

# Chromatin dynamics at the replication fork: there's more to life than histones

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Before each division, eukaryotic cells face the daunting task of completely and accurately replicating a heterogeneous, chromatinized genome and repackaging both resulting daughters. Because replication requires strand separation, interactions between the DNA and its many associated proteins — including histones — must be transiently broken to allow the passage of the replication fork. Here, we will discuss the disruption and re-establishment of chromatin structure during replication, and the consequences of these processes for epigenetic inheritance.

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**Current Opinion in Genetics & Development** 2013, **23**:140–146

This review comes from a themed issue on **Genome architecture and expression**

Edited by **Genevive Almouzni** and **Frederick Alt**

For a complete overview see the [Issue](#) and the [Editorial](#)

Available online 21st January 2013

0959-437X/\$ – see front matter, Published by Elsevier Ltd.

<http://dx.doi.org/10.1016/j.gde.2012.12.007>

## Introduction: the epigenetic challenge

In our understanding, the term epigenetics is used to describe the inheritance of patterns of gene expression that are not based on changes in DNA sequence [1]. Epigenetic processes need not rely on histones or post translational modifications [2], but they should fulfill three key requirements: first, to regulate gene expression; second, to be maintained through cell division; and third, to template their own duplication; thus, epigenetic factors must be heritable in the absence of ongoing inducing signals [3].

Chromatin is frequently associated with epigenetics; chromatin structure is determined by myriad histone modifications, histone variants, transcription factors, remodeling enzymes, and RNAs [4–8]. The primary repeating unit of chromatin is the nucleosome, which contains ~147 bp of DNA wrapped around an octameric core composed of two copies of each of the core histone proteins: H2A, H2B, H3, and H4 [9]. The core histones interact to form H2A–H2B and H3–H4 dimers; two H3–H4 dimers associate to form the (H3–H4)<sub>2</sub> tetramer, and

H2A–H2B dimers bind on either side of the tetramer to form the octamer. The large number of histone post-translational modifications found on histones within transcriptionally active or repressed chromatin [10,11], and the associated ‘histone code’ hypothesis [8], leads intuitively to the attractive possibility that histones may act as global carriers of regulatory information to control gene expression through generations. However, if histone modifications are to serve as epigenetic vectors they must surmount the challenge of DNA replication, in which chromatin is disassembled ahead of the replication fork and reassembled onto two daughter genomes.

Eukaryotic replisomes contain up to five distinct histone chaperones (extensively reviewed [12]), which together minimize the amount of non-nucleosomal DNA at the replication fork and ensure timely redeposition of nucleosomes on nascent DNA. After displacement, H3–H4 dimers and/or (H3–H4)<sub>2</sub> tetramers are passed to PCNA-associated CAF-1 [13,14] via FACT [15–17], Asf1 [18,19] or other chaperones [5] for deposition. Electron microscopy has suggested the disruption of 1–2 nucleosomes in front of the replication machinery [20], and we have shown that ongoing lagging-strand synthesis is intrinsically linked to the immediate redeposition of histones in *Saccharomyces cerevisiae* [21<sup>••</sup>], suggesting that the location of parental histones on the daughter strand to which they have segregated should closely correspond to their position within the parental genome. Consistent with this, a clever epitope switching method that allows parental and nascent histone H3 to be distinguished has shown that the parental histones are generally redeposited within ~400 bp of their prereplication position in *S. cerevisiae* [22<sup>•</sup>]. The reassembly of nucleosomes close to their parental locations renders theoretically feasible the inheritance of histone post-translational modifications within relatively discrete domains (H2A and H2B are deposited after H3 and H4 [23,24] and will not be discussed further).

Importantly, (H3–H4)<sub>2</sub> tetramers generally remain intact during replication fork passage [25<sup>•</sup>,26,27<sup>•</sup>] and appear to segregate randomly to the two daughter genomes [28]. Some tetramers, containing variants of H3 whose deposition is independent of S-phase [29], split [27] but the significance of this is presently unclear. It is notable that key histone chaperones [30,31] — including CAF-1 [32] — interact with two H3–H4 dimers most likely in the form of a (H3–H4)<sub>2</sub> tetramer [32]. The production of two daughter genomes necessitates that nucleosomes

containing parental (H3–H4)<sub>2</sub> cores are intermixed with an equal number containing newly synthesized H3–H4, devoid of parental modifications but carrying characteristic deposition-related marks [33,34]. *Trans*-acting maintenance modification complexes (see below), targeted by marks on parental histones, can subsequently modify newly synthesized histones to harmonize the chromatin modification state across domains in the two daughter genomes.

### Histone-mediated inheritance of chromatin domains

Histone modifications are clearly maintained in specific domains through the action of factors that target modification enzymes. Studies have shown that methylation of histone H3 lysine-9 (H3K9me) and lysine-27 (H3K27me) is catalyzed by methyltransferases that reside in complexes containing subunits that also bind methylated histone tails [35–38]. This provides a simple mechanism by which maintenance modification complexes can propagate a modification to nearby histones. The prototypic example of this behavior is provided by H3K9me3: this is bound by HP1, which recruits H3K9-specific methyltransferases and, by oligomerizing, can spread this modification across a domain [39]. Thus, provided that modifications recognized *in cis* can be propagated *in trans* to nearby nucleosomes, domains of modified chromatin can potentially be maintained via simple mechanisms that do not rely on signaling or sequence-specific DNA binding proteins, and that are robust in the face of twofold dilution during replication [40].

Recently, a series of elegant experiments by the Crabtree laboratory [41\*\*] has lent experimental support to the assertion that H3K9 methylation can be transmitted via truly epigenetic mechanisms. Through the small-molecule-induced recruitment of HP1 $\alpha$ , Hathaway and coworkers were able to induce H3K9me3 at the Oct4 locus in mouse embryonic stem cells (ES) and embryonic fibroblasts (MEFs); the ensuing methylation spanned ~10 kb, making it remarkably similar in size to endogenous H3K9me3 domains. Critically, in MEFs, which do not express ES cell pluripotency factors, the induced H3K9me3 domain was heritably transmitted through DNA replication and persisted through multiple cell divisions after removal of the HP1 $\alpha$  stimulus. Further *in silico* modeling demonstrated that a simple equilibrium between modification and replication-independent histone turnover or demethylation could give rise to stable, inherently bounded domains where spreading of the mark along the chromosome was opposed by removal of the modification [41,42] (see Figure 1a). According to the model proposed by the authors, steady state levels of H3K9me3 are the result of the opposing influences of histone methylation and turnover; this equilibrium can clearly be altered, which sits well with experiments in *Drosophila* that have shown that pericentromeric

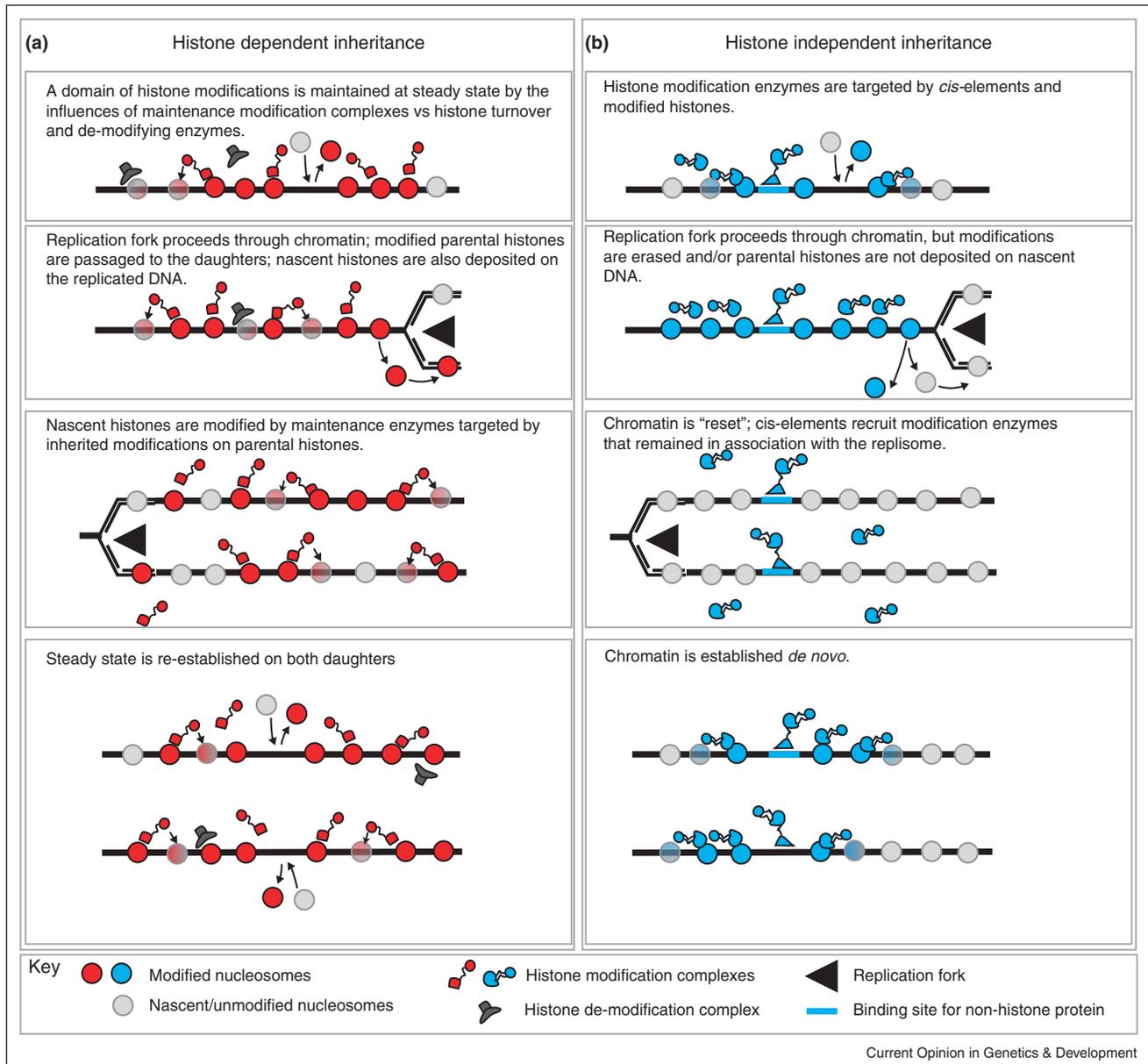
heterochromatin domains containing H3K9me3 are sensitive to the dosage of factors that either promote or oppose heterochromatin [43]. Consistent with this, targeting an activator to the ectopically methylated chromatin generated by Crabtree and colleagues was sufficient to disrupt the steady state and favor unmethylated H3K9 [41\*\*]. Earlier studies in human fibroblasts have described a similar scenario for the persistence of H3K27me3. Hansen *et al.* [36] showed that H3K27me3, which is catalyzed by the Polycomb Repressive Complex 2 (PRC2), is also bound by PRC2; and, using a heterologous reporter system, found that transient recruitment of PRC2 led to H3K27me3 and gene repression that persisted through cell divisions. Importantly, methylation of H3K27 appears to be important for the long-term persistence of the repressed state as targeting a catalytically inactive methyltransferase greatly diminished the long-term silencing of the reporter.

Euchromatin and heterochromatin are spatially distinct, and are replicated at different times in S-phase [44,45]. Recent work has indicated that the spatial organization of chromatin within the nucleus directly impacts the time at which it replicates, and that proximity to early-replicating or late-replicating domains [46] is by far the best predictor of when a given region of the genome will be replicated [47\*\*,48\*\*]. The broad division of the genome into early-replicating and late-replicating regions based on spatial proximity presumably provides a simple means to increase the robustness with which domains of common modifications can be inherited, as the binding of euchromatin-specific or heterochromatin-specific maintenance factors to the replisome can be both spatially and temporally controlled. Consistent with this hypothesis, microinjection experiments have shown that the assembly of transcriptionally competent chromatin is dependent upon the timing of the injection, with DNA injected early in S-phase being assembled into acetylated chromatin and expressed at higher levels [49,50]. Temporally separating the replication of active and repressed chromatin can therefore be sufficient to establish distinct chromatin types, even on naked DNA. Spatial segregation of chromatin domains likely adds a layer of redundancy, ensuring that chromatin domains are properly inherited by providing a high local concentration of certain modifying factors; replication in this context will likely ensure that the resulting chromatin on the daughters will be modified by physically associated factors, regardless of the fate of parental histones and their modifications.

### Inheritance mediated by nonhistone proteins

Although histones and their modifications are often described as epigenetic, in most cases there is in fact no direct evidence to support the assertion that modifications themselves are the key determinants of epigenetic processes. Indeed, several observations make it

Figure 1



Simplified schematic depiction of mechanisms by which chromatin domains may be maintained through DNA replication. (a) Histone-dependent inheritance: parental histones and modifications are retained on the daughter genomes following replication fork passage, but are diluted by nascent histones. Inheritance of modifications is mediated by maintenance enzymes targeted to inherited modifications. (b) Histone-independent inheritance: parental histones or modifications are not maintained; the chromatin state is re-established *de novo* by factors targeted to *cis*-acting elements in the DNA.

difficult to reconcile the generality of histones and epigenetics: first, modifications themselves appear to have little instructive capability and are often correlative rather than causative of basic processes such as gene transcription [4,51]; second, histone modifications such as acetylation and phosphorylation have very short half-lives [52]. Longer-lived modifications such as methylation may not necessarily persist through generations because the histones themselves could be replaced before the next cell cycle via replication-independent turnover [53,54].

On a global level, it has long been recognized that parental histones are passed to the daughter genomes during DNA replication, but whether such passing is consistent across the genome, and whether all histones maintain their modifications, often remain open questions. Recent experiments suggest that the assumption that all parental histone modifications are transferred to daughter genomes may be overly simplistic. Petruk *et al.*, asked whether H3K4me3, which is catalyzed by the trithorax protein Trx, or H3K27me3, catalyzed by the

polycomb group protein E(z), was associated with replicating DNA in *Drosophila* embryos [55<sup>••</sup>]. Surprisingly, the methylated histones in question were not detected during S-phase or found to be in proximity to the replication fork. Intriguingly, however, the methyltransferases Trx and E(z) were observed to colocalize with the replisome. These findings are consistent with other recent work that shows that the polycomb group complex PRC1 remains continuously associated with chromatin replicated *in vitro* [56,57]. Although one should not discount the possibilities that chromatin assembly in *Drosophila* embryos may be unusual, or that methylated histones may be present during S-phase in low quantities, these studies highlight important questions regarding the necessity of histones as mediators of epigenetic processes: propagation of histones and their modifications may not be required, provided that factors capable of establishing chromatin structures *de novo* are continually present (see Figure 1b).

The results of Hathaway *et al.* [41] and Petruk *et al.* [55] may represent two ends of a spectrum: if an appropriate balance between histone methylation and turnover is struck, then modifications can persist without establishment factors; however, if turnover/demethylation is dominant then establishment factors will be required. It is unclear why one mechanism should be favored over another, but clues may lie in the size and stability of the chromatin domain. Computational modeling has shown that a chromatin domain can persist if the propagation rate (i.e. methylation) is greater than the turnover rate [34]; a necessary consequence of this is a chromatin domain that may expand tens of kb. Indeed, in *Drosophila*, large domains of H3K27me3 spanning hundreds of kb are found in the bithorax complex region [58], which is a region of comparatively low histone turnover [59]. However, while such domains may be stable and easily persist through DNA replication, they may be too robust to allow rapid changes in gene expression or may simply be too large to be compatible with the intricate aspects of gene regulation mediated by fine-scale chromatin structure.

Evidently the association of chromatin modifying and remodeling factors with the replication fork is likely to aid the inheritance of chromatin structures through DNA replication. Petruk *et al.* suggest that Trx and E(z) may persist at the replication fork through interactions with single stranded DNA; HP1 $\alpha$  is also found at the replication fork in complex with the histone chaperone CAF-1 and the H3K9 methyltransferase SetDB1 [60] as are other remodeling factors such as SMARCD1 and histone deacetylases [61]. Furthermore, modifying complexes can travel with DNA polymerase  $\epsilon$  to progressively modify histones at the fork [62]. However, we should not discount the role of sequence-specific transcription factors, which may be retained through DNA replication

and are often fundamental to the establishment of chromatin states.

### Transcription factors and the limits of epigenetics

Recently, van Steensel and colleagues employed a DamID approach to map the localization of 53 broadly selected factors in *Drosophila*, identifying five distinct chromatin types that each contain a characteristic mixture of chromatin-associated factors [63]. Interestingly, many proteins are not confined to a single chromatin type, demonstrating that physiologically relevant chromatin structures are unlikely to be specified by the action of a single exclusive factor, but rather result from the combinatorial and/or redundant action of many activities. In addition, many regions of histone modification span only a few nucleosomes and are asymmetric [64]. Increased turnover (or restricted, hyperaccurate propagation by maintenance complexes) at these sites may be necessary to restrict the spread of the mark, but this also diminishes its ability to persist in the absence of the ongoing presence of inducing signals such as transcription factor binding, reducing the potential for epigenetic inheritance.

Similarly to nucleosomes, transcription factors must be removed from DNA before its replication. By analyzing Okazaki fragments in budding yeast, we have recently found that some transcription factors influence how Okazaki fragments are processed, which suggests that these factors are able to bind as rapidly as histones behind the replication fork [21]. Interestingly, the three proteins for which we observed rapid rebinding — Abf1, Reb1, and Rap1 — are key determinants of chromatin structure and help organize nucleosomes in gene promoters [65,66]. A recent study by the Lieb laboratory found that the occupancy times for even high-turnover Rap1 binding sites in *S. cerevisiae* are of the order of 30 min [67<sup>•</sup>] — almost as long as S-phase. Given that transcription factors, unlike histones, can be present at one local copy per two daughter genomes, asymmetric transcription factor binding behind the fork coupled with long residence could represent a means to establish distinct patterns of gene expression in daughter cells. In metazoan systems, rapid transcription factor reassociation with chromatin might play key regulatory roles at the promoters of tissue-specific genes during differentiation. For example, recent work has shown that the hematopoietic transcription factor GATA1 dissociates from DNA during S-phase, but rebinds to regulators of tissue-specific gene expression before M-phase. This ‘mitotic bookmarking’ [68] facilitates the rapid re-activation of these GATA1 targets in the subsequent G1, anchoring a complex of multiple transcription factors that do not reassociate in M after S-phase dissociation. The establishment of chromatin asymmetry by transcription factors, however, requires that protein binding be stable over time periods comparable with the

duration of S-phase, and is thus likely to be restricted to a subset of transcription factors with unusually long residence times on DNA.

## Conclusions and perspective

As the precise roles of histone modifications and sequence-specific DNA binding proteins in the perpetuation or alteration of gene expression state are further delineated, the answers to many important questions in the field of epigenetics may become clear. What is the fate during chromatin disruption and reassembly of the many histone modifications that are currently presumed to be directly inherited? For modifications such as H3K4me3 and H3K27me3 in *Drosophila* embryos, which are apparently erased during S-phase, it will be interesting to ascertain how this removal is accomplished and what, if any, is its biological purpose. Furthermore, the division of some cells — stem cells, for example — is accompanied by changes in expression state in a specific daughter. To what extent is this mediated by the asymmetric rebinding of transcription factors and/or modified histones behind the replication fork?

## Acknowledgements

We have highlighted only papers published in the past three years and apologize to colleagues whose work could not be cited due to space limitations. This work was supported by National Institute of Health Grant R01 GM102253 and an Alfred Bressler Scholars Endowment Award to I.W.; D.J.S. is a HHMI fellow of the Damon Runyon Cancer Research Foundation (DRG-#2046-10).

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