

ARTICLE

Received 25 Oct 2011 | Accepted 18 May 2012 | Published 19 Jun 2012

DOI: 10.1038/ncomms1915

Histone modifications and lamin A regulate chromatin protein dynamics in early embryonic stem cell differentiation

Shai Melcer¹, Hadas Hezroni¹, Eyal Rand², Malka Nissim-Rafinia¹, Arthur Skoultschi³, Colin L. Stewart^{4,5}, Michael Bustin² & Eran Meshorer¹

Embryonic stem cells are characterized by unique epigenetic features including decondensed chromatin and hyperdynamic association of chromatin proteins with chromatin. Here we investigate the potential mechanisms that regulate chromatin plasticity in embryonic stem cells. Using epigenetic drugs and mutant embryonic stem cells lacking various chromatin proteins, we find that histone acetylation, G9a-mediated histone H3 lysine 9 (H3K9) methylation and lamin A expression, all affect chromatin protein dynamics. Histone acetylation controls, almost exclusively, euchromatin protein dynamics; lamin A expression regulates heterochromatin protein dynamics, and G9a regulates both euchromatin and heterochromatin protein dynamics. In contrast, we find that DNA methylation and nucleosome repeat length have little or no effect on chromatin-binding protein dynamics in embryonic stem cells. Altered chromatin dynamics associates with perturbed embryonic stem cell differentiation. Together, these data provide mechanistic insights into the epigenetic pathways that are responsible for chromatin plasticity in embryonic stem cells, and indicate that the genome's epigenetic state modulates chromatin plasticity and differentiation potential of embryonic stem cells.

¹ Department of Genetics, The Alexander Silberman Institute of Life Sciences, The Hebrew University of Jerusalem, Edmond J. Safra campus 91904, Israel. ² Laboratory of Metabolism, National Cancer Institute, National Institutes of Health, 37 Convent Drive, Bldg. 37, Bethesda, Maryland 20892, USA. ³ Department of Cell Biology, Albert Einstein College of Medicine, Bronx, New York 10461, USA. ⁴ Institute of Medical Biology, 8A Biomedical Drive Singapore 138648. ⁵ Department of Biological Sciences National University of Singapore. Correspondence and requests for materials should be addressed to E.M. (email: meshorer@huji.ac.il).

Chromatin and nuclear architecture in embryonic stem cells (ESCs) have several unique properties when compared with somatic or differentiated cells. Undifferentiated ESCs possess fewer condensed heterochromatin foci, which are larger and more dispersed than those of early differentiating cells¹ and a less structured nuclear lamina², which is devoid of lamin A³ (LMNA). LMNA is directly associated with chromatin^{4,5} and may thus confer nuclear rigidity and chromatin stability in differentiated cells. In undifferentiated ESCs, the organization of chromatin seems to be more homogeneous, and the DNA is distributed more evenly throughout the nucleus as compared with somatic cells or neuronal progenitor cells (NPCs) derived from ESCs by *in vitro* differentiation⁶. Moreover, chromatin in ESCs is characterized by hyperdynamic binding of architectural chromatin proteins including heterochromatin protein 1 (HP1), histone H1 and core histones⁷. Significantly, restriction of this dynamic state, using a tightly bound H1 mutant (H1cc), which we previously generated, interferes with ESC self-renewal and differentiation potential⁷, suggesting that the hyperdynamic state of chromatin proteins is functionally important for the stem cell state. ESCs are also distinguished by a unique epigenetic landscape. Developmentally regulated genes are enriched with histone marks of both active (H3K4me) and inactive (H3K27me) chromatin^{8–10}, rendering them primed for the onset of a transcriptional program that will lead to differentiation^{11,12}. In addition, in undifferentiated ESCs, the levels of acetylated histones are generally increased, while the levels of heterochromatin-associated histone modifications, such as histone H3 tri-methylated on lysine 9 (H3K9me3), are generally reduced^{2,13}. Finally, silent chromatin marked by H3K9me2 accumulates in the form of large blocks during ESC differentiation¹⁴. Taken together, these data underline the importance of chromatin and its epigenetic landscape for the maintenance of both cellular pluripotency and the differentiated state, and call for studying the mechanisms regulating chromatin plasticity in ESCs.

To investigate the molecular mechanisms underlying the hyperdynamic nature of chromatin in ESCs, in this study we focussed on the dynamic mobility of histone H1 (H1-GFP fusion protein), which, as it has an intermediate level of dynamics between the highly dynamic HP1 proteins and the stably bound core histones^{7,15}, is an excellent indicator of chromatin plasticity in ESCs⁷. We analysed H1 dynamics and differentiation potential in several mutant ESCs that lack various chromatin-related proteins, including *Suv39h1/2*^{-/-}, *G9a*^{-/-}, *Lmna*^{-/-}, *Dnmt1*^{-/-}, *Dnmt3a/3b*^{-/-}, *Dnmt1/3a/3b*^{-/-} and *H1c/d/e*^{-/-}, as well as following treatments with DNA de-methylating agents, histone acetyltransferase inhibitor and histone deacetylase inhibitors (HDACi). We found that the epigenetic state of the genome, particularly histone acetylation and histone methylation, as well as LMNA expression, affect chromatin plasticity and the differentiation potential of ESCs. Specifically, increased histone acetylation is associated with increased plasticity and protein dynamics on euchromatin, while H3K9 methylation regulates both euchromatin and heterochromatin protein dynamics and ectopic expression of LMNA exclusively restricts heterochromatin protein dynamics. In contrast, DNA methylation and nucleosome repeat length have little effect on chromatin protein dynamics. These results provide mechanistic insights into the regulation of chromatin plasticity in ESCs and pave the way for the manipulation of pluripotency by chromatin-related mechanisms.

Results

Histone acetylation increases H1 dynamics in euchromatin.

Previously, we reported that both linker and core histones display hyperdynamic association with chromatin in undifferentiated ESCs⁷. To investigate the potential mechanisms leading to this property, we tested different cells and treatments using H1-GFP fusion protein as an indicator for chromatin plasticity. Undifferentiated ESCs exhibit elevated levels of acetylated histones compared with differentiated

cells^{6,7,16}, hence we investigated whether these higher acetylation levels contribute to the hyperdynamic nature of chromatin-associated proteins in ESCs⁷. To this end, we examined the effects of HDACi on chromatin protein dynamics in somatic mouse embryonic fibroblasts (MEFs) and in mouse ESCs. We treated H1-GFP-expressing cells with low levels (0.5 mM) of the HDACi valproic acid (VPA) for 24 h before subjecting them to fluorescence recovery after Photobleaching (FRAP)⁷, analysing euchromatin and heterochromatin separately¹⁵. We detected a significant increase in the dynamic association of H1-GFP with euchromatin in MEFs (Fig. 1a), but not with heterochromatin (Fig. 1b). Similar behaviour was observed in undifferentiated R1 ESCs, with a significant increase in euchromatin (Fig. 1c) but not in heterochromatin (Fig. 1d), although initial levels of H1 kinetics were higher in ESCs and the relative increase was lower (compare Fig. 1a with Fig. 1c). Similar results were observed for a 4-h treatment with either VPA or the HDACi trichostatin A (TSA) (Fig. 1e–l). Increased acetylation levels were verified by western blotting (Fig. 2a). To test whether changes in H1 dynamics mark a general chromatin hyperdynamics pattern, we examined chromatin association dynamics of HP1 and of HMGN1 in ESCs—the dynamic binding of these two proteins to chromatin has been studied in detail^{7,17}; once again, similar results were observed (Supplementary Fig. S1), ruling out the possibility that the effect is due to acetylation or other modifications of H1 itself. Increased dynamics was more significant in euchromatin than heterochromatin (Fig. 1), suggesting differences in the accessibility to HDACs or in the nucleosomes' basic interaction with chromatin proteins. Regardless, increased histone acetylation correlates with increased chromatin protein dynamics in euchromatin in both somatic cells and ESCs.

Next, we examined how HDACi affect the differentiation propensity of ESCs. Interestingly, E14 ESCs treated with VPA for 24 h demonstrate a reduced propensity to differentiate in the absence of leukaemia inhibitory factor (LIF). When induced to differentiate by LIF withdrawal, the number of the neuroectodermal marker nestin-positive E14 ESC colonies increased from ~5 to ~25%, but when treated with VPA, 90% of the colonies remained nestin-negative (Fig. 2b). To confirm these results at the population level and with a different differentiation method, we performed qRT-PCR on RNA extracted from undifferentiated ESCs ± retinoic acid (RA) and ± VPA for 48 h and measured messenger RNA levels of nestin. RA alone increased *nestin* levels by almost twofold, but this increase was prevented by VPA (Fig. 2c). VPA alone did not significantly change *nestin* mRNA levels (Fig. 2c). We also analysed the expression levels of additional mesodermal (*Brachyury*) and endodermal (*GATA4*) differentiation markers. RA significantly reduced *Brachyury* and significantly increased *GATA4*, but these trends were reversed in the presence of VPA (Supplementary Fig. S2). We also analysed levels of pluripotency markers. After 48 h of LIF withdrawal, *Oct4* intensity in ESCs was reduced to 35% of control levels (Fig. 2d). However, in the presence of VPA, *Oct4* levels were restored to 76% of original levels (Fig. 2d). This was confirmed independently with RA for 48 h, in the absence or presence of VPA. Once again, low levels of VPA reduced RA-induced differentiation (Fig. 2e). These results suggest that HDACi-induced histone hyperacetylation in ESCs supports the undifferentiated state. However, HDACi do not affect the expression levels of pluripotency genes in ESCs^{18,19}, suggesting an indirect effect, possibly through altering global chromatin structure to a more open state²⁰. Together, these data show that elevated histone acetylation may be an underlying mechanism regulating chromatin hyperdynamics in undifferentiated ESCs and suggest that persistent hyperdynamics may have an inhibitory effect on differentiation, implying a potential functional link between chromatin dynamics and differentiation capacity.

We next examined the effect of HDACi on ESC-derived NPCs, which have lower chromatin protein dynamics than undifferentiated

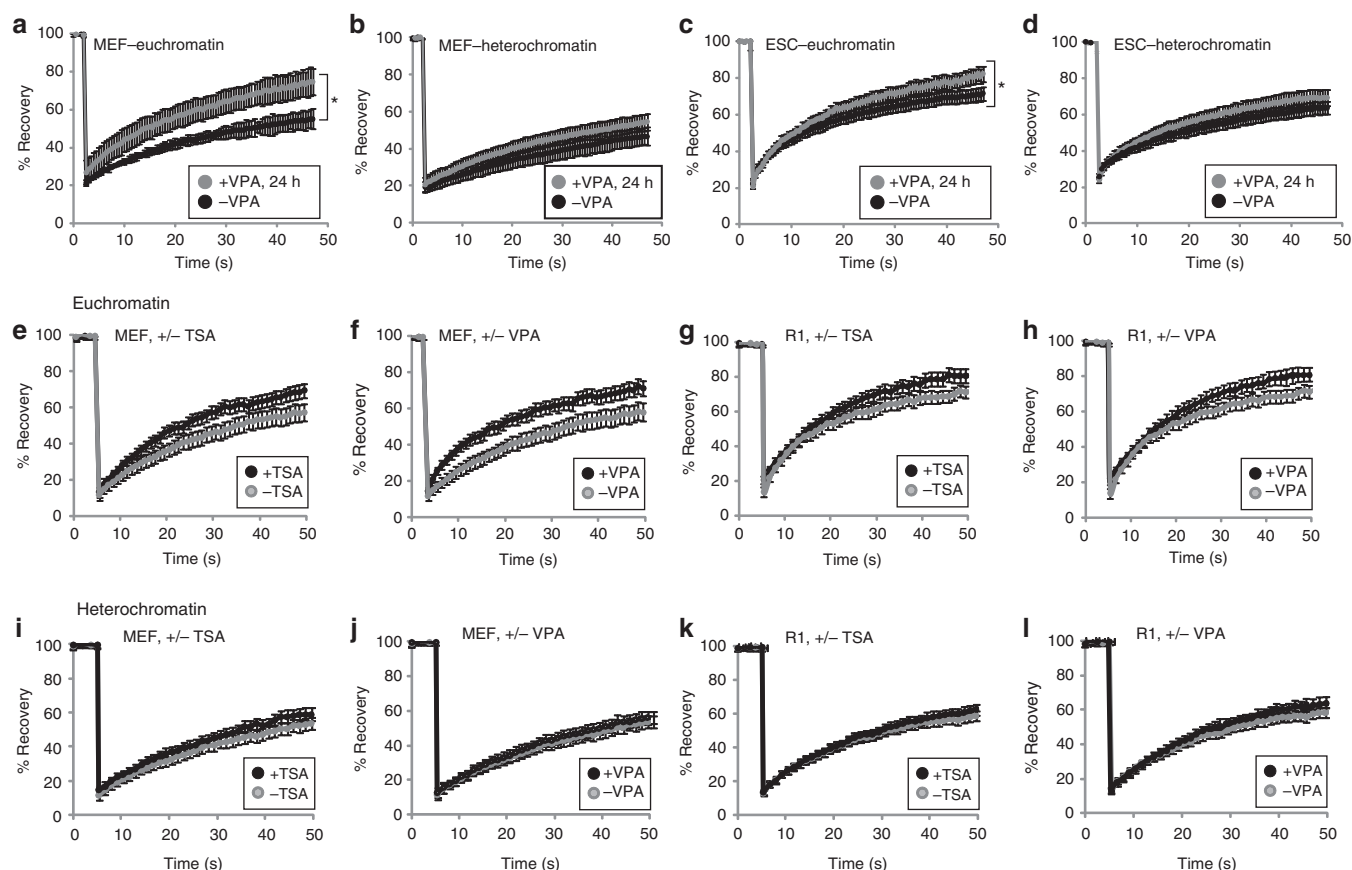


Figure 1 | Increased histone acetylation increases H1 dynamics. (a–d) FRAP analysis (\pm s.d.) of H1e-GFP in euchromatin (a,c) and heterochromatin (b,d) in MEFs (a,b) and in R1 mouse ESCs (c,d) treated (grey) or untreated (black) with the HDAC inhibitor VPA for 24 h. Changes for euchromatin are significant (a, $P < 0.005$; c, $P < 0.01$; d, $P > 0.05$, 2-tailed t -test). (e, f) FRAP analysis of H1e-GFP in euchromatin of MEFs treated (black) or untreated (grey) with TSA (e) or VPA (f) for 4 h. Changes are significant ($P < 0.005$ in both cases, 2-tailed t -test). (g, h) FRAP analysis of H1e-GFP in euchromatin of mouse ESCs treated (black) or untreated (grey) with TSA (g) or VPA (h) for 4 h. Changes are significant ($P < 0.05$ in both cases, 2-tailed t -test). (i, j) FRAP analysis of H1e-GFP in heterochromatin of MEFs treated (black) or untreated (grey) with TSA (i) or VPA (j) for 4 h. (k, l) FRAP analysis of H1e-GFP in heterochromatin of ESCs treated (black) or untreated (grey) with TSA (k) or VPA (l) for 4 h. Values are from at least 2 independent experiments, at least 20 cells each. Error bars represent s.d.

ated ESCs⁷. Indeed, increased H1–GFP dynamics was observed in NPCs treated with VPA for 24 h (Fig. 2f,g). Interestingly, whereas HDACi largely maintain ESCs in an undifferentiated state (Fig. 2b–e; Supplementary Fig. S2), VPA-treatment of NPCs slightly increased neuronal differentiation rather than halting it, as judged by the percentage of β -Tubulin-III (TUB1)-positive cells after 24 h (from $73.5 \pm 8.5\%$ in the untreated NPCs to $84.4 \pm 11.0\%$ in the VPA-treated NPCs; Fig. 2h). This result is consistent with a previous report, which demonstrated that, in hippocampal neural progenitor cells, VPA prevented gliogenesis and promoted neurogenesis²¹.

Finally, we analysed the effects of a histone acetyltransferase inhibitor anacardic acid (AA) on H1–GFP dynamics in ESCs. As expected, the dynamic association of H1–GFP with chromatin was significantly reduced in ESCs treated for 1 h with $50 \mu\text{M}$ of AA (Fig. 2i). We repeated these experiments in differentiating ESCs after 4 days of RA treatment and found a similar restriction of H1–GFP dynamics in both AA-treated and non-treated cells, as previously observed⁷. Taken together, these results show that histone acetylation regulates chromatin protein mobility in euchromatin, and apparently maintains ESCs undifferentiated while facilitating neuronal differentiation of NPCs.

H3K9 methylation affects H1 dynamics in ESCs. Relatively low H3K9 methylation levels are a hallmark of chromatin in undiffer-

entiated ESCs^{7,13,14} and thus a promising candidate for restricting chromatin plasticity during ESC differentiation. We tested previously characterized mutant ESCs that lack either G9a (ref. 22) or Suv39h1/2 (ref. 23), the H3K9 histone methyltransferases. As ESCs gain H3K9 methylation following differentiation, we compared between mutant and WT undifferentiated ESCs and differentiating NPCs. We detected no difference in the dynamic association of H1–GFP with chromatin in Suv39h1/2 mutant cells; however, in $G9a^{-/-}$ ESCs, H1–GFP dynamics were not restricted following differentiation by RA treatment for 4 days, as observed in WT ESCs⁷ (Fig. 3a–d). In $G9a^{-/-}$ ESCs with G9a re-introduced ($G9a$ -Add-WT)²⁴, normal kinetics were observed with reduced H1–GFP dynamics following differentiation (Fig. 3e,f). In contrast, when a mutant G9a lacking catalytic activity was re-introduced into $G9a^{-/-}$ ESCs ($G9a$ -Add-Mut)²⁴, the Null phenotype was observed (Fig. 3g,h), suggesting that G9a catalytic activity is required to restrict chromatin protein dynamics during ESC differentiation. Because G9a mutants can still form heterodimers with GLP and bind chromatin²⁴, chromatin binding is not sufficient to mediate restriction of chromatin dynamics.

To test the relationship between histone methylation and acetylation and to ask whether these pathways are linked in terms of chromatin plasticity, we treated $G9a^{-/-}$ ESCs with VPA for 24 h and measured H1–GFP mobility using FRAP. We found that VPA

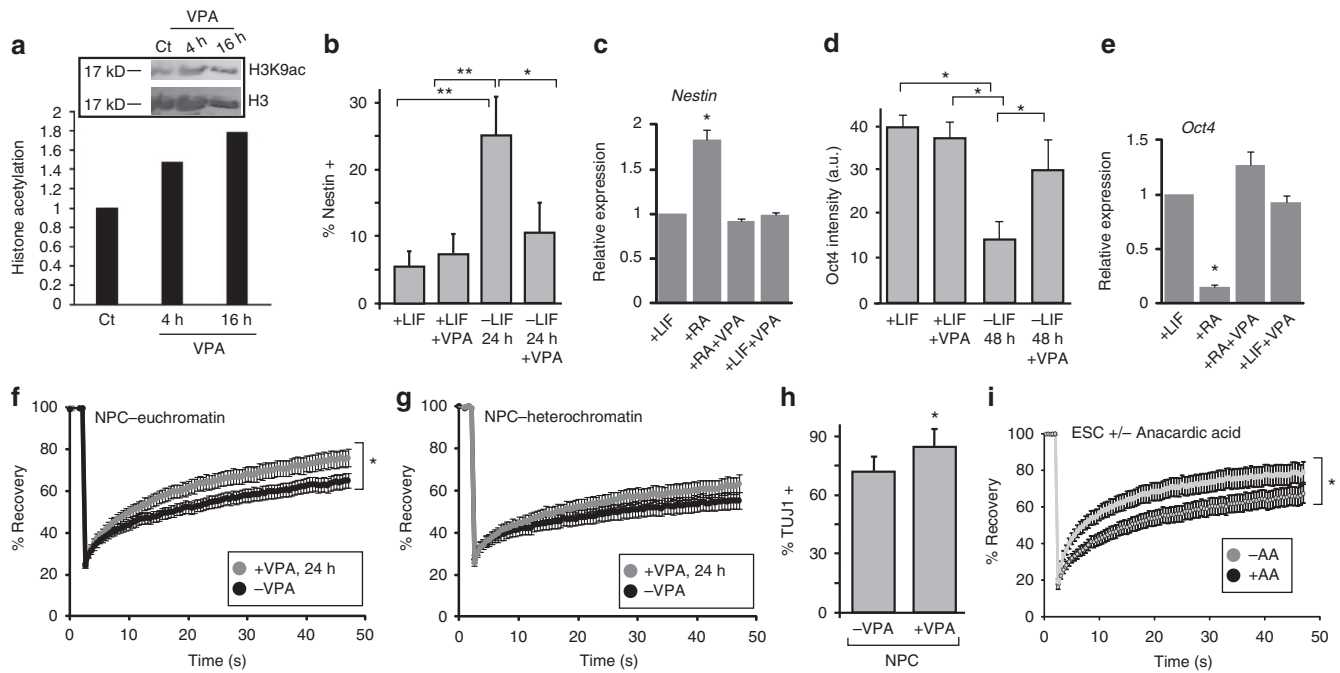


Figure 2 | Increased histone acetylation affects pluripotency and differentiation. (a) Western blot (inset) and densitometric quantification of histone acetylation levels following 4 and 16 h of VPA treatment. Unmodified histone H3 was used for normalization. (b) In the presence of LIF, ~5% of the ESC colonies display nestin-positive cells. When LIF is removed, ~25% of the cells become nestin-positive after 24 h ($P = 3 \times 10^{-8}$, 2-tailed t-test), but this number drops to ~10% when the cells are also treated with VPA ($P < 0.05$, 2-tailed t-test). y Axis denotes the % of nestin-positive colonies. (c) Quantitative real-time RT-PCR analyses of *nestin* expression in R1 ESCs treated or untreated with RA for 48 h in the absence or presence of VPA. (d) Over 95% of undifferentiated ESCs were Oct4-positive. Oct4 levels were significantly reduced 48 h after LIF withdrawal but largely restored when also treated with VPA ($P < 0.05$, 2-tailed t-test). At least 100 cells were analysed in 2 independent experiments. y Axis denotes fluorescence intensity. (e) Quantitative real-time RT-PCR analyses of Oct4 expression in R1 ESCs treated or untreated with RA for 48 h in the absence or presence of VPA. (f,g) FRAP analysis of H1e-GFP in 7-days NPCs differentiated from ESCs treated (grey) or untreated (black) with VPA in both euchromatin (f) and heterochromatin (g). Changes for euchromatin are significant ($P < 0.05$, 2-tailed t-test). (h) VPA treatment increases TUJ1 positive cells from ~74 to ~84% ($P < 0.05$, 2-tailed t-test). At least 100 cells were analysed in 2 independent experiments. (i) FRAP analysis of H1e-GFP in ESCs treated (black) or untreated (grey) with AA for 1 h. Changes are significant ($P < 0.05$, 2-tailed t-test). In panels a–e and h, values are from 2–3 biological replicates; in panels f–g and i, values are from at least 2 independent experiments, at least 20 cells each. Error bars represent s.d.

induced hyperdynamic binding of H1-GFP in $G9a^{-/-}$ ESCs (Fig. 3i), as we observed in WT ESCs (Fig. 1). We repeated the experiment in $G9a^{-/-}$ ESCs after 4 days of RA-induced differentiation. Once again, we found that VPA increased mobility of H1-GFP, although to a somewhat lesser extent than the undifferentiated ESCs (Fig. 3j). These data suggest that the two pathways of histone acetylation and $G9a$ -mediated methylation are not directly linked and that at least some HDACs bind chromatin in the absence of $G9a$.

We tested whether $G9a$ overexpression would have the opposite effect in ESCs. Overexpressing $G9a$ in WT ESCs for 72 h reduced H1-GFP dynamics (Supplementary Fig. S3). When subjected to *in vitro* neuronal differentiation, the $G9a^{-/-}$ cells failed to differentiate into stage IV and stage V neurons and died shortly before or shortly after they became stage III NPCs (Fig. 3k). In contrast, both $G9a$ -addback ($G9a$ -Add-WT) and $Suv39h1/2^{-/-}$ ESCs differentiated normally (Fig. 3k), although $Suv39h1/2^{-/-}$ ESCs tended to differentiate spontaneously and precociously in the absence of MEFs. As $G9a^{-/-}$ ESCs failed to generate mature neurons, we tested their differentiation potential *in vivo* by injecting $G9a^{-/-}$ ESCs into severe combined immunodeficient (SCID) mice and examining the teratomas 21 days later. We found mostly undifferentiated mesoderm and some residual endoderm but very little evidence of neural ectoderm (Fig. 3l). This suggests that $G9a$ might be essential not only for proper heterochromatin formation during ESC differentiation, but also for proper differentiation and/or pluripotency of ESCs.

Considering the possibility that pluripotency factors directly regulate $G9a$, we examined ChIP-seq data of pluripotency factors²⁶ at the $G9a$ locus, but found no association with the $G9a$ promoter. Also, overexpression of pluripotency factors in ESCs²⁷ did not alter $G9a$ expression levels, and the level and distribution of both H3K9 histone methyltransferases did not change significantly during early ESC differentiation^{6,28,29}. Finally, the levels of pluripotency genes in ESCs remained unaltered in the absence of $G9a$ (ref. 30). We conclude that $G9a$ is not directly bound or regulated by pluripotency factors in ESCs. Taken together, these data suggest that increased H3K9 methylation in differentiating ESCs restricts chromatin protein dynamics and that $G9a$ is required for proper differentiation of ESCs.

Lamin A partially restricts heterochromatin H1 dynamics. We hypothesized that LMNA may be involved in the restriction of chromatin dynamics following ESC differentiation based on several facts. First, LMNA is not detected in undifferentiated ESCs³ (Fig. 4a), and their nuclear lamina, marked by lamin B, appears less structured than in differentiated cells^{7,31–33}. Moreover, LMNA appears early during differentiation, approximately when chromatin-binding proteins become less mobile³. Finally, heterochromatin domains are frequently found at the nuclear periphery and LMNA directly associates with chromatin by binding core histones^{4,5}. We therefore tested the relationship between LMNA expression and chromatin plasticity.

To test the contribution of LMNA to chromatin dynamics restriction during ESC differentiation, we ectopically expressed

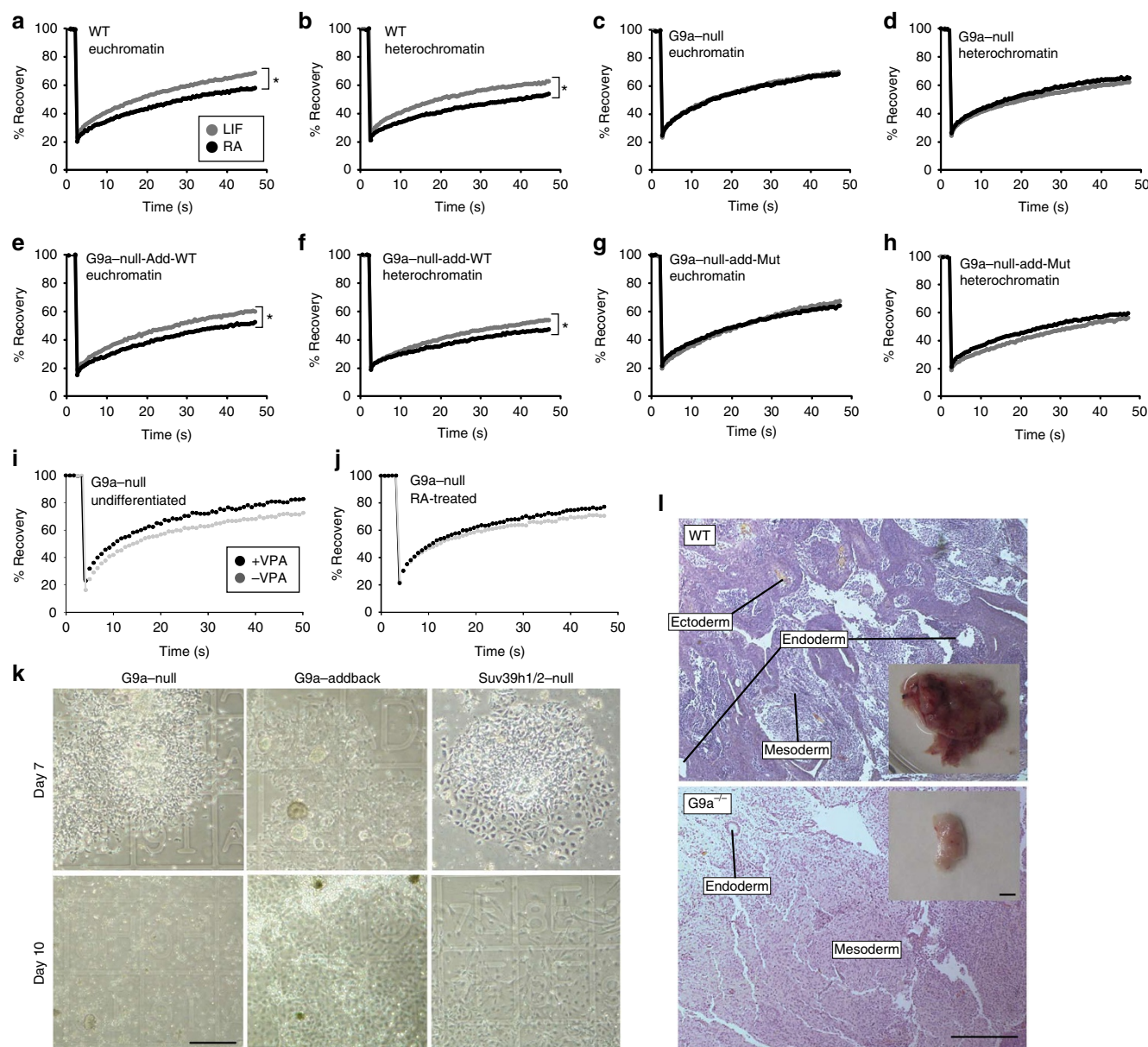


Figure 3 | H3K9 methylation affects H1 dynamics and differentiation. (a,b) H1e-GFP FRAP analysis in euchromatin (a) and heterochromatin (b) of WT ESCs, undifferentiated (LIF, grey) or differentiated with retinoic acid for 4 days (RA, black). Changes are significant ($P < 0.01$, 2-tailed t -test). (c,d) H1e-GFP FRAP analysis in euchromatin (c) and heterochromatin (d) of G9a^{-/-} ESCs, undifferentiated (grey) or differentiated with RA for 4 days (black). No restriction of H1 dynamics is observed following differentiation. (e,f) H1e-GFP FRAP analysis in euchromatin (e) and heterochromatin (f) of G9a^{-/-} ESCs in which WT G9a was added-back (G9a-Null-add-WT), undifferentiated (grey) or differentiated with RA for 4 days (black). Changes are significant ($P < 0.05$, 2-tailed t -test). (g,h) H1e-GFP FRAP analysis in euchromatin (g) and heterochromatin (h) of G9a^{-/-} ESCs in which a mutant form of G9a lacking catalytic activity was added-back (G9a-Null-add-Mut), undifferentiated (grey) or differentiated with RA for 4 days (black). No restriction of chromatin protein dynamics is observed following differentiation. (i,j) H1e-GFP FRAP analysis in G9a^{-/-} ESCs, undifferentiated (i) and differentiated for 4 days with RA (j), treated (black) or untreated (grey) with 0.5 mM VPA for 24 h. In both cases, changes are significant ($P < 0.005$, 2-tailed t -test). (k) Neuronal differentiation of G9a^{-/-}, G9a-addback and Suv39h1/2^{-/-} ESCs. All cells reach the NPC stage at day 7 (top) but the G9a^{-/-} cells die between day 7 and day 10 while the G9a-addback and the Suv39h1/2^{-/-} cells continue to differentiate normally. Scale bar, 100 μ m. (l) Teratoma analysis in SCID mice injected with WT (top) or G9a^{-/-} (bottom) ESCs. WT ESCs displayed normal differentiation with apparent endoderm, mesoderm and ectoderm. However, teratomas made from G9a^{-/-} ESCs were significantly smaller (inset) and did not show neuroectodermal differentiation. Scale bar, 0.5 mm; inset scale bar, 5 mm. Values are from at least 2 independent experiments, at least 20 cells each.

LMNA together with H1e-GFP in undifferentiated ESCs and analysed H1e-GFP dynamics. Ectopic LMNA properly localized in the ESC nuclear lamina. This was viewed in live cells expressing Cherry-tagged human- and mouse-LMNA (Supplementary Fig. S4) and in fixed cells that expressed untagged LMNA, detectable by anti-LMNA antibodies (Fig. 4b). We used untagged LMNA

because anti-LMNA antibodies failed to recognize Cherry-tagged LMNA. Lamin B staining revealed the less structured nuclear lamina in undifferentiated as opposed to differentiated ESCs². H1e-GFP FRAP experiments were performed in ESCs with or without co-expression of LMNA (Fig. 4b). When comparing H1e-GFP dynamics in cells co-expressing LMNA with cells transfected only

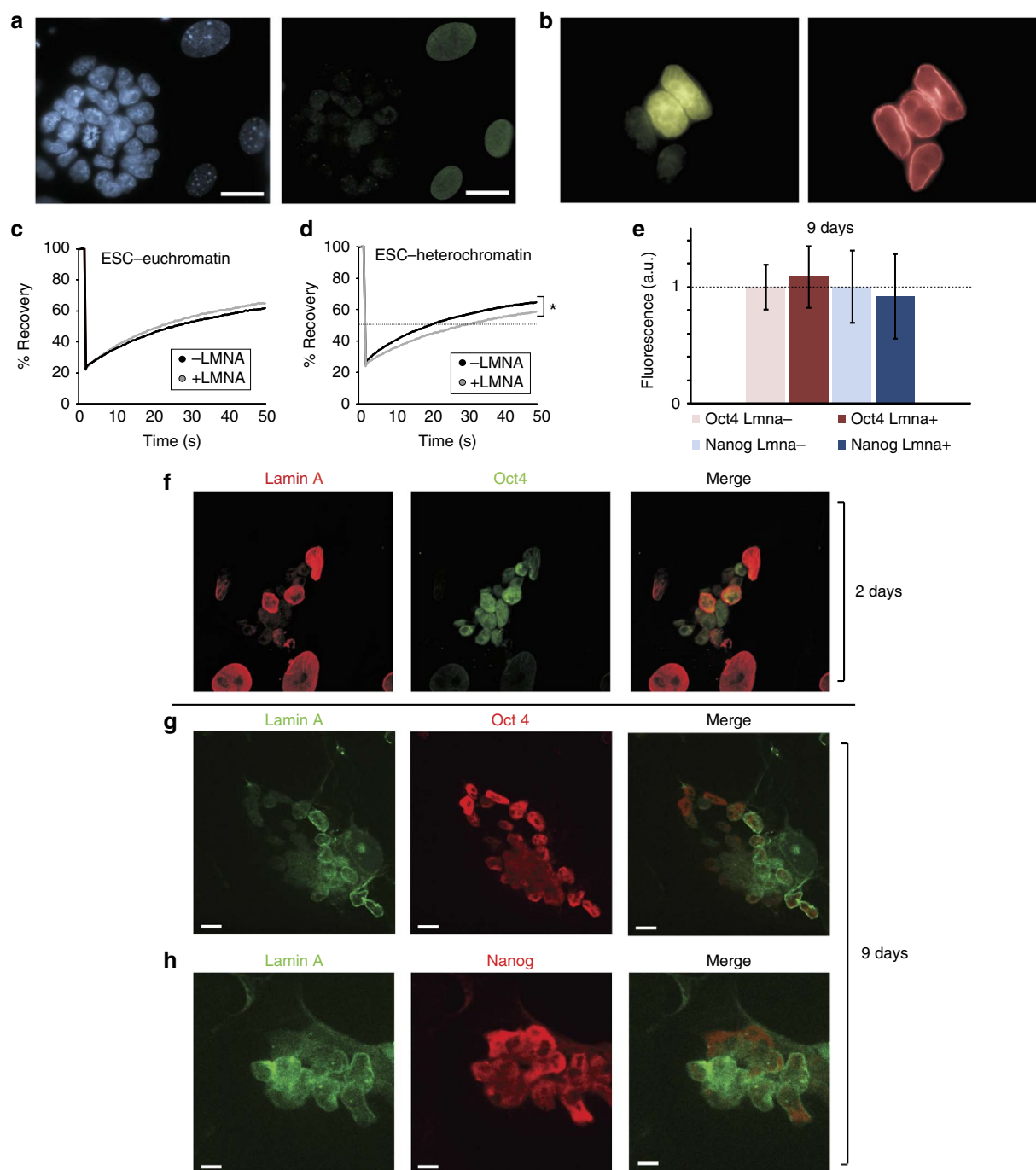


Figure 4 | LMNA partially restricts H1 dynamics in heterochromatin. **(a)** Stained with DAPI (blue), an R1 mouse ESC colony is surrounded by three fibroblasts. Fibroblasts stain positive with anti-LMNA antibodies (green); ESCs do not. **(b)** R1 ESCs were co-transfected with both *Lmna* and *H1e-GFP* at a 10:1 ratio. Anti-LMNA antibodies (red) recognize the exogenously expressed LMNA. Because of the transfection ratio between the two plasmids, all *H1e-GFP*-positive cells were LMNA-positive. **(c,d)** *H1e-GFP* FRAP analysis in R1 ESCs ectopically expressing (grey) or not expressing (black) WT mouse LMNA. No significant changes were observed in euchromatin **(c)** but changes in heterochromatin **(d)** are significant ($P < 0.05$, 2-tailed *t*-test). **(e)** Quantitative analysis of immunofluorescence levels of Oct4 and Nanog is shown before and after 9 days of ectopic LMNA expression. **(f)** Forty-eight hours after *Lmna* transfection, all R1 ESCs stain positive for Oct4 (green), regardless of whether they stain positive or negative for LMNA (red). The large nuclei are MEFs that served as LMNA-positive controls and Oct4-negative controls. **(g,h)** Nine days after *Lmna* transfection, all R1 ESCs stain positive for Oct4 and Nanog (red), regardless of whether they stain positive or negative for LMNA (green). In all panels, values are from 2–3 biological replicates. Error bars represent s.d.

with *H1e-GFP*, we found a slight reduction in chromatin protein dynamics (Supplementary Fig. S4e).

We repeated the LMNA and *H1e-GFP* co-transfection experiments and analysed recovery rates in euchromatin and heterochromatin,

separately. Whereas *H1e-GFP* dynamics displayed similar kinetics in euchromatic regions of LMNA-expressing and non-expressing cells (Fig. 4c), a significant reduction in H1 mobility was recorded in discrete heterochromatin foci (Fig. 4d). Reduced plasticity was not

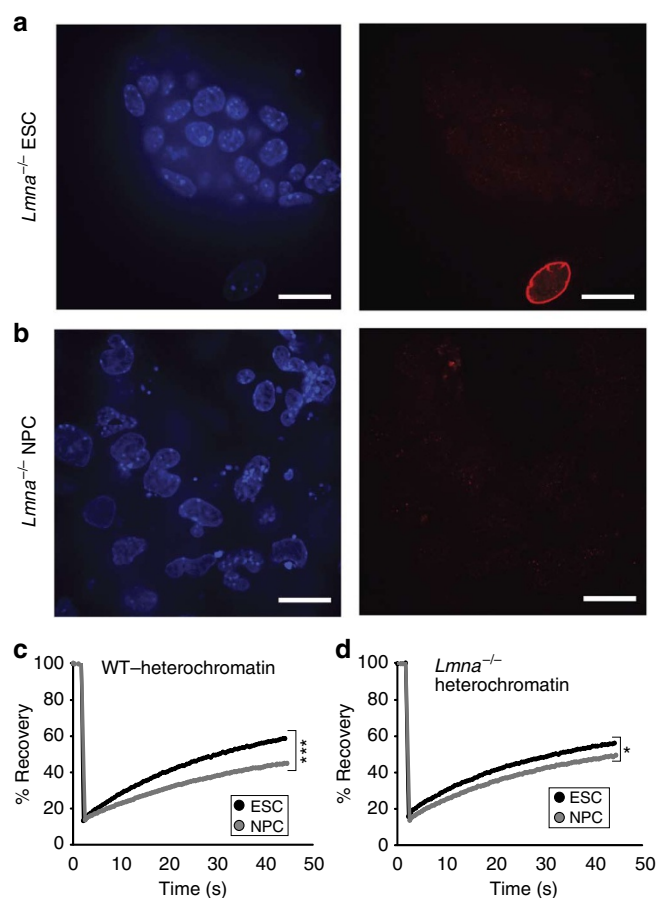


Figure 5 | *Lmna*^{-/-} NPCs exhibit reduced heterochromatin restriction.

(a, b) *Lmna*^{-/-} ESCs (a) and NPCs (b) stain negative for LMNA, unlike the fibroblast that expresses LMNA (red). DAPI is shown in blue. Note aberrant nuclear shapes of *Lmna*^{-/-} NPCs. (c) H1e-GFP FRAP analysis in heterochromatin of undifferentiated R1 ESCs (black) and NPCs (grey). Changes are significant ($P < 0.001$, 2-tailed t -test). (d) H1e-GFP FRAP analysis in heterochromatin of undifferentiated *Lmna*^{-/-} ESCs (black) and NPCs derived from *Lmna*^{-/-} ESCs (grey). Changes are significant ($P < 0.05$, 2-tailed t -test). Values are from at least 2 independent experiments, at least 20 cells each.

caused by forced differentiation of LMNA-expressing ESCs because they did not alter their pluripotency markers, 2 or 9 days after transfection (Fig. 4e–h). These results indicate that ectopic expression of LMNA in ESCs restricts chromatin protein dynamics exclusively in heterochromatin.

We next tested whether LMNA is necessary to restrict chromatin protein dynamics during ESC differentiation. We generated *Lmna*^{-/-} NPCs (Fig. 5a,b) and measured H1e-GFP dynamics in heterochromatin and euchromatin, separately. Interestingly, H1 dynamics restriction, characteristic of the NPC stage or earlier⁷ (Fig. 5c), was significantly, albeit not entirely, abolished in *Lmna*^{-/-} NPCs (Fig. 5d). Additionally, whereas H1 kinetics in heterochromatin are usually slower than in euchromatin^{7,15,34,35} (Fig. 1), in *Lmna*^{-/-} cells, H1 kinetics in heterochromatin were similarly hyperdynamic as in euchromatin (Supplementary Fig. S4f–g). This interesting observation reflects the hyperdynamic heterochromatin in *Lmna*^{-/-} NPCs, suggesting perturbed heterochromatin formation or maintenance in *Lmna*^{-/-} ESC differentiation. Confirming this notion, nuclear shape in *Lmna*^{-/-} NPCs was highly irregular (Figs 5b and 6b,d), and the number of conspicuous heterochromatin foci was dramatically lower than in WT NPCs (Fig. 5e,f).

Finally, to check how LMNA affects differentiation potential, we stained *Lmna*^{-/-} NPCs at day 6 for early and late neuronal markers nestin and β -tubulin-III (TUJ1), respectively. Whereas nestin was detected in both WT and *Lmna*^{-/-} NPCs, β -tubulin-III was only detected in WT NPCs and was completely absent from *Lmna*^{-/-} NPCs (37% versus 0%, $P < 0.005$, 2-tailed t -test).

These observations did not recur in *Lmna*^{-/-} MEFs derived from *Lmna*^{-/-} mice³⁶. Comparing H1e-GFP dynamic association with chromatin in WT versus *Lmna*^{-/-} MEFs, we detected no significant differences in either euchromatin or heterochromatin regions (Supplementary Fig. S5). *Lmna*^{-/-} MEFs may harness other mechanisms to maintain normal heterochromatin and lamina-chromatin association and function as they exhibit similarly abundant and conspicuous heterochromatin foci as in WT MEFs, whereas differentiating *Lmna*^{-/-} ESCs fail to organize heterochromatin (Fig. 5e,f).

Taken together, these results suggest that LMNA affects chromatin plasticity during ESC early differentiation, restricting chromatin protein dynamics exclusively in heterochromatin, and is required for normal heterochromatin organization during ESC differentiation. In the absence of LMNA, differentiating cells fail to restrict chromatin protein dynamics, fail to obtain normal nuclear conformation and heterochromatin organization and display altered differentiation propensity.

DNA methylation does not affect H1 dynamics in ESCs. Obtained from the developing embryo's inner cell mass, ESCs have relatively low DNA methylation levels on derivation and they subsequently acquire DNA methylation during culturing³⁷. Therefore, we hypothesized that DNA methylation might influence chromatin protein dynamics. To determine whether global DNA methylation levels contribute to the hyperdynamic nature of chromatin proteins in ESCs, we treated ESCs stably expressing H1-GFP with the DNA demethylating agents 5-Azacytidine (5-Aza) and Zebularine (Zeb) (Supplementary Fig. S6a) for 5 days, and performed FRAP analyses. We determined global DNA methylation levels using the nearest-neighbour method. We found that DNA methylation was reduced by ~30% following either 5-Aza or Zeb treatment (Supplementary Fig. S6b). Despite the decrease in DNA methylation levels, H1-GFP dynamics showed no significant changes following the different treatments (Fig. 7).

To reach lower levels of DNA methylation, we further analysed chromatin protein dynamics in ESCs lacking one or more DNA methyltransferase (DNMT) genes. We tested ESCs lacking the maintenance DNMT gene *Dnmt1*³⁸ and ESCs lacking both *de novo* DNMT genes, *Dnmt3a* and *Dnmt3b*³⁹. Nearest-neighbour analysis demonstrated that global DNA methylation levels were reduced by ~80% in *Dnmt1*^{-/-} cells and by more than 90% in *Dnmt3* double knockout cells (Supplementary Fig. S6b). We note that this method gives a gross approximation, not absolute values, but our levels are in good agreement with previous reports⁴⁰. FRAP analyses showed a very slight, statistically insignificant, increase in H1e-GFP dynamics in both *Dnmt1*^{-/-} (Fig. 7b,f) and *Dnmt3a/3b*^{-/-} (Fig. 7c,g) ESCs tested. To exclude the possibility that DNMTs are functionally redundant with respect to chromatin dynamics, we further analysed ESCs lacking all three DNMT genes (*Dnmt1/3a/3b* triple knockout (TKO))⁴⁰. In these *Dnmt*-TKO ESCs, we observed a slightly more pronounced increase in H1-GFP mobility, albeit not statistically significant (Fig. 7d,h). HP1-GFP and HMGN1-GFP both displayed similar kinetics between WT ESCs and the different mutants used (*Dnmt1*^{-/-}, *Dnmt3*^{-/-} and *Dnmt*-TKO), as did H1-GFP (Supplementary Fig. S7).

As DNA methylation levels had very little effect on chromatin plasticity in undifferentiated ESCs, we searched for an effect during early differentiation. Although embryoid body (EB) formation kinetics were faster in *Dnmt*^{-/-} ESCs than in WT cells, the double

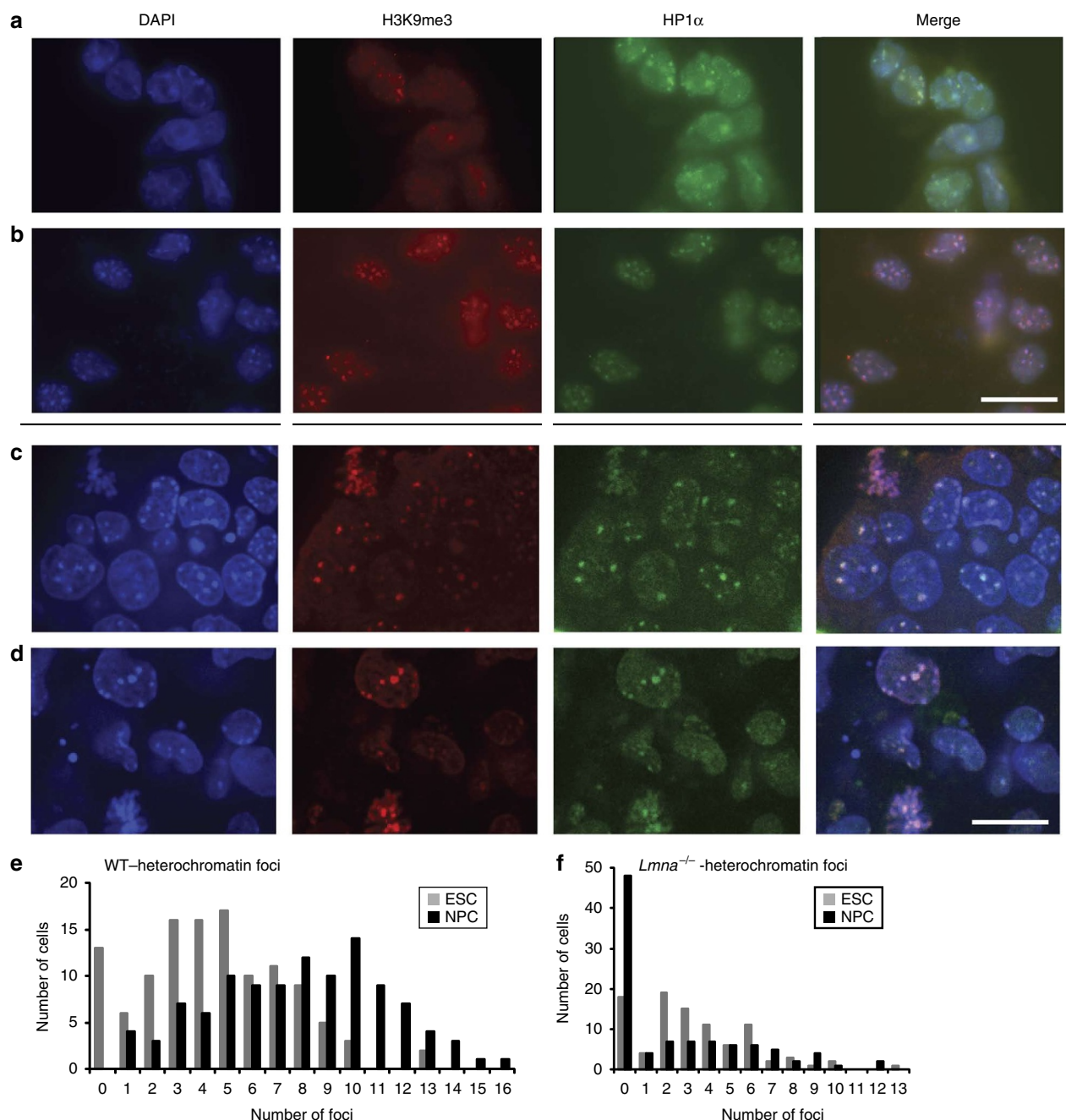


Figure 6 | *Lmna*^{-/-} NPCs exhibit abnormal heterochromatin structure. (a,b) DAPI (blue) and immunostaining for heterochromatin markers H3K9me3 (red) and HP1α (green) in WT mouse ESCs (a) and NPCs (b). Scale bar, 20 μm. (c,d) DAPI (blue) and immunostaining for heterochromatin markers H3K9me3 (red) and HP1α (green) in *Lmna*^{-/-} ESCs (c) and in *Lmna*^{-/-} NPCs (d). Scale bar, 20 μm. (e) Distribution of heterochromatin foci number per nucleus in WT ESCs (grey) and NPCs (black). The average number of foci per nucleus increased from 4.5 ± 5.9 in ESCs to 7.9 ± 4.1 in NPCs. (f) Distribution of heterochromatin foci number per nucleus in *Lmna*^{-/-} ESCs (grey) and NPCs (black). The average number of foci per nucleus decreased from 3.3 ± 6.8 in ESCs to 2.6 ± 12.1 in NPCs. At least 100 cells were counted in 2 independent experiments.

mutants failed to differentiate properly into NPCs (Supplementary Fig. S8a–b). We therefore induced differentiation, using RA, and analysed H1e–GFP dynamics after 24 and 48 h. Within 24 h of RA treatment, we observed no appreciable differences in H1e dynamics (Supplementary Fig. S8c) but within 48 h of RA treatment, we observed a notable restriction in H1e dynamics in *Dnmt3*^{-/-} ESCs when compared with WT ESCs (Supplementary Fig. S8d). This demonstrates that *de novo* DNA methylation is required during ESC differentiation, but suggests no involvement of DNA methylation in restriction of chromatin plasticity.

Considering potential secondary effects on histone modifications in *Dnmt* knockout cells⁴¹, we examined histone acetylation and H3K9 methylation levels in *Dnmt*-TKO and WT cells, and found no significant differences (Supplementary Fig. S8e). These data indicate that with the exception of a small contribution to H1 dynamics, global levels of DNA methylation do not account for the altered dynamic nature of chromatin proteins in ESCs.

Nucleosome spacing does not affect H1 dynamics in ESCs. We then asked whether global changes in chromatin architecture, such

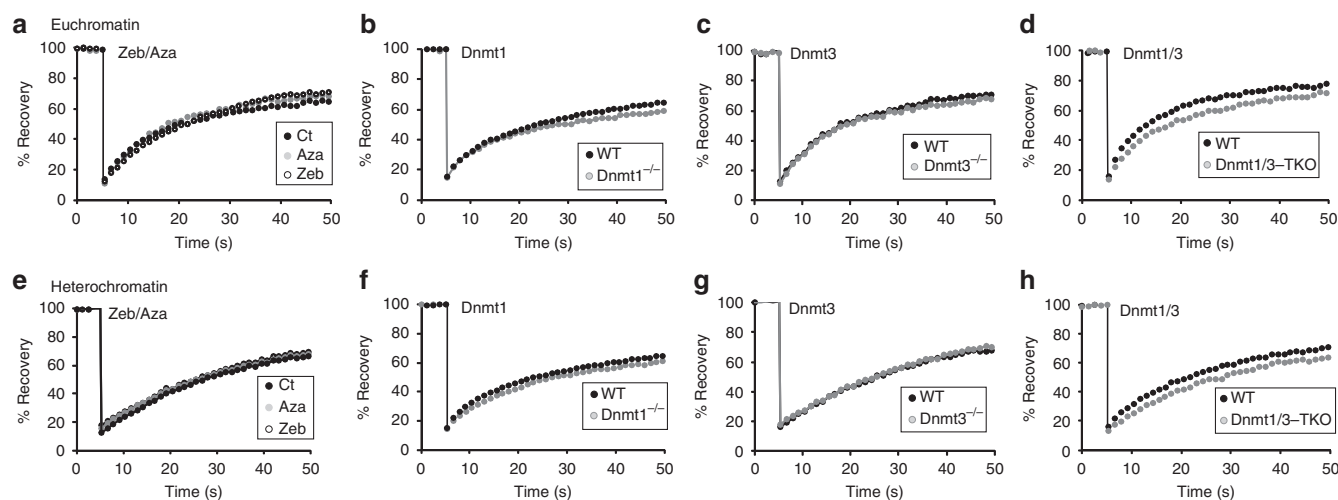


Figure 7 | DNA methylation does not affect H1 dynamics in ESCs. (a) H1e-GFP FRAP analysis in euchromatin of R1 ESCs, treated for 5 days with either 100 mM 5-Aza (grey) or 1 mM Zeb (empty) or untreated (Ct, black). (b) H1e-GFP FRAP analysis in euchromatin of Dnmt1 knockout ESCs (Dnmt1^{-/-}, grey) compared with wild-type controls (WT, black). (c) H1e-GFP FRAP analysis in euchromatin of Dnmt3a/b double-knockout ESCs (Dnmt3^{-/-}, grey) compared with wild-type controls (WT, black). (d) H1e-GFP FRAP analysis in euchromatin of Dnmt3a/b and Dnmt1 triple-knockout ESCs (Dnmt1/3-TKO, grey) compared with wild-type controls (WT, black). Changes are not significant ($P = 0.066$, 2-tailed t -test). (e) H1e-GFP FRAP analysis in heterochromatin of R1 ESCs, treated for 5 days with either 100 mM 5-Aza (grey) or 1 mM Zeb (empty) or untreated (Ct, black). (f) H1e-GFP FRAP analysis in heterochromatin of Dnmt1 knockout ESCs (Dnmt1^{-/-}, grey) compared with wild-type controls (WT, black). (g) H1e-GFP FRAP analysis in heterochromatin of Dnmt3a/b double-knockout ESCs (Dnmt3, grey) compared with wild-type controls (WT, black). (h) H1e-GFP FRAP analysis in heterochromatin of Dnmt3a/b and Dnmt1 triple-knockout ESCs (Dnmt1/3-TKO, grey) compared with wild-type controls (WT, black). Changes are not significant ($P = 0.07$, 2-tailed t -test). Values are from at least 2 independent experiments, at least 20 cells each.

as altered nucleosome spacing, would influence chromatin protein dynamics. We examined ESCs lacking three different H1 genes, *H1c*, *H1d* and *H1e* (H1 triple-knockout, H1-TKO)⁴², which have reduced chromatin repeat length and altered chromatin structure⁴³. FRAP analysis demonstrated that the absence of these H1 variants did not affect H1-GFP dynamics (Fig. 8). H1-TKO ESCs lack 50% of WT H1 levels⁴², so ectopically expressed H1-GFP might partially or fully restore H1 levels, rescue the H1 deficiency phenotype and thus obscure its effect on chromatin dynamics. We therefore compared HP1-GFP and HMGN1-GFP dynamics in H1-TKO cells versus WT ESCs, but neither protein had detectably different chromatin dynamics in these cells (Fig. 8). These results suggest that reduced histone H1 levels in H1-TKO cells and the resulting altered chromatin structure do not affect chromatin protein dynamics and do not explain the observed differences in chromatin plasticity between ES and differentiating cells.

Discussion

Here we investigate mechanisms underlying hyperdynamic and plastic properties of chromatin observed in undifferentiated ESCs⁷. We find that histone hyper-acetylation enhances euchromatin dynamics in the undifferentiated state, H3K9 methylation affects both euchromatin and heterochromatin dynamics, and LMNA restricts heterochromatin dynamics in early differentiation. These observations are in agreement with the fact that ESCs are devoid of LMNA³ and that ESC chromatin has relatively high levels of histone acetylation and relatively low levels of H3K9 methylation^{6,7,13,14,16,18,19}. Importantly, these observations suggest a direct link between chromatin dynamics and ESC pluripotency. This notion is supported by the fact that increased histone acetylation caused by HDACi treatment, promotes and facilitates reprogramming of somatic cells into pluripotent cells and helps maintain ESCs in an undifferentiated state^{44,45}. Moreover, re-expression of Oct4 during cell reprogramming has been shown to be inhibited by G9a-mediated H3K9 methylation²⁵ whereas the small molecule BIX-01294 (BIX), a G9a

inhibitor, improves reprogramming efficiency of primary NPCs transduced with *Oct4* and *Klf4* alone to a level comparable to 4-factor transduction (*Oct4*, *Klf4*, *Sox2* and *cMyc*)⁴⁶. It thus seems that both histone modification pathways we identified here as important for chromatin plasticity are also important for pluripotency. Notwithstanding, Suv39h1/2 knockout ESCs displayed no phenotype regarding chromatin plasticity or differentiation when induced to differentiate into neurons. It is likely that Suv39h1/2 participate in alternate differentiation routes rather than neuronal. Supporting this notion is a recent report showing the specific targeting of Suv39h1 to bivalent genes on trophoblast lineage commitment⁴⁷.

LMNA presents an additional pathway to restrict chromatin plasticity. When ectopically expressed in WT ESCs, LMNA restricted heterochromatin dynamics, perhaps by its direct association with core histones^{4,5}. Our results also raise the possibility of a connection between LMNA and pluripotency, differentiation, and reprogramming. One such potential link may be through restricting pervasive transcription during differentiation^{48,49}. Tethering a genomic locus to the nuclear lamina in somatic cells, which was shown to restrict transcription⁵⁰, was accompanied by LMNA accumulation at the tethered site. It would therefore be interesting to test whether LMNA expression restricts pervasive transcription in ESCs^{6,51}.

It is likely that additional mechanisms have a role in regulating chromatin plasticity in ESCs. ATP-dependent chromatin remodeling proteins, prevalent in undifferentiated ESCs⁶, are good candidates. Recently, Chd1 was shown to regulate pluripotency in ESCs⁵². We found that stable downregulation of Chd1 expression increased heterochromatin formation and reduced chromatin protein dynamics⁵², suggesting that Chd1 facilitates an open chromatin conformation in undifferentiated ESCs. It would be interesting to decipher the role(s) that other chromatin remodellers have in stem cell biology because accumulating evidence suggest they are highly important for the stem cell state^{6,52–54}.

Interestingly, we found that DNA methylation levels, although crucial for ESC differentiation, have very little role, if any, in

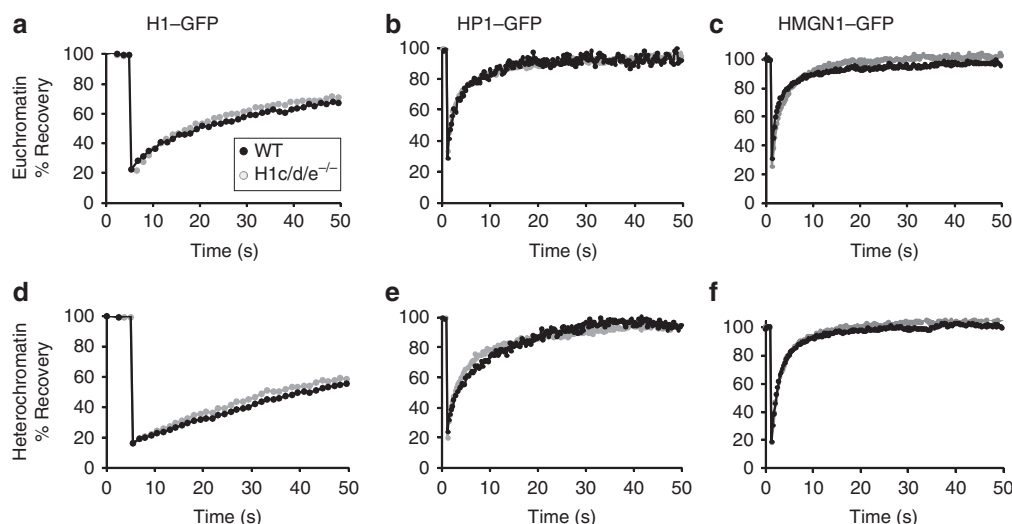


Figure 8 | Nucleosome spacing does not affect chromatin protein dynamics in ESCs. (a) H1e-GFP FRAP analysis in euchromatin of H1c/d/e triple-knockout ESCs (H1c/d/e^{-/-}, grey) compared with wild-type controls (WT, black). (b) HP1-GFP FRAP analysis in euchromatin of H1c/d/e triple-knockout ESCs (grey) compared with wild-type controls (black). (c) HMGN1-GFP FRAP analysis in euchromatin of H1c/d/e triple-knockout ESCs (grey) compared with wild-type controls (black). (d) H1e-GFP FRAP analysis in heterochromatin of H1c/d/e triple-knockout ESCs (grey) compared with wild-type controls (black). (e) HP1-GFP FRAP analysis in heterochromatin of H1c/d/e triple-knockout ESCs (grey) compared with wild-type controls (black). (f) HMGN1-GFP FRAP analysis in heterochromatin of H1c/d/e triple-knockout ESCs (grey) compared with wild-type controls (black). No significant changes were observed for any of the tested proteins. Values are from at least 2 independent experiments, at least 20 cells each.

regulating chromatin protein dynamics in ESCs. In another study⁵⁵, a slight decrease in linker histone kinetics was observed in *Dnmt3*^{-/-} ESCs. Although these observations are in line with our measured kinetics in differentiating *Dnmt3*^{-/-} ESCs, we observed no differences in the undifferentiated state. This could be owing to differences in GFP-tag location, the nature of the linker histone used or differences between different ESC lines. Regardless, our studies indicate that DNA methylation is not a major factor responsible for the dynamic nature of chromatin proteins in ESCs.

Perhaps more surprising is the fact that global chromatin structure had no apparent effect on chromatin protein dynamics in ESCs. ESCs lacking three linker histones displayed reduced local chromatin compaction and decreased global nucleosome spacing⁴² but their chromatin dynamics remained unaltered. It is possible that local, not global, chromatin dynamics have been altered, but this cannot be detected using FRAP.

To conclude, we provide mechanistic insight into our previous observations that chromatin proteins are hyperdynamic in ESCs. We demonstrate that reduced histone acetylation, increased H3K9 methylation and LMNA expression, all act to restrict chromatin plasticity during ESC differentiation. Thus, undifferentiated ESCs, which possess high levels of histone acetylation, relatively low levels of H3K9 methylation and no LMNA, are able to maintain a plastic chromatin state that supports pluripotency.

Methods

Cells. Mouse R1 and E14 ESCs were from ATCC; *Dnmt1*^{-/-} ESCs from R. Jaenisch (Cambridge, MA, USA); *Dnmt3a*/b^{-/-} from E. Li (Cambridge, MA, USA); *Dnmt1/3a*/b^{-/-} from M. Okano (Kobe), *Suv39h1/2*^{-/-} from T. Jenuwein (Freiburg); *G9a*^{-/-}, *GW1* (*G9a* addback), *G44* (*G9a* mutant-addback) and *TT2* (the parental strain of the 3 *G9a* derivatives) from Y. Shinkai (Kyoto). Pluripotent ESCs were grown on γ -irradiated or mitomycin-C-treated MEFs in DMEM, 10–15% ESC-grade FCS, 1 mM sodium pyruvate, 0.1 mM nonessential amino acids, 0.1 mM β -mercaptoethanol, and 1,000 U ml⁻¹ LIF. LIF withdrawal differentiation was performed by separating ESCs from MEFs, plating them on gelatin-coated dishes and culturing in ES medium without LIF. RA-induced differentiation was performed in a similar way with the addition of 1 μ M RA. For NPC differentiation, ESCs were

separated from MEFs and grown in suspension on bacterial culture dishes without LIF for 4 days to allow for EB formation. EBs were replated on poly-L-ornithine/fibronectin (Sigma)-coated plates in DMEM/F12 medium supplemented with ITS (5 mg ml⁻¹ insulin, 50 mg ml⁻¹ transferrin, 30 nM selenium chloride) and fibronectin (5 mg ml⁻¹) and grown for 2–6 days longer until NPC day 6–10, respectively. All cell culture media were supplemented with 2 mM glutamine, 100 U ml⁻¹ penicillin, and 50 mg ml⁻¹ streptomycin. All cell culture reagents were purchased from GIBCO-BRL (Invitrogen; Carlsbad, CA, USA). DNA transfections were performed with TransIt (Mirus) and Lipofectamine-2000, according to the manufacturer's instructions. For photobleaching experiments, cells were grown in 8-well μ -Slides (ibidi, Munich, Germany) or in chambered cover glasses (Lab-Tek; Rochester, NY, USA) or in glass-bottom culture dishes (MatTek; Ashland, MA, USA). 5-Aza, Zeb, TSA, VPA and AA were added to the cells at a final concentration of 0.5 μ M, 200 μ M, 50 nM, 0.5 mM and 50 μ M, respectively. 5-Aza and Zeb were applied for 5 days. TSA and VPA were applied for 4 h, 16 h or 24 h. AA was applied for 1 h.

Plasmids. H1e-GFP, H1-GFP, HP1 α -GFP, HMGN1-GFP and FLAG-HA-G9a fusion proteins have been previously described^{7,15,17,22,56,57}. Stable R1 clones expressing GFP fusion proteins were generated by introducing the expression vectors via electroporation, selection for resistance to G418, followed by identification of expressing clones by direct observation under epifluorescence. For mouse *Lmna* (*mLmna*) cloning, total RNA was extracted from MEFs using the RNeasy Mini Kit (Qiagen) and complementary DNA was prepared using an RTase kit (Applied Biosystems). The *mLmna* coding sequence was amplified by PCR with the following DNA oligos: 5'-cgatctcgaggagtgagacccgtcacagcg-3'; 5'-cgatggatcctcatgatgctgcagcttctggg-3' (Syntezza Bioscience, Israel). Products were separated in agarose gels and ligated into the T-Easy Vector system (Promega). The *mLmna* coding sequence was inserted into the pmCherry-C2 vector (Clontech), using EcoRI, and orientation was verified by sequencing. To create a WT tag-less LMNA A expression vector, mCherry was excised with AgeI and BglII and the *mLmna*-containing vector was treated with Klenow and re-ligated.

Antibodies and immunofluorescence. The following antibodies were used at the corresponding concentrations: Nestin, 1:100 whole serum antibody (rabbit polyclonal), kindly provided by R. McKay (National Institutes of Health, Bethesda, MD, USA); LMNA, 2 μ g ml⁻¹ (rabbit polyclonal, sc-20680: Santa Cruz Biotechnologies, Santa Cruz, CA, USA); TUJ1, 1:200 ascites fluid (mouse monoclonal, MAB1637: Millipore); Oct4, 1 μ g ml⁻¹ (N19 goat polyclonal, sc-8628: Santa Cruz); Nanog, 10 μ g ml⁻¹ (Goat polyclonal, AF2729: R&D systems); HP1 α :1:750 ascites fluid (Mouse monoclonal, 2HP-1H5-As: Euromedex); H3K9me3, immunofluorescence (IF) 1:100, WB 1:1,000 serum antibody (Rabbit polyclonal), kindly provided by T. Jenuwein; H3Ac, 1 μ g ml⁻¹ (Rabbit polyclonal, 06-599: Upstate). Detection for WB was with anti-rabbit or anti-mouse antibodies conjugated to HRP

(Rockland) and for IF with anti-rabbit, anti-mouse or anti-goat antibodies conjugated to Alexa488 or Alexa568 (Molecular Probes, Carlsbad, CA, USA), Cy3 or FITC (Jackson ImmunoResearch, West Grove, PA, USA). For IF, cells, grown either in 8-well μ -Slides (ibidi, Munich, Germany) on cover-slips, were fixed in 4% paraformaldehyde (15 min, room temp.), washed 3 times in PBS (5 min, room temp.), permeabilized with 0.5% Triton X-100 in PBS (5 min, room temp.) and incubated with the primary antibodies (1 h, room temp. or 4 °C overnight). Cells were then washed 3 times in PBS (5 min, room temp.), incubated with secondary antibodies (1 h, room temp.), washed again in PBS (5 min, room temp.), DAPI stained (5 min, room temp.), washed again in PBS (5 min, room temp.) and mounted on a microscope slide with anti-fade (Dako, Glostrup, Denmark).

Microscopy and photobleaching. Imaging was done as described⁷ or using a Revolution spinning disk (CSUX, Yokogawa) imaging system (Andor, UK) with solid state lasers 405, 488, 561 and 640 nm (50 mW each) mounted on an Olympus IX81 fully automated microscope equipped with an automated stage, autofocus device (Z-drift compensation) and an environmental chamber (LIS, Switzerland) controlling oxygen, humidity, CO₂ and temperature. FRAP was done using a specialized FRAPPA module (Andor, UK) at 50–100% laser intensity as described⁵⁸. Bleaching was confined to square areas of 15×15 pixels in MEFs and 12×12 pixels in ESCs and NPCs using a ×60 oil lens with a numerical aperture of 1.4. We used an EMCCD iXon+ camera (Andor, UK) with a window size of 512×512 pixels. When separately analysing euchromatin and heterochromatin, bleach areas were discretely positioned away from GFP-positive foci or directly on them, respectively. All FRAP analyses were performed on 15–30 cells from at least 2 independent experiments. Data analyses were as described⁷. Heterochromatin foci quantification was performed by manually counting DAPI dense foci that co-localize with HP1 α and H3K9me3 immunostaining. This method was verified semi-automatically with ImageJ threshold analysis.

Teratoma formation. Ten million cells suspended in 35 μ l ESC medium and 15 μ l MatrigelTM (BD Biosciences) were injected subcutaneously into the dorsal flank of SCID mice (C.B-17/lcrHsd-SCID-bg). Three weeks after the injection, mice were killed and teratomas were surgically dissected. Samples were weighed, fixed in PBS containing 4% paraformaldehyde, embedded in paraffin and sliced into 7 μ m-thin sections. Sections were stained with haematoxylin and eosin.

The joint ethics committee (IACUC) of the Hebrew University and Hadassah Medical Center approved the study protocol for animal welfare. The Hebrew University is an AAALAC International accredited institute. All animal experiments were conducted in accordance with the Hebrew University's animal committee, ethical approval # IACUC:NS-09-11616-4.

Nearest-neighbour analysis. DNA methylation levels were determined using MboI digestion and end labelling as described in detail⁵⁹.

Statistical analyses. Two-tailed Student's *t*-test was performed to compare the kinetics of the different FRAP curves. *t*-test was used to adhere to our previous analyses to allow comparisons⁷. We note that *t*-test in this regard is a conservative test and that other statistical tests (for example, Fisher's Omnibus), which take into account entire graphs may result in lower *P*-values. Graphs were compared at time *t* = 40 s, and *P*-values were calculated on these comparable time points, as described⁵⁶.

References

- Aoto, T., Saitoh, N., Ichimura, T., Niwa, H. & Nakao, M. Nuclear and chromatin reorganization in the MHC-Oct3/4 locus at developmental phases of embryonic stem cell differentiation. *Dev. Biol.* **298**, 354–367 (2006).
- Meshorer, E. & Misteli, T. Chromatin in pluripotent embryonic stem cells and differentiation. *Nat. Rev. Mol. Cell Biol.* **7**, 540–546 (2006).
- Constantinescu, D., Gray, H. L., Sammak, P. J., Schatten, G. P. & Csoka, A. B. Lamin A/C expression is a marker of mouse and human embryonic stem cell differentiation. *Stem Cells* **24**, 177–185 (2006).
- Mattout, A., Goldberg, M., Tzur, Y., Margalit, A. & Gruenbaum, Y. Specific and conserved sequences in D. melanogaster and C. elegans lamins and histone H2A mediate the attachment of lamins to chromosomes. *J. Cell Sci.* **120**, 77–85 (2007).
- Taniura, H., Glass, C. & Gerace, L. A chromatin binding site in the tail domain of nuclear lamins that interacts with core histones. *J. Cell Biol.* **131**, 33–44 (1995).
- Efroni, S. *et al.* Global transcription in pluripotent embryonic stem cells. *Cell Stem Cell* **2**, 437–447 (2008).
- Meshorer, E. *et al.* Hyperdynamic plasticity of chromatin proteins in pluripotent embryonic stem cells. *Dev. Cell* **10**, 105–116 (2006).
- Azuara, V. *et al.* Chromatin signatures of pluripotent cell lines. *Nat. Cell Biol.* **8**, 532–538 (2006).
- Bernstein, B. E. *et al.* A bivalent chromatin structure marks key developmental genes in embryonic stem cells. *Cell* **125**, 315–326 (2006).
- Pan, G. *et al.* Whole-genome analysis of histone H3 lysine 4 and lysine 27 methylation in human embryonic stem cells. *Cell Stem Cell* **1**, 299–312 (2007).
- Guenther, M. G., Levine, S. S., Boyer, L. A., Jaenisch, R. & Young, R. A. A chromatin landmark and transcription initiation at most promoters in human cells. *Cell* **130**, 77–88 (2007).
- Rasmussen, T. P. Developmentally-poised chromatin of embryonic stem cells. *Front. Biosci.* **13**, 1568–1577 (2008).
- Hawkins, R. D. *et al.* Distinct epigenomic landscapes of pluripotent and lineage-committed human cells. *Cell Stem Cell* **6**, 479–491 (2010).
- Wen, B., Wu, H., Shinkai, Y., Irizarry, R. A. & Feinberg, A. P. Large histone H3 lysine 9 dimethylated chromatin blocks distinguish differentiated from embryonic stem cells. *Nat. Genet.* **41**, 246–250 (2009).
- Misteli, T., Gunjan, A., Hock, R., Bustin, M. & Brown, D. T. Dynamic binding of histone H1 to chromatin in living cells. *Nature* **408**, 877–881 (2000).
- Krejci, J. *et al.* Genome-wide reduction in H3K9 acetylation during human embryonic stem cell differentiation. *J. Cell. Physiol.* **219**, 677–687 (2009).
- Catez, F., Brown, D. T., Misteli, T. & Bustin, M. Competition between histone H1 and HMGN proteins for chromatin binding sites. *EMBO Rep.* **3**, 760–766 (2002).
- Hezroni, H. *et al.* H3K9 histone acetylation predicts pluripotency and reprogramming capacity of ES cells. *Nucleus* **2**, 300–309 (2011).
- Hezroni, H., Sailaja, B. S. & Meshorer, E. Pluripotency-related, valproic acid (VPA)-induced genome-wide histone H3 lysine 9 (H3K9) acetylation patterns in embryonic stem cells. *J. Biol. Chem.* **286**, 35977–35988 (2011).
- Gaspar-Maia, A., Alajem, A., Meshorer, E. & Ramalho-Santos, M. Open chromatin in pluripotency and reprogramming. *Nat. Rev. Mol. Cell Biol.* **12**, 36–47 (2011).
- Hsieh, J., Nakashima, K., Kuwabara, T., Mejia, E. & Gage, F. H. Histone deacetylase inhibition-mediated neuronal differentiation of multipotent adult neural progenitor cells. *Proc. Natl Acad. Sci. USA* **101**, 16659–16664 (2004).
- Tachibana, M. *et al.* G9a histone methyltransferase plays a dominant role in euchromatic histone H3 lysine 9 methylation and is essential for early embryogenesis. *Genes Dev.* **16**, 1779–1791 (2002).
- Peters, A. H. *et al.* Loss of the Suv39h histone methyltransferases impairs mammalian heterochromatin and genome stability. *Cell* **107**, 323–337 (2001).
- Tachibana, M., Matsumura, Y., Fukuda, M., Kimura, H. & Shinkai, Y. G9a/GLP complexes independently mediate H3K9 and DNA methylation to silence transcription. *EMBO J.* **27**, 2681–2690 (2008).
- Feldman, N. *et al.* G9a-mediated irreversible epigenetic inactivation of Oct-3/4 during early embryogenesis. *Nat. Cell Biol.* **8**, 188–194 (2006).
- Chen, X. *et al.* Integration of external signaling pathways with the core transcriptional network in embryonic stem cells. *Cell* **133**, 1106–1117 (2008).
- Nishiyama, A. *et al.* Uncovering early response of gene regulatory networks in ESCs by systematic induction of transcription factors. *Cell Stem Cell* **5**, 420–433 (2009).
- Krouwels, I. M. *et al.* A glue for heterochromatin maintenance: stable SUV39H1 binding to heterochromatin is reinforced by the SET domain. *J. Cell Biol.* **170**, 537–549 (2005).
- Guttman, M. *et al.* Chromatin signature reveals over a thousand highly conserved large non-coding RNAs in mammals. *Nature* **458**, 223–227 (2009).
- Yokochi, T. *et al.* G9a selectively represses a class of late-replicating genes at the nuclear periphery. *Proc. Natl Acad. Sci. USA* **106**, 19363–19368 (2009).
- Bhattacharya, D., Talwar, S., Mazumder, A. & Shivashankar, G. V. Spatio-temporal plasticity in chromatin organization in mouse cell differentiation and during Drosophila embryogenesis. *Biophys. J.* **96**, 3832–3839 (2009).
- Meshorer, E. Chromatin in embryonic stem cell neuronal differentiation. *Histol. Histopathol.* **22**, 311–319 (2007).
- Pajerowski, J. D., Dahl, K. N., Zhong, F. L., Sammak, P. J. & Discher, D. E. Physical plasticity of the nucleus in stem cell differentiation. *Proc. Natl Acad. Sci. USA* **104**, 15619–15624 (2007).
- Phair, R. D. & Misteli, T. High mobility of proteins in the mammalian cell nucleus. *Nature* **404**, 604–609 (2000).
- Phair, R. D. *et al.* Global nature of dynamic protein-chromatin interactions in vivo: three-dimensional genome scanning and dynamic interaction networks of chromatin proteins. *Mol. Cell Biol.* **24**, 6393–6402 (2004).
- Sullivan, T. *et al.* Loss of A-type Lamin Expression Compromises Nuclear Envelope Integrity Leading to Muscular Dystrophy. *J. Cell Biol.* **147**, 913–920 (1999).
- Pannetier, M. & Feil, R. Epigenetic stability of embryonic stem cells and developmental potential. *Trends Biotechnol.* **25**, 556–562 (2007).
- Chen, R. Z., Pettersson, U., Beard, C., Jackson-Grusby, L. & Jaenisch, R. DNA hypomethylation leads to elevated mutation rates. *Nature* **395**, 89–93 (1998).
- Okano, M., Bell, D. W., Haber, D. A. & Li, E. DNA methyltransferases Dnmt3a and Dnmt3b are essential for de novo methylation and mammalian development. *Cell* **99**, 247–257 (1999).
- Tsumura, A. *et al.* Maintenance of self-renewal ability of mouse embryonic stem cells in the absence of DNA methyltransferases Dnmt1, Dnmt3a and Dnmt3b. *Genes Cells* **11**, 805–814 (2006).

41. Cedar, H. & Bergman, Y. Linking DNA methylation and histone modification: patterns and paradigms. *Nat. Rev. Genet.* **10**, 295–304 (2009).
42. Fan, Y. *et al.* Histone H1 depletion in mammals alters global chromatin structure but causes specific changes in gene regulation. *Cell* **123**, 1199–1212 (2005).
43. Woodcock, C. L., Skoultschi, A. I. & Fan, Y. Role of linker histone in chromatin structure and function: H1 stoichiometry and nucleosome repeat length. *Chromosome Res.* **14**, 17–25 (2006).
44. Ware, C. B. *et al.* Histone deacetylase inhibition elicits an evolutionarily conserved self-renewal program in embryonic stem cells. *Cell Stem Cell* **4**, 359–369 (2009).
45. Zhou, H. *et al.* Generation of induced pluripotent stem cells using recombinant proteins. *Cell Stem Cell* **4**, 381–384 (2009).
46. Shi, Y. *et al.* Induction of pluripotent stem cells from mouse embryonic fibroblasts by Oct4 and Klf4 with small-molecule compounds. *Cell Stem Cell* **3**, 568–574 (2008).
47. Alder, O. *et al.* Ring1B and Suv39h1 delineate distinct chromatin states at bivalent genes during early mouse lineage commitment. *Development* **137**, 2483–2492 (2010).
48. Melcer, S. & Meshorer, E. The silence of the LADs: dynamic genome-lamina interactions during ESC differentiation. *Cell Stem Cell* **6**, 495–497 (2010).
49. Melcer, S. & Meshorer, E. Chromatin plasticity in pluripotent cells. *Essays Biochem.* **48**, 245–262 (2010).
50. Reddy, K. L., Zullo, J. M., Bertolino, E. & Singh, H. Transcriptional repression mediated by repositioning of genes to the nuclear lamina. *Nature* **13**, 243–247 (2008).
51. Efroni, S., Melcer, S., Nissim-Rafinia, M. & Meshorer, E. Stem cells do play with dice: a statistical physics view of transcription. *Cell Cycle* **8**, 43–48 (2009).
52. Gaspar-Maia, A. *et al.* Chd1 regulates open chromatin and pluripotency of embryonic stem cells. *Nature* **460**, 863–868 (2009).
53. Ho, L. *et al.* An embryonic stem cell chromatin remodeling complex, esBAF, is an essential component of the core pluripotency transcriptional network. *Proc. Natl Acad. Sci. USA* **106**, 5187–5191 (2009).
54. Yan, Z. *et al.* BAF250B-associated SWI/SNF chromatin-remodeling complex is required to maintain undifferentiated mouse embryonic stem cells. *Stem Cells* **26**, 1155–1165 (2008).
55. Gilbert, N. *et al.* DNA methylation affects nuclear organization, histone modifications, and linker histone binding but not chromatin compaction. *J. Cell Biol.* **177**, 401–411 (2007).
56. Cheutin, T. *et al.* Maintenance of stable heterochromatin domains by dynamic HP1 binding. *Science* **299**, 721–725 (2003).
57. Gerlitz, G. *et al.* Migration cues induce chromatin alterations. *Traffic* **8**, 1521–1529 (2007).
58. Nissim-Rafinia, M. & Meshorer, E. Photobleaching assays (FRAP & FLIP) to measure chromatin protein dynamics in living embryonic stem cells. *J. Vis. Exp.* **52**, pii 2696 (2011).
59. Mills, K. I. & Ramsahoye, B. H. DNA methylation protocols. Overview. *Methods Mol. Biol.* **200**, 1–7 (2002).

Acknowledgements

We thank Andreas Nagy (Toronto), Rudolf Jaenisch (Cambridge, MA), En Li (Cambridge, MA), Masaki Okano (Kobe), Thomas Jenuwein (Freiburg) and Yoichi Shinkai (Kyoto) for kindly providing cells and reagents; Cecilia Ostlund and Howard Worman (New York) for kindly providing human LMNA plasmids, Adi Alajem (Jerusalem) for help with experiments, and Tom Misteli for critically reviewing the manuscript. E.M. is the Joseph H. and Belle R. Braun senior Lecturer in Life Sciences. This work was supported by the Safra Foundation, the Israel Science Foundation (ISF 215/07 and 943/09 to EM), the European Union (IRG-206872 and ERC-281781 to EM) and NIH Grant CA079057 (to A.S.). S.M. and H.H. are Edmond J. Safra fellows. The research was partly supported by the Center for Cancer Research, intramural program of the NCI, NIH.

Author contributions

S.M., H.H., E.R., M.N.R. and E.M. performed the experiments. S.M., H.H., E.R., M.N.R. and E.M. analysed the results. H.H. performed computational analyses. A.S. and C.L.S. provided reagents. S.M., E.R., M.B. and E.M. conceived the experiments. S.M. and E.M. wrote the manuscript.

Additional information

Supplementary Information accompanies this paper at <http://www.nature.com/naturecommunications>

Competing financial interests: The authors declare no competing financial interests.

Reprints and permission information is available online at <http://npg.nature.com/reprintsandpermissions/>

How to cite this article: Melcer, S. *et al.* Histone modifications and lamin A regulate chromatin protein dynamics in early embryonic stem cell differentiation. *Nat. Commun.* **3**:910 doi: 10.1038/ncomms1915 (2012).