technologies) and 2μM/ml Lithium Chloride (Sigma) in 0.5% FBS (Hyclone), 200mM GlutaMax (Life Technologies), MEM Non-Essential Amino Acid (Life Technologies), 55μM 2-mercaptoethanol in RPMI medium (Life Technologies).

Teratoma formation

Teratoma formation assay was performed as a service by Harvard Genome Modification Facilities. Briefly, hESCs were injected into the kidney capsules of 3 immuno-suppressed mice as 1 million cells per animal to form teratomas. The formed teratomas were fixed in 4% paraformaldehyde (PFA) pH7.4, embedded in paraffin, sectioned 10–12μm thickness, and stained with Hematoxylin and Eosin (H&E) for examination. Histology examination was performed by histology core facility at Harvard Stem Cell Institute.

Hepatoblast Differentiation

Differentiation was performed as described previously[61] with some modifications. Briefly, cells were dissociated with Accutase and seeded onto geltrex-coated dishes with ROCK-inhibitor at high confluence in mTESR1. Within twenty-four hours after plating, definitive

endoderm differentiation was started and cells were cultured in RPMI/2% B27 (without Insulin), 100ng/ml Activin A, 10ng/ml BMP4, and 20ng/ml FGF2 for 48 hours. For days 3–5 of differentiation, cells were cultured in RPMI/2% B27 (without Insulin), 100ng/ml Activin A. For days 6–10 of differentiation, cells were cultured in RPMI/2% B27 (with Insulin), plus 20ng/ml BMP4 and 10ng/ml FGF2. Media was changed every twenty-four hours.

Cell cycle assay

Alive cells were incubated with Hoechst 33342 in the incubator for 15min. Then cells were trypsinized into single cells, washed with PBS once and analyzed by FACs directly.

Dot blot assay

Isolated genomic DNA was denatured in 0.4M NaCl and 10mM EDTA for 10 min at 99°C. The denatured DNA was spotted in a 2-fold serial dilution on a Zeta Membrane (BioRad) using a Bio-Dot apparatus (BioRad). The blotted membrane was rinsed in 2xSSC, air-dried and UV-crosslinked at 120,000 μJ cm^{-2}. The membrane was blocked in 1xTBS containing 5% nonfat dry milk for 30min. Anti-5-methylcytosine monoclonal antibody (Diagenode, #33D3) was diluted 1:250 in TTBS, containing 0.3% Tween 20 and 5% nonfat dry milk, and was incubated overnight at 4°C. The membrane was washed in 3x 5min in TBS and was incubated for 1 hour at room temperature with HRP-conjugated goat anti-mouse IgG (Jackson Lab, #115-025-174). The membrane was washed 3x in TBS and visualized by chemiluminescence using the Gel Doc XR (BioRad). The same membrane was incubated 30min with SYBR Gold Nucleic Acid stain (Invitrogen, #S-11494) in 1xTAE buffer, total DNA loading was visualized by UV transillumination.

Scorecard Assay and Data Analysis

Experiment was carried out using TaqMan® hPSC Scorecard™Kit (Life Technologies, A15872). Data could be analyzed through TaqMan® hPSC Scorecard™ Analysis Software through Life Technologies website.

RRBS and WGBS Library Construction

RRBS library construction was described in Ref [62]. WGBS library construction was well described in Ref [9].

RRBS and WGBS Data Processing and Analysis

WGBS raw sequencing reads were aligned using MAQ in bisulfite mode against human genome version hg19/GRCh37, discarding duplicate reads. DNA methylation calling was performed based on an extended custom software pipeline published previously for RRBS[63]. Briefly, the methylation level at a CpG or CpA site was the number of reads with that site methylated divided by the total number of reads covering the site.

To ensure comparability of region DNA methylation levels across all samples, only CpGs covered by ≥5x in all samples qualified for the computation of region DNA methylation levels. We defined the average methylation for a genomic region as the coverage-weighted mean of the methylation levels of the individual CpGs within the region. Subsequently, we

averaged a region's DNA methylation level over replicates. Differentially methylated regions were identified by using a two sample weighted t-test using the methylation values of the CpGs within each regions as individual sampling events (minimum number of CpGs in region for each sample is 2). We performed multiple testing correction using a method defined in Ref [64] using the R qvalue package.

We used two sets of CpG island annotations. The first uses a bioinformatics definition and designates a region as a CpG island if it has a GC content higher than 0.5 and a ratio greater than 0.6 of observed CpG dinucleotides to what would be expected based on the GC content over a length of 700 bp. The second is a set of experimentally verified CpG islands[41].

We defined promoters as the regions 2000 bp upstream and 500 bp downstream of the Refseq TSS set. These were divided into high CpG promoters (HCP), intermediate CpG promoters (ICP), and low CpG promoters (LCP) as published previously[65]. Intergenic regions, exons and introns were defined on the basis of the RefSeq gene models downloaded from the UCSC browser.

We used GREAT (Genomic Region Enrichment of Annotation Tool)[45] to associate DMRs with genes. We limited the associations to genes within 20kb of DMRs and used an FDR threshold of 10^{-6} and a fold enrichment threshold of 2. The complete set of enriched terms across all ontologies tested is in Supplementary Table 2.

Fidelity of DNMT1 was calculated by fitting an exponential model to the mean DNA methylation of different genomic features using the nls function in R. The exponential model has the form $y=a^x$, where "a" is the fidelity and "x" is the estimated number of population doublings based on passage number, days per passage (8) and doubling time (28.8 hours).

The proportion of discordant reads (PDR) was calculated for each CpG by examining the individual reads covering that CpG and classifying them into concordant (all CpGs methylated or all CpGs unmethylated), filtering out reads with less than 3 CpGs covered.

We downloaded the conservative set of uniform peaks called by the ENCODE project[43] and computed overlaps of these peaks with DMRs identified in $DNMT3A^{-/-}$, $DNMT3B^{-/-}$ and $DNMT3A^{-/-}3B^{-/-}$. We calculated the significance of the enrichment using Fisher's exact test, using the DMR set as the foreground and the set of non-repetitive 1kb tiles with a methylation level of at least 0.4 as the background.

Human imprinted control regions were taken from Ref. [66].

Mouse $Dnmt3a^{-/-}$, $3b^{-/-}$ and $3a^{-/-}3b^{-/-}$, $Dnmt1{-}/{-}$ and $3a^{-/-}3b^{-/-}$ with a $Dnmt1$ knockdown[16] data were aligned to mm9 and otherwise processed similar to the human RRBS data.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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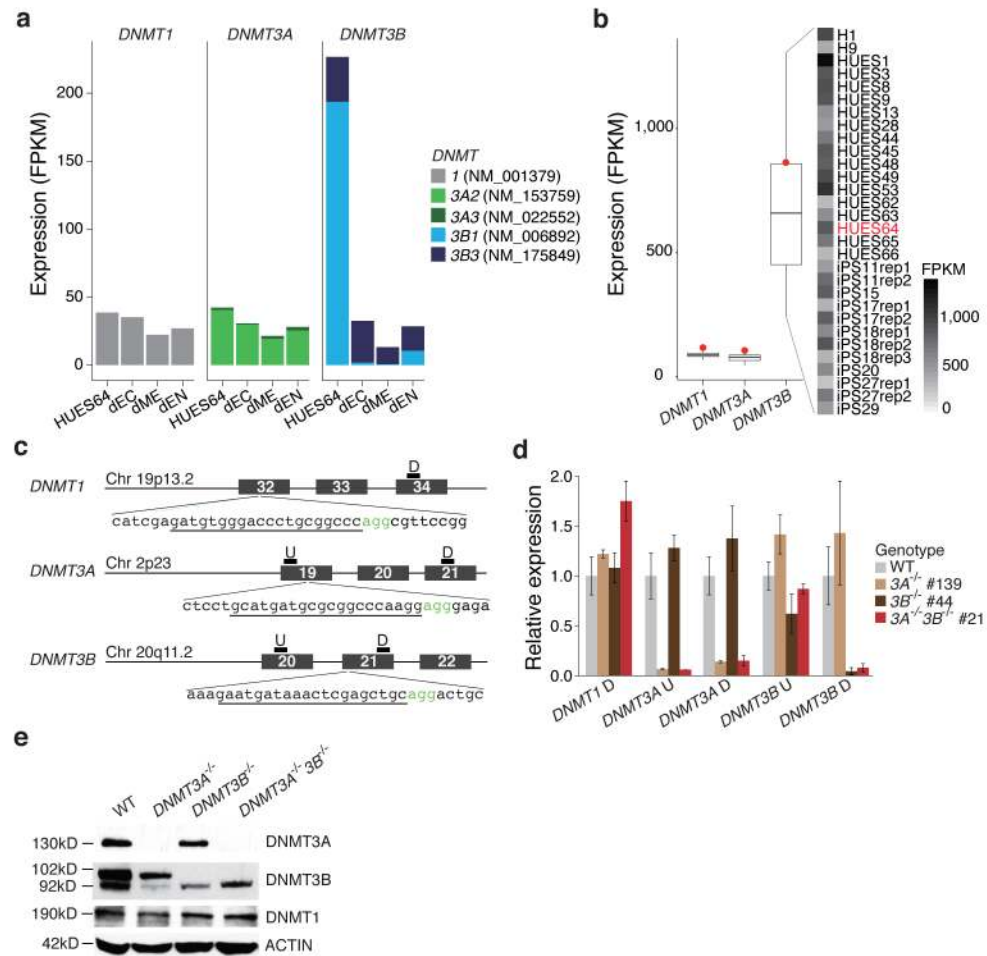
Figure 1. Targeted deletion of *DNMT1*, *3A* and *3B* in human ESCs

(a) Expression levels of *DNMT1*, *3A* and *3B* in undifferentiated HUES64 ESCs and their derivatives, ectoderm (dEC), mesoderm (dME) and endoderm (dEN) in FPKM (Fragments Per Kilobase per Million fragments mapped) are shown. Only expression of the major isoforms is shown (see Supplementary Fig. 1a for all). (b) Left: Expression of *DNMT1*, *3A*, *3B* for 25 pluripotent (ESC and iPSC) lines (line: median, box: IQR, whiskers: furthest point within 1.5xIQR, red dot: HUES64). Right: Cell lines and *DNMT3B* expression. There is substantial variation of *DNMT3B* expression even among biological replicates ("rep").(c) Overview schematic of the Cas9/gRNA-target sites. Genomic coordinates are shown on the right. The gRNA-targeting sequence is underlined, and the Protospacer-Adjacent Motif (PAM) sequence is labeled in green. Position of qPCR primers for RNA expression validation is shown on the top of the exons. P: primer pair; U: upstream; D: downstream.(d) RT-qPCR analysis for *DNMT1*, *3A* and *3B* in HUES64, *DNMT3A*$^{-/-}$ (#139), *DNMT3B*$^{-/-}$ (#44) and *DNMT3A*$^{-/-}$*3B*$^{-/-}$ (#21). Primer details are in Fig. 1c. Error bars were generated from biological triplicates and represent one standard error.

(e) Western blot analysis for DNMT3A, 3B and DNMT1 in HUES64, *DNMT3A*$^{-/-}$ (#139), *DNMT3B*$^{-/-}$ (#44) and *DNMT3A*$^{-/-}$*3B*$^{-/-}$ (#21). Arrow indicates the major DNMT3B isoform (NM_006892). The smaller band in the DNMT3B western blotting result is likely
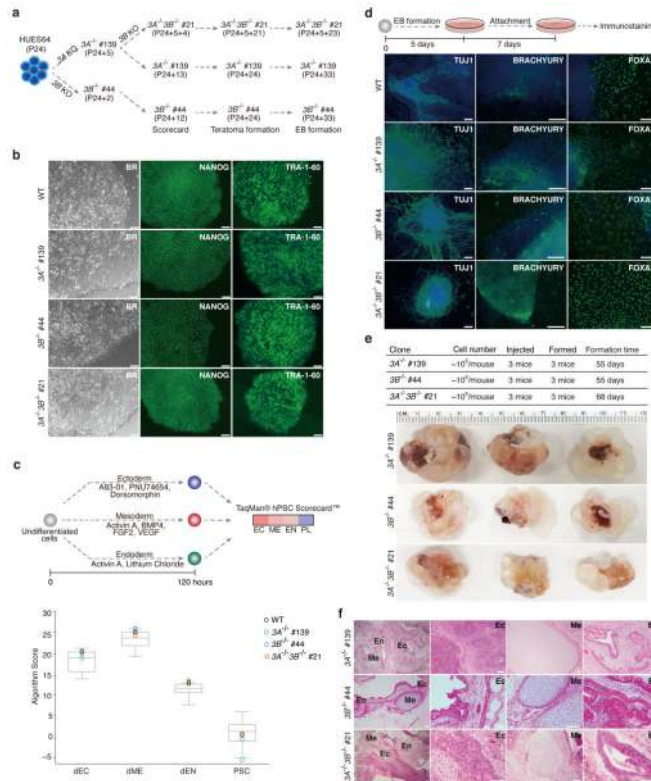
isoform 3 (NM_175849) as reported in Ref. [55], which was not targeted by our knockout strategy.

Figure 2. Assessing the differentiation potential of the *DNMT3 knockouts*

(a) Overview of clones and passage numbers used for the various assays. HUES64 passage 24 (P24) was used to generate *DNMT3A* and *3B* single knockouts. The number after "+" represents the passage number post targeting. The *DNMT3A$^{-/-}$* clone #139 was passaged five times (P24+5) and then targeted using the *DNMT3B* gRNA. The *DNMT3A$^{-/-}$3B$^{-/-}$* passage numbers are then counted from there.

(b) Wild-type and *DNMT3A$^{-/-}$*, *3B$^{-/-}$* and *3A$^{-/-}$3B$^{-/-}$* cells show comparable morphology and expression of characteristic markers. Representative bright field (BR) and pluripotency-associated marker (NANOG, TRA-1-60) immunostaining images are shown. 10X magnification is shown. Scale bars, 100μm.

(c) TaqMan hPSC Scorecard analysis of direct differentiation potential for WT, *DNMT3A$^{-/-}$*, *3B$^{-/-}$* and *3A$^{-/-}$3B$^{-/-}$* cells. Top: Schematic overview of the conditions used for direct differentiation into ectoderm, mesoderm and endoderm. Bottom: Differentiation scores for WT, *DNMT3A$^{-/-}$*, *3B$^{-/-}$* and *3A$^{-/-}$3B$^{-/-}$* cells. Gray box represents the score range of 11 reference human PSC lines.

(d) Top: Schematic overview for the embryonic body (EB) formation. Bottom: Representative immunostaining images for TUJ1 (ectoderm), BRACHYURY (mesoderm) and FOXA2 (endoderm). 10X magnification is shown. Scale bars, 100μm.

(e) Top: Summary of injection and teratoma formation results. Bottom: Images of the nine teratomas collected. Ruler on top provides a size reference.

(f) Representative images of Hematoxylin and Eosin (H&E) staining of the teratoma from **panel e**. Neuroepithelium (ectoderm) (EC), cartilage with ossification center (mesoderm) (ME), and gut-like epithelium (EN). Scale bars, 50μm.
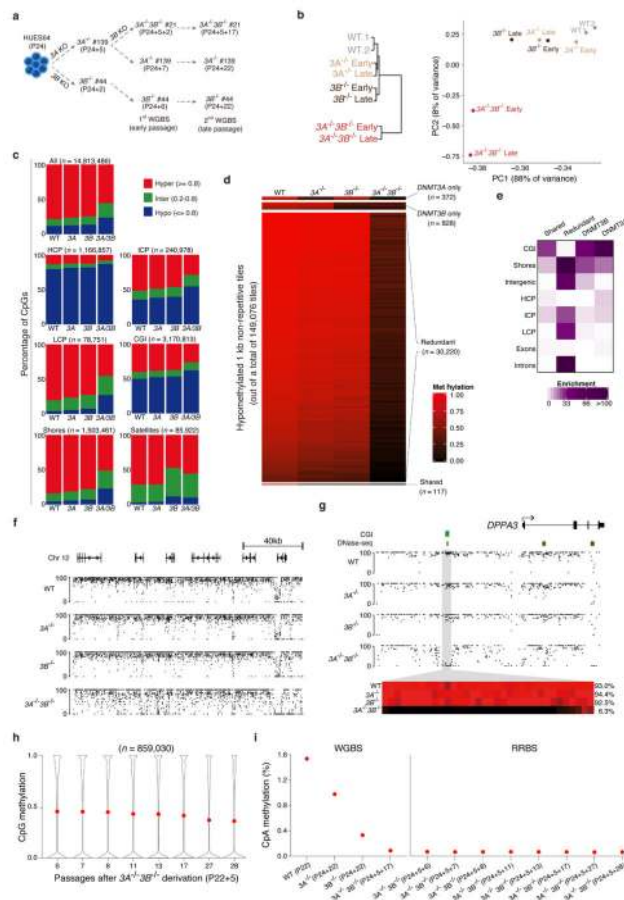
Figure 3. Global DNA methylation dynamics

(a) Overview of the samples used for whole genome bisulfite sequencing (WGBS).

(b) Left: Hierarchical clustering using Euclidean distance based on mean DNA methylation levels of 1 kb tiles across the human genome. Right: Principal component analysis (PCA) based on mean CpG methylation levels for 1 kb tiles.

(c) Fraction of CpGs with high (≥0.8, red), intermediate (inter, >0.2 and <0.8, green) and low (≤0.2, blue) methylation values. Top left panel shows all CpGs while the other panels show CpG methylation distribution within different genomic features, including high CpG Promoters (HCP), intermediate CpG promoters (ICP), low CpG promoters (LCP), CpG islands (CGI), CpG island shores, and satellite repeats. The total number of CpGs associated with each feature is shown above each bar plot.

(d) Heatmap of non-repetitive differentially methylated 1kb tiles (q-value < 0.05 and methylation difference greater than 0.4).

(e) Enrichment [−log10(hypergeometric p-value)] of genomic features in the target classes identified in Fig. 3d.

(f) Genome browser tracks covering approximately 160kb in the different lines.

(g) Higher resolution view of the *DPPA3* (*STELLA*) locus. The heat map below shows the DNA methylation values of individual CpGs within the grey region. The average DNA methylation value for the entire highlighted region is shown on the right.

(h) CpG methylation levels over several passages of the *DNMT3A*$^{-/-}$*3B*$^{-/-}$ cells. Red dot indicates mean methylation.

(i) Global mean CpA methylation levels as assayed by WGBS and over several passages of the *DNMT3A*$^{-/-}$*3B*$^{-/-}$ cells by RRBS.
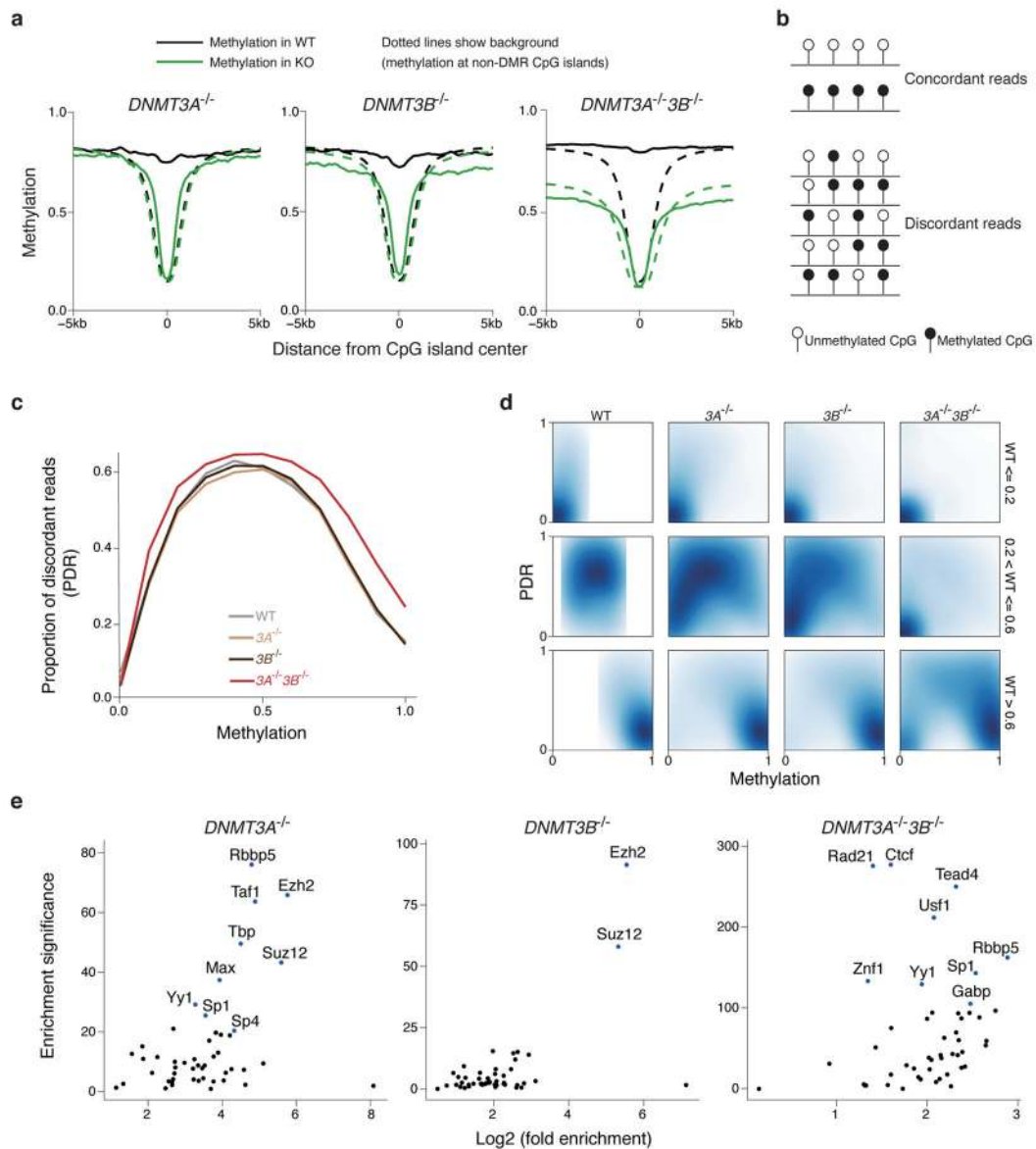
Figure 4. Characterization of targets for DNMT3A and DNMT3B

(a) Composite plot of methylation around CpG islands in WT and knockouts. The solid lines show the mean methylation of differentially methylated CpG islands, while the dotted lines show the mean methylation of all other CpG islands.

(b) Schematic for the definition of concordant and discordant reads.

(c) Proportion of discordant reads (PDR) in WT, *DNMT3A$^{-/-}$*, *3B$^{-/-}$* and *3A$^{-/-}$3B$^{-/-}$* cells.

(d) Two-dimensional density plots of PDR vs. methylation for CpGs with low (<= 0.2), medium(> 0.2 and <= 0.6) and high(> 0.6) methylation in WT cells.

(e) Enrichment of experimentally determined transcription factor binding sites in H1 ESCs generated by the ENCODE project for DMRs in *DNMT3A$^{-/-}$*, *3B$^{-/-}$* and *3A$^{-/-}$3B$^{-/-}$* cells compared to WT. Enrichment significance was defined as −log10(p-value) of the Fisher's exact test, with the background being non-repetitive 1kb tiles that had at least 0.4 enrichment in WT.
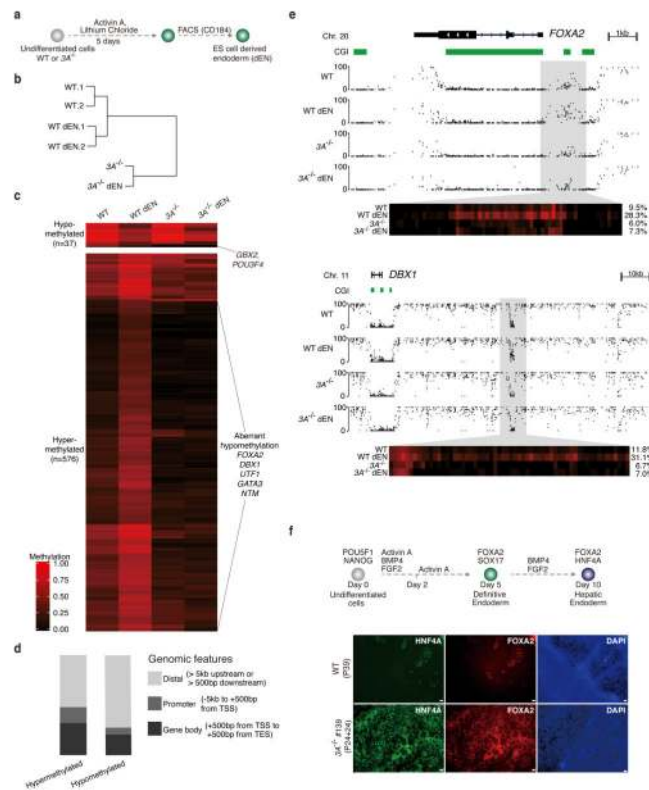
Figure 5. Effect of *DNMT3A* deletion on endoderm differentiation

(a) Schematic for the generation of CD184+ endodermal progenitor cells (dEN).

(b) Global analysis for the WGBS data of undifferentiated and dEN cells from WT (P23 and P26) and *DNMT3A*$^{-/-}$ (P24+13). Hierarchical clustering based on mean DNA methylation levels of 1 kb tiles across the human genome using Euclidean distance. 1 and 2 are two biological replicates. The passage number of the replicates is within 2 passages.

(c) Heat map of methylation levels (black, 0; red, 1) in WT, WT dEN, *DNMT3A*$^{-/-}$, and *DNMT3A*$^{-/-}$ dEN of DMRs that change methylation significantly (q-value < 0.05 and methylation difference of at least 0.2) between WT and WT dEN.

(d) Classification of endoderm DMRs into promoters, gene bodies and distal regions as defined in the legend.

(e) Genome browser tracks of 1kb and 10kb regions highlight global and local methylation level for *FOXA2* and *DBX1*. The heat map below shows the DNA methylation values of individual CpGs within the highlighted region. The average DNA methylation value for the entire highlighted region is shown on the right.

(f) Hepatoblast differentiation of WT (P39) and *DNMT3A*$^{-/-}$ (P24+24) cells. Top: Schematic of hepatoblast differentiation. Bottom: Representative images from two individual biological replicates are shown. 4X magnification of hepatoblast markers FOXA2, HNF4A immunostaining images for HUES64 (WT), *DNMT3A*$^{-/-}$ (#139) are shown. Scale bars, 100μm.
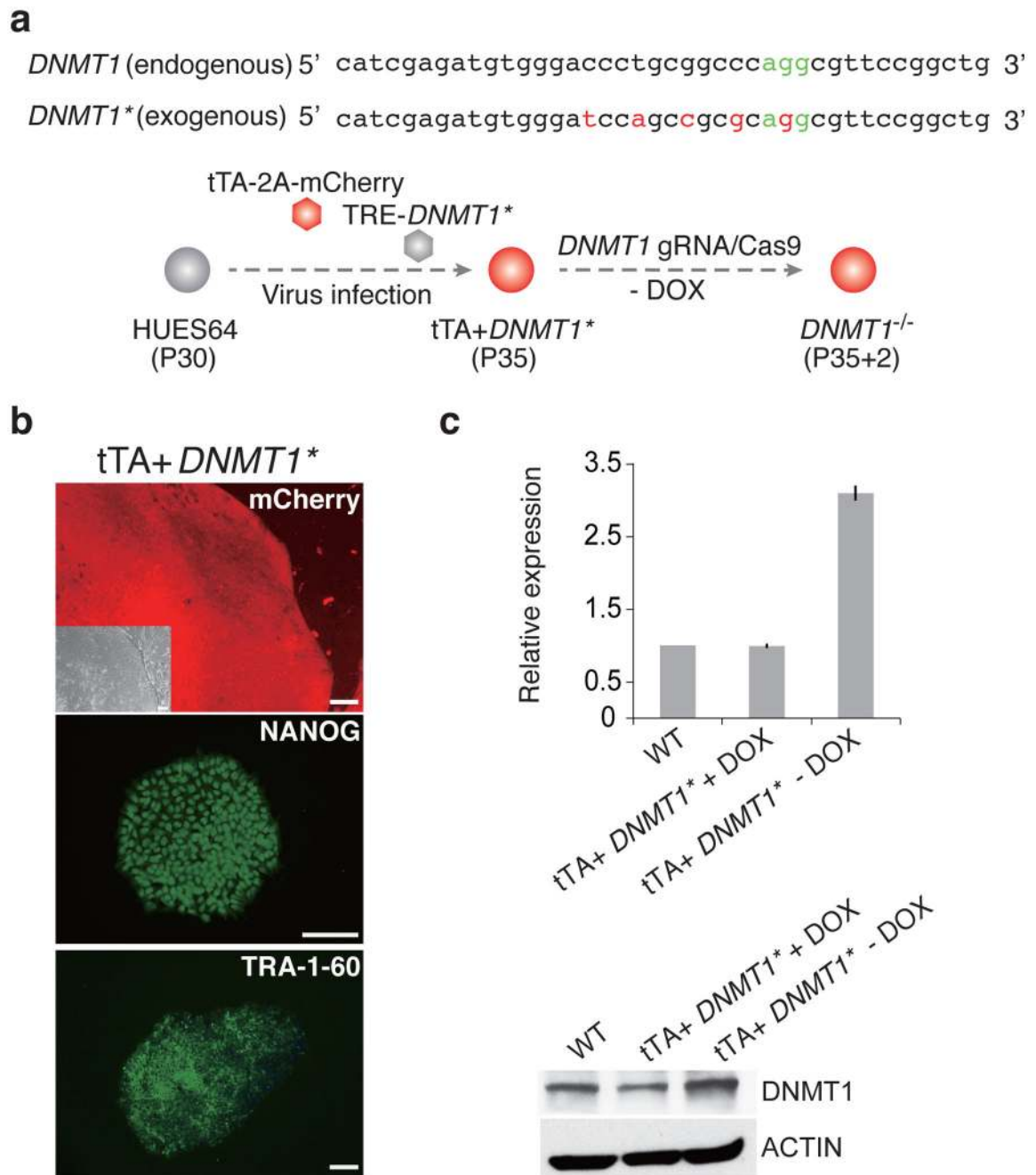
Figure 6. *DNMT1* knockout strategy and targeting efficiency

(a) Strategy to obtain *DNMT1* homozygous clones. Top: We created mutations within the exogenous *DNMT1* sequence that would prevent the targeting by our gRNA, but not alter the protein (Supplementary Fig. 6d). Asterisk (*) is used to distinguish the exogenous modified *DNMT1** from endogenous *DNMT1*. Bottom: The *DNMT1* rescue line was established by infecting HUES64 with two separate lenti virus, tTA-2a-mCherry and TRE-*DNMT1** at passage 30. Then, *DNMT1* was targeted as before into the tTA-*DNMT1** line while the exogenous *DNMT1** was expressed (no DOX).

(b) Representative bright field (BR), mCherry expression and pluripotent markers NANOG, TRA-1-60 immunostaining images for rescue line tTA+*DNMT1**. Scale bars, 100μm.

(c) *DNMT1* expression levels in our *DNMT1** TET/OFF system. Left: RT-qPCR analysis for *DNMT1*. Primers amplify both endogenous and exogenous *DNMT1*. Error bars were generated from biological replicates and represent one standard error. Right: Western blotting analysis for DNMT1 under the control of DOX. The antibody can detect both endogenous and exogenous DNMT1.
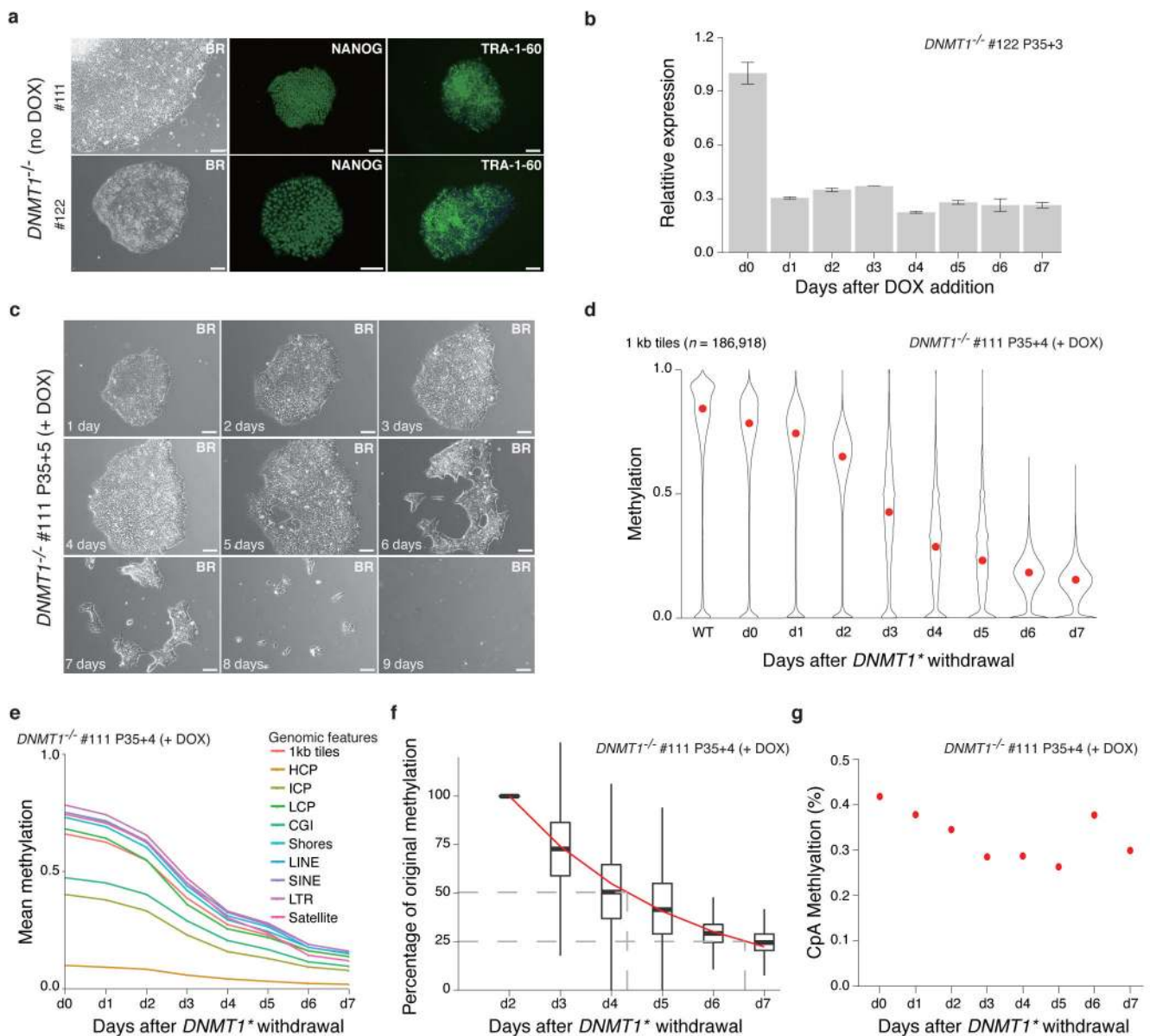
Figure 7. Loss of *DNMT1* causes global demethylation and cell death

(a) Representative bright field (BR) and NANOG/TRA-1-60 immunostaining for *DNMT1*−/− cells with the exogenous *DNMT1** expressed (no DOX). 10X magnification. Scale bars, 100μm.

(b) RT-qPCR analysis after *DNMT1** withdrawal relative to d0 level. Error bars are based on two replicates and represent one standard error.

(c) *DNMT1** withdrawal causes massive cell death. Representative 10X magnification bright field (BR) images are shown. Scale bars: 100μm.

(d) Violin plots of the mean methylation (measured by RRBS) of 1 kb tiles across the human genome after *DNMT1** withdrawal. Mean is indicated by red dot.

(e) Global reduction of DNA methylation in different genomic features. Overall CpGs (1kb tile) and selected features including CpG Islands (CGI), Shores, HCPs, ICPs, LCPs, LINEs, SINEs and satellite repeats are shown.

(f) Exponential model of methylation decay for 1kb tiles. Methylation at day 2 was normalized to 1 and methylation levels on subsequent days were calculated as a percentage of the day 2 value. An exponential model was fitted starting at day 2 and is shown by the red line. The dashed grey lines mark the timepoints when 50% and 25% of the day 2 methylation levels remain. Days 0 and 1 were excluded from this model to avoid noise from any remaining DNMT1 protein.

(g) Mean CpA methylation levels after *DNMT1** withdrawal.