Higher-order chromatin structure is tightly linked to gene expression and therefore cell identity. In recent years, the chromatin landscape of pluripotent stem cells has become better characterized, and unique features at various architectural levels have been revealed. However, the mechanisms that govern establishment and maintenance of these topological characteristics and the temporal and functional relationships with transcriptional or epigenetic features are still areas of intense study. Here, we will discuss progress and limitations of our current understanding regarding how the 3D chromatin topology of pluripotent stem cells is established during somatic cell reprogramming and maintained during cell division. We will also discuss evidence and theories about the driving forces of topological reorganization and the functional links with key features and properties of pluripotent stem cell identity.

Principles and Mechanisms of 3D Chromatin Organization

The three-dimensional (3D) organization of the genome has been shown to have major implications for gene regulation and therefore cell identity. Over the last three decades, technological advancements in microscopy and genome-wide chromatin topology assays (Kempfer and Pombo, 2020) uncovered a hierarchical 3D genomic organization ranging from whole chromosome territories to (sub)megabase structures and fine-scale chromatin loops (Gibcus and Dekker, 2013; Yu and Ren, 2017). Individual chromosomes reside in discrete regions of the nucleus called chromosome territories (Cremer and Cremer, 2001). Within each chromosome, euchromatic and heterochromatic regions segregate into different subnuclear compartments named A and B, respectively (Lieberman-Aiden et al., 2009; Wang et al., 2016). The A compartments are gene rich, accessible, transcriptionally active, and largely occupy the nuclear interior, while B compartments are gene poor, inactive, and inaccessible regions, located at the nuclear periphery and overlapping with lamina-associated domains (LADs) (Guelen et al., 2008; Haarhuis et al., 2017). Due to their association with transcriptional activity, compartments are distinct across cell types (Dixon et al., 2012, 2015; Takebayashi et al., 2012). At (sub)megabase scale, we detect self-associating domains, known as topologically associating domains (TADs) (Dixon et al., 2016; Nora et al., 2017; Sexton et al., 2012) or contact domains (Tang et al., 2015). These are found in both A and B compartments and are demarcated by insulating boundaries, which prevent interactions with neighboring TADs. While TADs are largely conserved among cell types, the insulation strength of TAD boundaries and patterns of intra-TAD interactions can vary (Crane et al., 2015). Sub-TADs, or smaller insulated neighborhoods nested inside larger ones, have also been described (Phillips-Cremins et al., 2013; Rao et al., 2014). Although current nomenclature in the field is rather confusing (Dixon et al., 2016; Schoenfelder and Fraser, 2019), studies have provided strong evidence for the role of TADs and insulated neighborhoods in gene expression by restricting enhancer activity and specificity (Dowen et al., 2014; Lupiñez et al., 2015; Sun et al., 2019). Finally, at the finest level of architectural organization there are pairwise chromatin contacts between distal genomic loci, ranging from few kilobase pairs (kb) to several megabases (Mb) of linear distance. Different types of long-range genomic interactions have been identified based on the mediating protein factors and their potential impact on gene expression (Cheutin and Cavalli, 2019; Schoenfelder and Fraser, 2019; van Steensel and Furlong, 2019). These include active contacts between cis-regulatory elements (enhancer-promoter), Polycomb-mediated repressive/poising interactions (Cheutin and Cavalli, 2019; Denholtz et al., 2013; Eagen et al., 2017; Schoenfelder et al., 2015b), and CTCF/Cohesin structural loops (Handoko et al., 2011; Robson et al., 2019). Although structural loops are largely conserved among cell types, activating and repressive/poised contacts can be reorganized during cell fate transitions (Penalosa-Ruiz et al., 2019).

Two main forces are thought to work together and against each other to shape the layers of the 3D genome: loop extrusion and compartmental segregation, as extensively reviewed recently (Beagan and Phillips-Cremins, 2020; Nuebler et al., 2018; Rada-Iglesias et al., 2018; Rowley et al., 2017). In loop extrusion, the loaded Cohesin complex keeps sliding/extruding DNA through its ring until it encounters convergently oriented CTCF sites (Fudenberg et al., 2016; Sanborn et al., 2015). Loop extrusion is thus responsible for the formation of CTCF/Cohesin-mediated loops, TADs, and sub-TADs (Beagan and Phillips-Cremins, 2020), as well as the recently described “stripes” or “flames” (Fudenberg et al., 2016).
et al., 2016; Hsieh et al., 2020; Rao et al., 2014; Vian et al., 2018). On the other hand, A and B compartmentalization occurs through increased “self-attraction”/affinity among chromatin loci with similar transcriptional and chromatin states, such as homotypic histone modifications and transcription factors (TFs)/cofactors. Liquid/liquid-phase separation is one mechanism of chromatin self-organization/segregation (Palikyras and Papantonis, 2019), and it results in the formation of subnuclear droplet-like condensates or membraneless organelles, such as nucleoli or nuclear speckles. A large number of TFs (Boija et al., 2018; Chong et al., 2018), epigenetic modulators (BRD4, Polycomb Repressive Complex 2 [PRC2]) (Sabari et al., 2018; Tatavolsian et al., 2019), and transcriptional cofactors (Mediator, Pol II) (Cho et al., 2018; Zamudio et al., 2019) have been reported to form large nuclear condensates, which may mediate activating or repressive chromatin contacts in a loop-extrusion-independent manner. There is increasing evidence for the distinct but interconnected nature of these two guiding mechanisms, as knock-down or degradation of CTCF or Cohesin subunits disrupts TADs, stripes, and structural chromatin loops but has minimal effect on transcription and even reinforces compartmentalization (Haarhuis et al., 2017; Nora et al., 2017; Rao et al., 2017; Schwarzer et al., 2017; Wutz et al., 2017). Importantly, the expression levels and/or binding patterns of these critical factors (CTCF, Cohesin, TFs, cofactors, histone modifications) is often distinct across cell types and may thus be responsible for driving cell-type-specific chromatin architecture.

3D Chromatin (Re)Organization during Acquisition and Maintenance of Pluripotency

Pluripotent stem cells (PSCs) are endowed with the remarkable capacity to self-renew indefinitely in culture (self-renewal) while preserving their potential to differentiate into all somatic cell types upon proper stimulation in vitro or upon blastocyst injection in vivo (pluripotency) (Evans, 2011; Tabar and Studer, 2014). PSCs encompass both embryonic stem cells (ESCs), derived from early embryos, and induced PSCs (iPSCs), generated through somatic cell reprogramming (Takahashi and Yamanaka, 2006). The cellular properties of PSCs are supported by their unique epigenetic landscape, characterized by overall low levels of DNA methylation and heterochromatin marks and a prevalence of Polycomb-dependent repressive (H3K27me3) or bivalent (H3K4me3/H3K27me3) chromatin around lineage-related genes (extensively reviewed; see Di Giannattino and Apostolou, 2016; Harikumar and Mesehrer, 2015; Papp and Plath, 2013). This “open” and permissive chromatin environment becomes progressively restrained during differentiation as potency decreases (Sexton and Cavalli, 2015), and must be reset during reprogramming. The 3D genomic organization of PSCs is also characterized by distinct architectural features, such as the prevalence of A versus B compartments and the presence of PRC1/2-driven long-range interactions (Denholtz et al., 2013; Joshi et al., 2015; Schoenfelder et al., 2015b). Importantly, the 3D chromatin features of PSCs have largely been characterized in established, asynchronously cycling PSCs and often in comparison with other steady-state somatic cell types. In the following sections we will discuss recent work that sheds light on the dynamic nature of these features and particularly on how PSC-specific 3D chromatin landscape is established upon acquisition of pluripotency and propagated during cell division (Figure 1). We will particularly focus on findings from mouse naive PSCs that represent a pre-implantation pluripotent state, unless otherwise specified. Our goal is to highlight key factors and forces involved in building and maintaining PSC fate through reprogramming and self-renewal, respectively.

3D Chromatin Reorganization upon Acquisition of Pluripotency during Somatic Cell Reprogramming

Since the first derivation of iPSCs by Takahashi and Yamanaka (2006), cellular reprogramming has proved to be a powerful system to study mechanisms that dictate cell fate changes (Takahashi and Yamanaka, 2006). Ectopic expression of four TFs (OCT4, SOX2, KLF4, and c-MYC) can convert somatic cells into iPSCs with the genetic and epigenetic characteristics of stem cells, capable of self-renewing and differentiating into all germ layers in vivo. As somatic cells reprogram, they undergo cellular, transcriptional, metabolic, and epigenetic changes that result in erasure of the somatic program and establishment of pluripotency-defining properties (Apostolou and Hochedlinger, 2013; Brumbaugh et al., 2019; Papp and Plath, 2013). 3D chromatin architecture is also drastically reorganized during reprogramming, as shown by comparing various somatic cells with the resultant iPSCs, through profiling long-range interactions around specific loci (4C-seq or 5C) (Apostolou et al., 2013; Beagan et al., 2016; Denholtz et al., 2013; Wei et al., 2013) or in a genome-wide fashion by Hi-C (Krüger et al., 2016). Although most of these studies examined 3D chromatin architecture between start and endpoint fates, our review will focus on recent work that has begun to characterize the dynamic topological reorganization during reprogramming as well as underlying mechanisms and temporal associations with transcriptional and epigenetic changes.

3D Chromatin Architecture Is Drastically Reorganized during Somatic Cell Reprogramming along with Epigenetic and Transcriptional Changes

Compartment Switching

Reprogramming of multiple different mouse somatic tissue types (pre-B cells, macrophages, neural stem cells, mouse...
embryonic fibroblasts (MEFs)) resulted in iPSCs with 99% compartment similarity to one another and to naive ESCs (Krijger et al., 2016). This highlights the tight link between 3D organization and cell identity, and demonstrates that reprogramming promotes a rather faithful re-establishment of the pluripotency-associated chromatin architecture. To understand the kinetics of chromatin changes during reprogramming, a study from the Graf group characterized reprogramming intermediates at a genome-wide scale. They captured the dynamic topological reorganization that mouse B cells undergo during reprogramming to iPSCs using a highly efficient and synchronized system (Stadhouders et al., 2018). They discovered that compartmentalization changes occur gradually during reprogramming, with up to 20% of the genome switching compartments between any two intermediate time points. Surprisingly, many loci experienced multiple compartment switches throughout the time course, suggesting the emergence of transient, intermediate topologies and states. Whether these represent byproducts or critical steps of the reprogramming process remains unknown. Importantly, compartmentalization changes are associated with, and often occurred prior to, robust transcriptional changes. About 30%–40% of genes related to either B-cell or PSC identities underwent compartment switches during the course of reprogramming. Interestingly, pluripotency genes that were initially residing in A compartments were upregulated early during reprogramming, in parallel with the silencing of somatic genes, while those that switched from B to A (e.g., Dppa3) were activated at later stages. The delayed activation of genes embedded into repressive B compartments likely reflects the requirement for extensive epigenetic and topological remodeling prior to transcriptional upregulation. In support of this, integrative analysis with chromatin immunoprecipitation sequencing (ChIP-seq) data for the active histone mark H3K4me2 revealed that epigenetic remodeling occurs either concomitant with or prior to compartment switches, indicating that histone marks and/or the associated epigenetic modulators play active roles in large-scale architectural reorganization.

**TAD Reorganization**

In concordance with the reported conservation of TAD boundaries across cell types (Dixon et al., 2012, 2015), the vast majority of TAD boundaries remained constant throughout B-cell reprogramming. Genes found near the few lost or gained boundaries experienced variable effects on expression, with no clear correlation between gain/
loss and upregulation/downregulation. Despite minimal qualitative changes, 50%–80% of TADs experienced quantitative changes either in the insulation strength of their boundaries (frequency of interactions with neighboring TADs) or their self-connectivity (number of interactions within TAD). Interestingly, changes in TAD boundary insulation occurred early in reprogramming, broadly preceding epigenetic and compartmental reorganization. In agreement with reports in other cellular systems and during differentiation (Bonev et al., 2017; Crane et al., 2015; Van Bortle et al., 2014), insulation changes during reprogramming associated with transcriptional changes likely by facilitating or preventing enhancer-promoter communication. For example, both the Nanog and Sox2 loci underwent drastic TAD reorganization by losing or gaining a strong boundary, respectively. Weakening of the Nanog boundary enabled interaction with an upstream insulated enhancer, while the de novo boundary at the Sox2 locus generated a new insulated regulatory unit that included only Sox2 and its super-enhancer, potentially preventing aberrant/competing interactions. Importantly, the authors describe an inverse correlation between timing of transcriptional activation during reprogramming and degree of topological reorganization around the gene locus (Stadhouders et al., 2018). This suggests that the requirement for 3D chromatin reorganization may act as a limiting step for gene activation and, thus, as a reprogramming roadblock. Therefore, characterization of chromatin topology around critical genes may allow us to predict and overcome important architectural barriers during cell fate transition.

Rewiring of Chromatin Loops and Hubs

In parallel with this large-scale 3D chromatin reorganization, long-range chromatin contacts also change drastically. Early 4C-seq analyses revealed a large number of PSC-specific chromatin interactions around critical stem cell genes, such as Nanog, Oct4, and Dppa2/3 (Apostolou et al., 2013; Denholtz et al., 2013; Wei et al., 2013). They also captured a dynamic rewiring of chromatin loops at different reprogramming stages, which associated with and often preceded transcriptional activation of linked genes. In contrast with these studies, the recent Hi-C analysis of B-cell reprogramming captured only a small fraction of dynamic chromatin contacts (<20% of total loops), which surprisingly did not associate with B-cell- or PSC-specific genes or with transcriptional changes (Stadhouders et al., 2018). This is likely due to technical limitations and biases of the Hi-C approach and analysis, which preferentially detects structural CTCF/Cohesin-mediated loops rather than enhancer-promoter contacts (Bonev et al., 2017; Hsieh et al., 2020; Mumbach et al., 2016; Rao et al., 2014). To overcome this limitation, a recent study employed H3K27ac HiChIP (Mumbach et al., 2016, 2017), an approach that combines Hi-C with chromatin immuno-precipitation (Di Giammartino et al., 2019) to generate 3D reference maps of active enhancers and promoter connectomes in MEFs and PSCs. This approach revealed tens of thousands of cell-type-specific regulatory contacts, strongly associated with the transcriptional activity of linked genes, suggesting that drastic 3D enhancer rewiring occurs during reprogramming. Importantly, 4C-seq analysis independently validated the abrogation of enhancer-promoter contacts around silenced somatic genes (e.g., Ets1, Jag1) and de novo establishment around activated enhancers and genes (e.g., Tbx3, Mycn, Sox2). This could support an instructive role of enhancer-promoter looping for gene activation. However, due to the biased nature of Hi-ChIP, the authors cannot exclude that some differential Hi-ChIP loops may simply reflect changes in H3K27ac strength in preexisting loops rather than actual 3D reorganization. Future high-resolution assays, such as Capture-C or Micro-C (Hsieh et al., 2020; Krietenstein et al., 2020), will further dissect the degree and kinetics of 3D rewiring during reprogramming and its association to gene expression.

Intriguingly, H3K27ac HiChIP analysis also revealed a number of highly interacting enhancers, which the authors call 3D enhancer hubs (Di Giammartino et al., 2019). PSC-specific enhancer hubs, which partly overlap with previously described super-enhancers (Whyte et al., 2013), interact with multiple genes that are coordinately activated during reprogramming at a much higher probability compared with gene pairs of similar linear distance or within the same TAD. Indeed, CRISPR-mediated genetic or epigenetic perturbation of enhancer hubs resulted in downregulation of multiple hub-connected genes, supporting the role of these elements in gene coregulation. Importantly, meta-analysis and advanced modeling of the B-cell reprogramming Hi-C data revealed the assembly of similar 3D chromatin hubs that associate with robust transcriptional activation (Di Stefano et al., 2020). The presence and importance of 3D chromatin hubs has been described in many other cellular contexts (originally in the context of globin locus; de Laat and Grosveld, 2003), usually in association with super-enhancers, genes critical for cell identity, and/or high transcriptional activity (recently reviewed in Di Giammartino et al., 2020). Although most of these multiconnected hubs likely represent different, high-frequency, dynamic contacts within the cell population, simultaneous multi-way contacts in the same allele and cell have been detected by imaging or other specialized chromatin topology assays, albeit at low frequency. Specifically, genome architecture mapping (GAM) in mouse PSCs captured a large number of three-way contacts, especially at highly transcribed regions and super-enhancers (Beagrie et al., 2017), further supporting a functional link between degree of 3D connectivity and levels of transcriptional activity.
Taken together, these results document drastic 3D chromatin rewiring during reprogramming, occurring alongside or prior to transcriptional changes (Figure 2). These studies also identify 3D enhancer hubs as critical architectural centers of cell identity, whose reorganization during cell fate transitions might coordinate large-scale transcriptional changes. More studies are needed to understand how 3D hubs assemble and function: Do they represent dynamic and/or heterogeneous pairwise interactions or large topological assemblies? To what degree do they represent competition or synergy among hub-connected genes and enhancers? Are they formed (or disrupted) in an all-or-nothing or a stepwise manner during cell fate change and what are the driving forces? Dissecting the organizational principles of such architectural features might allow us to interfere with their assembly and thus to promote or prevent cell fate change.

**TFs Are Major Drivers of 3D Chromatin Reorganization during Reprogramming**

Somatic cell reprogramming is primarily driven by the activity of the ectopically expressed TFs, which bind the somatic genome and orchestrate many molecular changes, including transcriptional, post-transcriptional, and epigenetic remodeling, as summarized in recent reviews (Apostolou and Stadtfeld, 2018; Brumbaugh et al., 2019). The role of TFs in promoting/maintaining 3D chromatin topological features is increasingly appreciated and supported by computational and genomics analyses as well as imaging and functional experiments (recently reviewed in Di Giammartino et al., 2020; Kim and Shendure, 2019; Stadhouders et al., 2019). By combining this knowledge, we can envision various mechanisms of OKSM-driven architectural reorganization during reprogramming.

TF binding and subsequent recruitment of epigenetic modulators and chromatin remodelers can induce changes of chromatin accessibility and state, resulting in quantitative compartmental changes (e.g., strengthening or weakening) or compartment switches through self-association/segregation, as described in the introduction. Indeed, integration of Hi-C and ATAC-sequencing data during B-cell reprogramming showed that OCT4 and KLF4 binding (inferred from TF motifs around newly accessible sites) associated with early and stable B-to-A compartment switches (~5% of the genome) even prior to transcriptional changes (Stadhouders et al., 2018). The pioneering properties of reprogramming TFs (Soufi et al., 2012; 2015) that enable binding on nucleosomal DNA followed by remodeling may contribute to early compartmental changes. However, dense heterochromatic regions were shown to be refractory to early OKSM binding (Soufi et al., 2012), suggesting that additional factors and remodeling events are required for opening and compartment switching, resulting in delayed transcriptional activation of associated genes. TF-induced silencing of the somatic program, either by direct displacement of somatic TFs, recruitment of repressive complexes (e.g., HDACs) (Chronis et al., 2017),

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**Figure 2. Dynamic Changes in Chromatin Architecture during Somatic Cell Reprogramming**

Changes in local chromatin state and TAD insulation are the earliest architectural changes that have been reported during iPSC reprogramming. Compartment switching begins to occur in tandem with new loop formation and continued changes in TAD insulation. Finally, transcriptional changes often occur concurrently or after topological reorganization.
and/or activation of critical co-repressors such as SIN3A (Li et al., 2017), may contribute to the opposite A-to-B switches.

OKS have been shown to directly interact and co-occupy chromatin with multiple architectural protein factors, including subunits of Cohesin, Mediator, and PRC1/2 complexes (Apostolou et al., 2013; Denholtz et al., 2013; Kagey et al., 2010; Pardo et al., 2010; Schoenfelder et al., 2015a; van den Berg et al., 2010; Wang et al., 2006; Wei et al., 2013). TFs and recruited coactivators at enhancer or promoter regions may mediate “stalling” of the extruding Cohesin ring resulting in de novo loop formation. The overlapping binding patterns among OKS and Cohesin subunits support this notion (Dowen et al., 2014; Kagey et al., 2010). Furthermore, highly active genes and (super)enhancers have been reported to localize within strong boundaries or on Hi-C stripes/flames (Fudenberg et al., 2016; Hsieh et al., 2020; Rao et al., 2014; Vian et al., 2018). This suggests that OKSM binding may directly induce changes in TAD organization, such as new or stronger boundaries (Stadhouders et al., 2018), and create de novo contacts (Di Giammartino et al., 2019). Obviously, TAD reorganization during reprogramming may also be induced by altered CTCF occupancy, due to DNA methylation changes (Doerge et al., 2012; Gao et al., 2013; Polo et al., 2012; Sardina et al., 2018) around CTCF motifs (Bell and Felsenfeld, 2000; Hark et al., 2000; Maurano et al., 2015), or through its RNA-binding properties (Hansen et al., 2019; Saldaña-Meyer et al., 2019).

Reprogramming TFs are also involved at the finest level of topological reorganization as supported by their enriched binding at anchors of rewired chromatin contacts (Apostolou et al., 2013; Denholtz et al., 2013; Stadhouders et al., 2018). In addition, genetic perturbation experiments in established PSCs induced gain or abrogation of interactions around specific loci (de Wit et al., 2013; Novo et al., 2018; Wei et al., 2013). At a genome-wide scale, inducible depletion of KLF4 (and related KLF2 and KLF5) in PSCs resulted in a global loss of 3D enhancer connections with a preferential effect on 3D enhancer hubs (Di Giammartino et al., 2019). Similarly, CRISPR/Cas9-mediated mutation of a single KLF4 binding site within selected hubs was sufficient to induce partial disassembly of the hubs and downregulation of hub-connected genes (Di Giammartino et al., 2019). These experiments support an important role of KLF4 in the maintenance and function of 3D enhancer contacts. To also address its role in the de novo establishment/rewiring of long-range contacts, the authors performed an integrative analysis of KLF4 HiChIP, H3K27ac HiChIP, and transcriptional data at different reprogramming stages. This revealed that KLF4 binding associates both with silencing/disassembly of somatic enhancer-promoter contacts and with de novo formation of PSC enhancer-promoter contacts and hubs. KLF4 binding usually preceded or coincided with chromatin looping and enhancer/gene activation, but was not always sufficient to induce looping and/or transcription, highlighting the need for additional factors as described above. Another interesting finding was that KLF4 participates in two kinds of chromatin contacts: (1) activating enhancer-promoter loops and hubs, enriched in enhancer marks and transcriptional activators; and (2) repressive/pausing contacts enriched in H3K27me3 and PRC1/2 subunits, which were shown to promote long-range interactions among poised lineage-specific genes (de Wit et al., 2013; Denholtz et al., 2013; Pachano et al., 2019; Schoenfelder et al., 2015a; Schoenfelder and Fraser, 2019). Interestingly, many of these activating or repressive protein cofactors as well as the reprogramming TFs themselves are characterized by large intrinsically disordered regions (IDRs) (Xue et al., 2012), which were shown or presumed to mediate weak, low-affinity, multivalent interactions leading to the formation of subnuclear biomolecular condensates (Boija et al., 2018; Cho et al., 2018; Chong et al., 2018; Hnisz and Young, 2017; Sabari et al., 2018; Shrinivas et al., 2019). Such condensates may mediate not only pairwise chromatin interactions but also larger multiconnected active or repressive regulatory hubs, which are preferentially bound at high density by these protein factors (Beagrie et al., 2017; Di Giammartino et al., 2019; Hsieh et al., 2020; Rosencrance et al., 2020; Schmitt et al., 2016). Evidence that either supports or contradicts this notion has been extensively discussed in a recent review (Di Giammartino et al., 2020).

Future experiments will be critical to determine OKSM-dependent and OKSM-independent mechanisms of local and global topological reorganization during reprogramming. Another interesting direction would be to capture the dynamic formation of other PSC-characteristic structures, such as the reported extremely long-range chromatin contacts (Joshi et al., 2015; Novo et al., 2018) and the clustering of PRC-mediated bivalent genes (Denholtz et al., 2013; Schoenfelder et al., 2015b) and their significance in the establishment of pluripotency.

3D Chromatin Reorganization upon Maintenance of Pluripotency

Cell division poses a challenge for cell identity, since it is accompanied by transcriptional silencing (Gottesfeld and Forbes, 1997; Taylor, 1960), degradation or dissociation of most TFs and cofactors (Martínez-Balbás et al., 1995), widespread alterations in histone modifications (Wang and Higgins, 2013), and a dramatic reorganization of 3D genome architecture (Naumova et al., 2013) (Figure 3). PSCs are also characterized by a rapid cell cycle and short G1 phase compared with somatic cells (Savatier et al., 2002),
necessitating efficient mechanisms for faithful re-establishment of PSC-specific transcriptional and chromatin landscapes upon mitotic exit. In this chapter, we will discuss our current understanding of the 3D chromatin changes that cells undergo during mitosis and G1 entry, the association with transcriptional resetting, and presumed mechanisms that ensure timely and faithful heritability of PSC fate.

**Reorganization of Chromatin Architecture during Mitotic Entry and Exit**

The complex hierarchical and cell-type-specific 3D chromatin topology described in the previous chapters has been shown to drastically change during mitosis. Compartments, TADs, and loops collapse and chromatin acquires a cell-type-invariant conformation that consists of a compressed array of consecutive loops (Dekker, 2014; Naumova et al., 2013) (see Figure 1). This metaphase folding state is formed through a discrete series of steps: in prophase, formation of a consecutive loop array; in prometaphase, production of a helical, spiral-staircase-like structure with a condensin II scaffold; and in metaphase, further compaction into smaller clusters by condensin I and self-interactions (Batty and Gerlich, 2019; Gibcus et al., 2018). This mitosis-specific structure has been extensively characterized by microscopy (Earnshaw and Laemmli, 1983; Liang et al., 2015b; Marsden and Laemmli, 1979; Ou et al., 2017) and 3C-based methods (Dekker, 2014; Oomen and Dekker, 2017). The metaphase chromatin structure must unfold and reset during G1 entry, in concert with recruitment of TFs, cofactors, and transcriptional reactivation.

Several recent studies have begun characterizing the dynamic resetting of chromatin architecture upon mitotic exit in different cell types (Abramo et al., 2019; Kang et al., 2020; Naumova et al., 2013; Zhang et al., 2019), including mouse naive PSCs (Nagano et al., 2017; Pelham-Webb et al., 2020). Nagano et al. (2017) utilized single-cell Hi-C in ~3,000 asynchronous PSCs and performed cell cycle phasing to align them to a trajectory along the cell cycle. They observed a nonsynchronous manner of resetting upon mitotic exit, in which different levels of compartmental segregation continued throughout G1. TAD boundaries were also rapidly reset by G1, with boundary insulation reaching its maximum strength globally at G1 (Nagano et al., 2017), as previously observed (Naumova et al., 2013).

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| Cell Cycle Phase | Mitosis | G1 | S/G2 |
|-----------------|---------|----|------|
| Transcription   | Shut down | Partial resume | Full recovery |
| Transcription Factor (TF) binding | Only few (bookmarking TFs) | More TFs | All TFs |
| Cofactor binding | Dissociation | Recruitment | Full recovery |
| Epigenetic Marks | Variable loss, gain, or maintenance | Reestablishment | Full recovery |
| Chromatin conformation | Collapse to cell-type invariant structure | Reorganization | Full recovery |

Figure 3. Disruption of Cell-Type-Specifying Features Causes Temporary “Identity Crisis” during Mitosis

Summary of the molecular changes that challenge cell identity during cell division. Transcription largely shuts down and most TFs and cofactors are dissociated from mitotic chromatin, except for a select few mitotic bookmarking factors. Histone modifications are variably lost (acetylation), gained (phosphorylation), or selectively maintained (bookmarked). Finally, the 3D chromatin architecture collapses from a cell-type-specific structure to a cell-type-invariant metaphase state. These features then must all be faithfully reset during G1 in the daughter cells in order to maintain proper cell identity.
Nagano et al. (2017) also assessed chromatin loop dynamics during cell cycle, focusing on a small number (~2,000) of contacts with convergently oriented CTCF-bound motifs between 200 kb and 1 Mb apart. Although these few structural loops appeared stable throughout interphase, global analysis showed that cells in G1 had a lower percentage of short-range interactions than cells in S or G2, suggesting variability and asynchrony of contact reformation during mitotic exit (Nagano et al., 2017). However, the low resolution of Hi-C in this study and small number of cells captured in M-to-G1 transition make it difficult to dissect the kinetics of loop re-establishment and leave many questions unanswered. What types of chromatin contacts are established earlier or later during mitotic exit and what determines their differential kinetics? How does architectural reorganization associate with resetting of the PSC transcriptional program and identity?

More recent studies have shed light on these remaining questions in various mouse and human cell lines (Abramo et al., 2019; Kang et al., 2020; Zhang et al., 2019), including mouse PSCs (Pelham-Webb et al., 2020), by performing bulk Hi-C in various time points after mitotic arrest and release. Despite the technical differences, these reports also observed a rapid but heterogeneous resetting of genome architecture, with many similarities to the single-cell Hi-C study (Nagano et al., 2017). Compartments were detectable by ana-/telophase, but increasingly strengthened and expanded during G1, gaining longer-range A-A and B-B interactions (Abramo et al., 2019; Zhang et al., 2019). TAD boundaries were reset early, again reaching maximum insulation in early G1 (Pelham-Webb et al., 2020). Interestingly, Zhang et al. (2019) observed a “bottom-up” re-building of TADs in which small sub-TADs form first before merging into larger structures.

The increased resolution of Hi-C in these studies enabled better tracking of chromatin loop reformation kinetics, which appeared more heterogeneous than TADs or compartments (Abramo et al., 2019; Pelham-Webb et al., 2020; Zhang et al., 2019). Although many chromatin contacts were detected before G1 entry, they did not reach their half maximums until much later (Abramo et al., 2019). Short-range chromatin contacts reformed before longer ones (Pelham-Webb et al., 2020; Zhang et al., 2019) and Hi-C stripes gradually lengthened during G1 entry (Zhang et al., 2019), suggesting that CTCF/Cohesin-mediated loop extrusion (Fudenberg et al., 2016; Sanborn et al., 2015) is a major mechanism of chromatin reorganization upon mitotic exit. Interestingly, enhancer-promoter contacts were generally reset prior to structural loops in mouse G1E erythroblasts and PSCs, which could be a universal principle of architectural resetting or specific to stem and progenitor cells whose chromatin is more active and plastic. The slower kinetics of structural loops were partly due to the requirement for Cohesin reloading upon mitotic exit (Zhang et al., 2019), suggesting that the faster formation of enhancer-promoter loops might be driven by Cohesin-independent affinity forces. Importantly, chromatin contacts involving PSC super-enhancers were among the first to be detected (Pelham-Webb et al., 2020), highlighting their critical role in cell identity maintenance. These studies also reported transient conformational changes upon mitotic exit. Abramo et al. (2019) reported a temporary lack of loops during the transition from mitotic condensin-driven to interphase Cohesin-driven folding, while Zhang et al. (2019) identified a small group of transient contacts in telophase that were often disrupted by new boundaries or loops forming in G1 (Abramo et al., 2019; Zhang et al., 2019). The degree to which these transient contacts have any implications on genome function and cell identity remains to be shown.

Overall, these studies begin to shape a model of architectural resetting during mitotic exit (Figure 4). Boundary insulation is immediately reset, with some sub-TADs building up to larger TADs later on. Compartments are designated early (A or B), but gradually expand and gain strength. Loops follow more heterogeneous dynamics, with enhancer-promoter contacts reforming prior to structural ones and a few transiently established contacts. Future investigation could dissect the relative kinetics of 3D organization around cell-type-specific genes, enhancers, and hubs and their importance in cell fate determination.

Coordination of Transcriptional Reactivation with 3D Chromatin Reorganization

Alongside the chromatin reorganization during mitosis, transcription is largely halted (Prescott and Bender, 1962), as poised RNA polymerase II (Pol II) is cleared from DNA and blocked from re-initiation (Liang et al., 2015a). A recent study showed that Cohesin unloading from chromosome arms plays a role in the dissociation of Pol II during mitosis, and that suppression and reactivation of transcription is important for resetting the transcriptional landscape in G1 (Perea-Resa et al., 2020). Given the complex relationship between chromatin architecture and gene expression in steady state, how is the resetting of these processes coordinated during mitotic exit?

Recent studies have begun to characterize transcriptional reactivation during mitotic exit on a genome-wide scale, although few have explored its coordination with 3D chromatin reorganization. Here we focus on those that measured nascent transcriptional activity rather than steady-state RNA levels, by assays such as PRO-seq (precision nuclear run on sequencing) (Core et al., 2008; Kwak et al., 2013) and EU-RNA-seq (ethyl uridine-labeled RNA sequencing) (Palozola et al., 2017). These studies consistently showed that transcription of
genes and enhancers is dramatically downregulated during mitosis but rapidly recovered during G1 entry (Hsiung et al., 2016; Kang et al., 2020; Palozola et al., 2017; Pelham-Webb et al., 2020; Teves et al., 2018). Studies in somatic cells also saw a global, transient spike in transcription during mitotic exit, although groups of genes showed distinct kinetics (Hsiung et al., 2016; Palozola et al., 2017). A similar hypertranscriptional spike during G1 was independently detected in mouse PSCs (Pelham-Webb et al., 2020), indicating that this is a general feature of transcriptional resetting. However, the relative timing of gene reactivation differed among cell types, with cell identity genes being slowly reactivated in somatic cells, such as hepatocytes (Kang et al., 2020; Palozola et al., 2017), while PSC-associated genes and enhancers were among the first reactivated in stem cells (Pelham-Webb et al., 2020), indicating different requirements for propagation/resetting of the PSC program. Interestingly, these studies also described a significant correlation in the reactivation kinetics between enhancers and their target genes (assigned by linear proximity or looping) (Hsiung et al., 2016; Palozola et al., 2017; Pelham-Webb et al., 2020),

Figure 4. Resetting of Chromatin Architecture during Mitotic Exit
Schematic showing the resetting of different layers of chromatin architecture during the cell cycle. All of these features are disrupted in prometaphase (prometa), yet they reform with variable kinetics during mitotic exit. Compartments begin reorganizing during anaphase and telophase (ana/telo) and continue strengthening throughout S and G2. TADs are also rapidly reset, with smaller TADs in ana/telo building up to larger TADs in G1. TAD boundary insulation is strongest in G1 before weakening in S phase with the onset of DNA replication. Loops reform gradually and heterogeneously, with enhancer-promoter (Enh-Prom) contacts forming prior to CTCF/Cohesin-mediated structural loops. A few transient enhancer-promoter contacts are also observed during mitotic exit that are disrupted or lost later in the cell cycle.
suggesting that transcriptional reactivation is coordinated locally and/or through looping. However, there were many examples of early reactivated genes prior to enhancer activation and/or physical association (Hsiung et al., 2016; Pelham-Webb et al., 2020), arguing for both enhancer-dependent and -independent mechanisms.

To explore the coordination of transcriptional and architectural resetting during mitotic exit on a genome-wide scale, two recent studies integrated Hi-C with either Pol II ChIP-seq in erythroblasts (Zhang et al., 2019) or PRO-seq in PSCs (Pelham-Webb et al., 2020). Zhang et al. (2019) found that transcriptional kinetics were comparable within early- and late-forming TADs, indicating a minimal effect of these structures on gene reactivation. In contrast, Pelham-Webb et al., 2020 showed that rapid transcriptional reactivation correlated with faster compartmentalization and stronger boundary insulation in early G1, providing a link between transcriptional activity and large-scale architectural reorganization after cell division. Interestingly, although transcriptional activity positively correlated with loop strength over time, both studies reported no significant difference in the reactivation kinetics between genes involved in early or late regulatory loops (Pelham-Webb et al., 2020; Zhang et al., 2019). Another recent study using EU-RNA-seq and Hi-C in U2OS cells found that transcription is largelyreset prior to loop reformation, supporting that transcriptional reactivation is independent from long-range interactions during mitotic exit (Kang et al., 2020). Furthermore, in mouse PSCs, depletion of Pol II during mitosis and mitotic exit had no effect on large-scale chromatin (TAD, compartment) reformation (Jiang et al., 2020). Together, these studies indicate a surprisingly weak correlation between transcriptional and architectural re-setting. However, given that Hi-C enriches for structural over regulatory loops (Bonev et al., 2017; Di Giammartino et al., 2019; Nishiyama et al., 2013; Owens et al., 2019; Oomen et al., 2019; Sekiya et al., 2017; Shen et al., 2015). Although CTCF is retained during mitosis, Cohesin is lost in somatic cell lines (Burke et al., 2005; Hsieh et al., 2016; Kang et al., 2018; Liu et al., 2017; Mumbach et al., 2016), perhaps Hi-C is simply an insufficient method to explore this coordination. Indeed, Capture-C in erythroblasts successfully tracked enhancer-promoter contact reformation upon mitotic exit (Hsiung et al., 2016), and 4C-seq in PSCs detected a significantly faster resetting of chromatin contacts around early-activated enhancers (including super-enhancers) compared with all other regions (Pelham-Webb et al., 2020). In addition, despite the heterogeneity/asynchrony in loop dynamics around viewpoints, this analysis detected at least one early enhancer-promoter contact around every rapidly reactivated gene. Future experiments utilizing high-resolution genome-wide assays, such as Micro-C, Capture-C, or H3K27ac HiChIP, to detect re-establishment of enhancer-promoter contacts upon mitotic exit will be required to more concretely understand the interplay between enhancer-promoter activation and physical communication.

Role of Mitotic Bookmarking Factors in Architectural Resetting during G1 Entry

The heterogeneous kinetics of topological and transcriptional resetting during mitotic exit suggest the presence of distinct architectural factors or combinations of factors that drive reorganization. We have discussed these driving forces in the previous chapters, but it is important to note that many relevant factors are dramatically altered during mitosis and G1 entry, as the epigenetic landscape and TF binding patterns of mitotic cells are distinct from interphase cells. In addition to the global topological changes during mitosis, local chromatin features, including chromatin accessibility, nucleosome positioning, and histone modifications, are also perturbed (Kelly et al., 2010; Liang et al., 2015a; Wang and Higgins, 2013). Furthermore, TFs and cofactors are largely degraded or evicted from mitotic chromatin (Martínez-Balbás et al., 1995). What chromatin features are maintained during mitosis and are they important to instruct the resetting of stem cell fate?

Recent studies using advanced proteomics, imaging, and genomics technologies have revealed milder chromatin changes during mitosis than originally reported or presumed. Despite the global compaction of chromatin, DNA accessibility is largely retained, especially at promoters of active genes (Blythe and Wieschaus, 2016; Hsiung et al., 2015; Teves et al., 2016; Xu et al., 2017). Although histone tails are globally hypomethylated in mitosis, there is widespread retention of acetylation at certain residues (Behera et al., 2019; Hsiung et al., 2016; Javasky et al., 2018; Liu et al., 2017). This selective, partial maintenance of local interphase structure in mitosis is consistent with the concept of mitotic bookmarking (Festuccia et al., 2017; Kadauke and Blobel, 2013; Palozola et al., 2019). Recent work has explored mitotic bookmarking in mouse PSCs, identifying master regulator TFs such as ESRRB (Festuccia et al., 2016), SOX2 (Deluz et al., 2016; Liu et al., 2017; Teves et al., 2016), OCT4 (Deluz et al., 2016; Liu et al., 2017), and KLF4 (Liu et al., 2017), retained at selected genomic regions. Other mitotically retained factors and marks in PSCs include CTCF (Owens et al., 2019), TBP (Teves et al., 2018), and H3K27ac (Liu et al., 2017).

The potential bookmarking role of CTCF is particularly interesting due to its critical role in loop extrusion together with Cohesin. Owens et al. (2019) showed that while CTCF was retained during mitosis in mouse PSCs, it was lost in somatic cell lines (NIH3T3 and C2C12), which could explain incongruent results from prior studies (Burke et al., 2005; Oomen et al., 2019; Sekiya et al., 2017; Shen et al., 2015). Although CTCF is retained during mitosis, Cohesin is lost from all interphase sites (Nishiyama et al., 2013; Owens et al., 2019), arguing that CTCF bookmarking alone is likely insufficient to drive loop reformation. In support of this,
while CTCF was either mitotically retained or rapidly re-associated with chromatin upon mitotic exit in G1E erythroblasts, loop recovery kinetics correlated instead with the timing of Cohesin re-recruitment (Zhang et al., 2019). Mitotic retention of CTCF may, however, promote faster boundary resetting by sustaining local chromatin accessibility and nucleosome positioning (Owens et al., 2019) at critical sites, such as TAD boundaries (Hsiieh et al., 2020). On the other hand, mitotic loss of CTCF may allow formation of transient contacts during telophase (Zhang et al., 2019) or G1, resulting in aberrant gene activation.

Additional factors involved in extrusion-independent 3D chromatin organization (e.g., phase separation) and transcriptional regulation have also been identified as bookmarking factors in mouse PSCs. For example, TBP, a component of the RNA Pol II transcription complex, was widely retained at active promoters and its degradation during mitosis abrogated transcriptional reactivation globally (Teves et al., 2018). ESRRB maintained interphase nucleosome positioning during mitosis at bookmarked sites (Festuccia et al., 2019), which are enriched for stem cell identity genes (Festuccia et al., 2016). Degradation of either SOX2 (Deluz et al., 2016) or OCT4 (Liu et al., 2017) during M-to-G1 affected stemness, highlighting their importance during this window. Whether mitotic retention of any of these factors is involved in faster resetting of chromatin contacts and regulatory hubs remains to be tested.

Mitotic bookmarking by H3K27ac, which preferentially marks enhancers and genes associated with cell identity (Hsiung et al., 2016; Liu et al., 2017), was positively correlated with robust post-mitotic transcriptional reactivation in multiple cell types, including PSCs (Behera et al., 2019; Hsiung et al., 2016; Kang et al., 2020; Pelham-Webb et al., 2020), and its depletion during mitosis resulted in aberrant transcription in G1 (Kang et al., 2020; Pelham-Webb et al., 2020). Interestingly, post-mitotic recovery rate of enhancer-promoter contacts in erythroblasts positively correlated with levels of active histone marks, including H3K27ac (Zhang et al., 2019). Mitotic retention of H3K27ac was also identified as the best predictive mark for faster loop reformation and stronger TAD boundary insulation in PSCs exiting mitosis (Pelham-Webb et al., 2020). Together, these results support the role of this bookmark in faster transcriptional and architectural resetting of PSC identity after cell division (Figure 5), although further experimental evidence is required to dissect its function and underlying mechanisms. One possibility is that mitotic retention of H3K27ac promotes faster assembly of A compartments and active hubs through increased self-attraction with other homotypic chromatin regions. Alternatively, retention of H3K27ac may enable rapid recruitment of TFs/cofactors and reassembly of multiprotein complexes and condensates. One such candidate cofactor could be the chromatin reader BRD4, which plays critical roles in transcriptional activation and chromatin architecture in interphase (reviewed in Devaiah et al., 2016). Although BRD4 is partly retained on mitotic chromatin (Dey et al., 2009), its inhibition by JQ1 during mitosis did not alter transcriptional reactivation in G1 (Behera et al., 2019). This argues against the direct bookmarking function of BRD4 and suggests instead that mitotically retained histone acetylation marks, such as H3K27ac, are sufficient for rapid recruitment of BRD4 and proper genome activation during mitotic exit (Mochizuki et al., 2008; Yang et al., 2008). Future experiments depleting H3K27ac or other candidate cofactors during mitosis and mitotic exit will be required to validate this model.

Together, these results indicate that mitotic bookmarking factors contribute to a faster and more faithful molecular resetting of cell identity in daughter cells. It is plausible that the extent, nature, and significance of bookmarking in epigenetic inheritance of cell identity varies among cell types, depending on their cell cycle properties and vulnerability to cell fate change. Mitotic bookmarking by various histone marks and protein factors is quite widespread in PSCs (Liu et al., 2017; Teves et al., 2016), although direct side-by-side comparison with self-renewing progenitors or terminally differentiated cells is still missing. Future investigation will be required to determine the extent to which this distinct mitotic landscape of PSCs relates to their characteristic cell cycle and/or increased epigenetic plasticity and how it contributes to their ability to self-renew while maintaining pluripotency.

**M-to-G1 as a Window for Cell Fate Change**

The balance between self-renewal and pluripotency is especially important during mitotic exit, as G1 is the critical window for PSCs to “decide” either to self-renew or respond to differentiation cues toward defined lineages (Boward et al., 2016; Dalton and Coverdell, 2015; Soufi and Dalton, 2016). Although the concept of G1 as a window for cell fate change has been around for decades (Mummery et al., 1987; Pierce et al., 1984; Wells, 1982), only recently it was shown, for both human and mouse PSCs, that G1-sorted cells respond more rapidly to differentiation cues than S- or G2-sorted cells, which do not respond until the next G1 phase (Calder et al., 2013; Coronado et al., 2013; Pauklin and Vallier, 2013; Sela et al., 2012). Studies have also reported “noisy” expression of developmental genes during G1 (Asenjo et al., 2020; Singh et al., 2013), which could represent a temporal priming step that enables PSCs to tip the balance from self-renewal to lineage specification. This finding was independently validated using PRO-seq analysis in mouse PSCs, which revealed a weak but consistent activation of various lineage-related genes and enhancers during early and late G1.
What is unique about the M-to-G1 phase that primes PSCs to respond to differentiation cues? And what factors are responsible for closing this window after G1 to ensure self-renewal?

Recent studies have attempted to address this question, supporting different mechanistic models for transient activation of developmental genes during G1. Studies in human PSCs have implicated signaling molecules (Smads, Cyclins), which naturally fluctuate during cell cycle, concluding that varying levels of these factors allow for differentiation to endoderm in early G1 and neuroectoderm in late G1 (Pauklin et al., 2016; Pauklin and Vallier, 2013). Others consider that transient relaxation of repressive mechanisms during M-to-G1, such as the recently described cell-cycle-dependent fluctuations of PRC2 subunits (Asenjo et al., 2020), could be sufficient to allow for gene de-repression. In agreement, promoters of transiently G1-activated genes are predominantly marked by PRC2 and bivalent histone marks (H3K4me3/H3K27me3) as well as transiently activated during G1. This could be caused by fluctuation of PRC2 components over the cell cycle, slower recruitment of other repressive factors, and/or transient miswiring of chromatin loops.

Figure 5. Role of Mitotic Bookmarking in Transcriptional and Architectural Resetting during G1 Entry

Model describing the role of mitotic bookmarking TFs and histone marks in the rapid resetting of pluripotency genes and the transient activation of lineage-specific genes. In PSCs, bookmarking factors (blue circles) are preferentially retained at stem cell-specific genes and enhancers during mitosis. This allows for fast recruitment of cofactors and transcriptional machinery, resetting of enhancer-promoter contacts, and rapid transcriptional reactivation in G1 (arrows). Non-bookmarking TFs (green circles) must be recruited back to the chromatin during G1, resulting in a slower resetting of these sites. Lineage-specifying genes, often marked by PRC2 and bivalent histone marks (H3K4me3/H3K27me3) (pink circles), are transiently activated during G1. This could be caused by fluctuation of PRC2 components over the cell cycle, slower recruitment of other repressive factors (maroon ovals), and/or transient miswiring of chromatin loops.
and transient gene activation and re-repression. For example, late formation of a TAD boundary could allow for temporary enhancer-promoter “miswiring”, as observed during telophase in erythroblasts (Zhang et al., 2019). Alternatively, slower re-establishment of repressive loops (e.g., PRC1/2-mediated contacts) could enable transient de-repression of poised developmental genes. Both categories of temporarily mis-wired or late-established contacts have been detected by 4C-seq around the transiently activated, lineage-specific genes Gata6 and Neurod1 (Pelham-Webb et al., 2020). Interestingly, this study also captured a strong contact between the Gata6 promoter and an endoderm enhancer, which was pre-established/ maintained in mitosis (Pelham-Webb et al., 2020). This suggests that developmentally relevant chromatin structures may be already established in PSCs, as shown before (Cruz-Molina et al., 2017; Rubin et al., 2017; Schoenfelder et al., 2015a), but also selectively maintained during mitosis to enable rapid G1 activation and differentiation upon proper stimulation. The degree to which such permissive and “hard-wired” chromatin loops exist on a genome-wide scale and their functional role for transcriptional priming toward successful differentiation remains to be tested. Follow-up experiments in single cells will be critical to determine whether transient activation occurs in all or a subset of cells during G1 and the functional impact on lineage preference.

CONCLUSIONS AND FUTURE PERSPECTIVES

PSCs are considered a powerful system for disease modeling and regenerative medicine. Their unique cell identity is determined and maintained by a specific gene expression program, unique epigenetic landscape, and characteristic 3D genomic organization. Delineating the complex interplay among these regulatory layers is crucial for understanding and modulating PSC fate and harnessing their full biomedical potential. As discussed above, significant progress has been made by characterizing the dynamic changes that result in the (re)establishment of PSC identity starting either from a defined somatic identity (reprogramming) or a mitotic state (during self-renewal). A deeper understanding of these processes by constructing high-resolution 4D molecular roadmaps that integrate multiple -omics analyses will help address the following important questions: what are the essential transcriptional and architectural milestones and bottlenecks toward establishment of PSC identity? Are there common or unique regulatory hierarchies associated with each process (e.g., different order of establishment of PSC-associated features) and what are the critical driving forces? Can we identify specific vulnerabilities that may enable modulation of the self-renewal versus differentiation balance and a controlled pluripotency exit toward new fates?

A fundamental and actively debated question regarding cell fate control is the functional interplay between 3D genome architecture and transcriptional regulation (Beagan and Phillips-Cremins, 2020; Gibcus and Dekker, 2013; Gorkin et al., 2014; Robson et al., 2019; van Steensel and Furlong, 2019). The integrative reprogramming studies described above demonstrated that architectural changes (compartment switches, TAD insulation changes, and loop reorganization) often precede or coincide with transcriptional changes. Although this could support a cause-and-effect relationship, additional experimental evidence is required. In vivo imaging of chromatin reorganization, by advanced CRISPR-based approaches such as LiveFISH (Wang et al., 2019) or CLING (Maass et al., 2018), will enable tracking of long-range chromatin contacts and their association to gene activation. More importantly, breaking or engineering the chromatin loops/hubs predicted to be critical for PSC identity with technologies such as CRIPSR-GO (Wang et al., 2018) or LADL (Kim et al., 2019) will directly test their functional relevance. Previous studies in different systems have supported an instructive role of 3D chromatin architecture on gene regulation, showing that topological reorganization (altered insulation and/or looping) caused transcriptional changes of linked or nearby genes (Deng et al., 2012; Lupiániz et al., 2015; Schoenfelder et al., 2015a). Other studies suggest a permissive role, where preexisting chromatin loops enabled gene activation only upon recruitment of protein factors (Cruz-Molina et al., 2017; Ghavi-Helm et al., 2014; Jin et al., 2013; Melo et al., 2013; Rubin et al., 2017; Schoenfelder et al., 2015a). Finally, there is evidence for a reciprocal interplay and positive feedback between transcription and topological organization (van Steensel and Furlong, 2019). For example, transcriptional inhibition was shown to alter enhancer-promoter contacts (Hsieh et al., 2020) and to weaken CTCF-mediated loops that require its RNA-binding domain (Saldaña-Meyer et al., 2019), while genomic insertion of a transcription start site (TSS) was sufficient to induce local chromatin folding (Zhang et al., 2020). Moreover, RNA per se has been extensively documented to shape various subnuclear compartments and chromatin interactions (Quinodoz et al., 2020; Rinn and Guttman, 2014). Importantly, in most studies, perturbation of candidate factors and features happened in asynchronous cells and for variable time periods, often over many cell divisions. Therefore, it is hard to uncouple direct from indirect effects as well as their role in maintenance versus establishment of a molecular state (e.g., gene activation or loop formation). We argue that systematic interrogation of candidate factors specifically during mitosis or M-to-G1 may help provide more
definitive answers on the role of these regulators. To this direction, advanced technologies that enable temporal and reversible protein degradation (Kadauke et al., 2012; Nabet et al., 2018) or genetic/epigenetic engineering combined with reporters that enable prospective isolation or tracking of cells at defined cell cycle stages (Sakae-Sawano et al., 2008; Zerjatke et al., 2017) will be critical. Finally, an important consideration when performing or interpreting such studies is the degree of cell-type specificity of the molecular principles that are uncovered. Therefore, systematic comparison of multiple cell types or dynamic cell fate transitions are extremely valuable for dissecting universal and cell-type-specific mechanisms.

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