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Citation for published version:

Digital Object Identifier (DOI):
10.1101/gr.212308.116

Link:
Link to publication record in Edinburgh Research Explorer

Document Version:
Peer reviewed version

Published In:
Genome Research

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Constrained release of lamina-associated enhancers and genes from the nuclear envelope during T-cell activation facilitates their association in chromosome compartments

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Abstract

The 3D organization of the genome changes concomitantly with expression changes during hematopoiesis and immune activation. Studies have focused either on lamina-associated domains (LADs) or on topologically-associated domains (TADs), defined by preferential local chromatin interactions, and chromosome compartments, defined as higher-order interactions between TADs sharing functionally similar states. However, few studies have investigated how these affect one another. To address this, we mapped LADs using Lamin B1-DamID during Jurkat T-cell activation, finding significant genome re-organization at the nuclear periphery dominated by release of loci frequently important for T-cell function. To assess how these changes at the nuclear periphery influence wider genome organization, our DamID datasets were contrasted with TADs and compartments. Features of specific repositioning events were then tested by fluorescence in situ hybridization during T-cell activation. First, considerable overlap between TADs and LADs was observed with the TAD repositioning as a unit. Second, A1 and A2 sub-compartments are segregated in 3D space through differences in proximity to LADs along chromosomes. Third, genes and a putative enhancer in LADs that were released from the periphery during T-cell activation became preferentially associated with A2 sub-compartments and were constrained to the relative proximity of the lamina. Thus, lamina-associations influence internal nuclear organization and changes in LADs during T-cell activation may provide an important additional mode of gene regulation.
Introduction

The genome is organized within the three-dimensional interphase nucleus. Regulatory elements influence target genes hundreds of kbp to many Mbp distal by moving proximal through looping in three-dimensional space (Sanyal et al. 2012; Shen et al. 2012; Arner et al. 2015; Schoenfelder et al. 2015). Chromosomes are also organized along their length into discrete regions displaying preferential internal interactions detected by chromosome conformation capture (Hi-C) approaches, termed topologically-associated domains (TADs) or contact domains (CDs) depending on the algorithm used to calculate them (Dixon et al. 2012; Hou et al. 2012; Nora et al. 2012; Sexton et al. 2012). As TADs and CDs likely represent orthogonal measures of the same structural units, we refer to them as TADs unless specifically referring to published datasets employing the CD approach. Topological constraints imposed by TADs favor looping interactions to cis-regulatory element:gene pairings within the TAD (Lupianez et al. 2015; Franke et al. 2016; Hnisz et al. 2016). However, a higher level of genome organization also occurs where multiple TADs many Mbp distal on one chromosome or on different chromosomes can be adjacent and share a functional state (Lieberman-Aiden et al. 2009). Such associations form compartments, which segregate into two classes; A, transcriptionally active, and B, transcriptionally repressed (Lieberman-Aiden et al. 2009). While TAD composition is largely invariant, compartment composition changes during differentiation and between cell types correlating with gene activity (Dixon et al. 2015; Fraser et al. 2015); however, how compartment interactions are controlled during differentiation or other cellular transitions is largely unexplored.

The nuclear periphery in contrast contributes radial genome organization. Genome regions proximal to the lamin intermediate filament polymer lining the inner nuclear membrane are called lamina-associated domains (LADs) (Guelen et al. 2008). However, while implicated in genome organization (Mewborn et al. 2010; Mattout et al. 2011; Solovei et al. 2013), it remains unclear if lamins themselves are absolutely required for tethering LADs to the periphery (Amendola and van Steensel 2015; Zheng et al. 2015). The nuclear periphery is generally repressive with genes in
LADs tending to contain silencing marks (Guelen et al. 2008). Constitutive LADs are defined as unchanging, but the number defined as facultative, changing with the cell cycle or between cell types, is constantly increasing (Peric-Hupkes et al. 2010; Kind et al. 2013; Meuleman et al. 2013; Chandra et al. 2015; Robson et al. 2016). The importance of facultative LADs is underscored by the fact that their disruption has profound negative consequences for tissue differentiation (Robson et al. 2016). Tellingly, such facultative LADs move both to and from the periphery during differentiation and often contain genes requiring fine-tuned regulation (Peric-Hupkes et al. 2010; Robson et al. 2016).

Though LADs correlate with repressed compartments and TADs (Dixon et al. 2012; Rao et al. 2014; Fraser et al. 2015), few studies have investigated how alterations to one can influence the other. Moreover, this correlation is imperfect and is insufficient, for example, to predict TADs from LADs or vice versa. Several questions come to mind. Does a TAD as a functional unit move between the periphery and interior and thus perhaps encompass a facultative LAD? Can a new LAD that forms during differentiation influence the organization of TADs/compartments or vice-versa?

Facultative LAD repositioning tends to associate with repression or activation of genes within them. In neurogenesis and myogenesis 5-15% of genes move from the nuclear periphery to the interior while another 5-10% move from the interior to the periphery (Peric-Hupkes et al. 2010; Robson et al. 2016). Though lacking LAD datasets, lymphocyte-specific gene-positioning changes are reported such as IGH locus release from the periphery upon induction of V-D-J recombination (Kosak et al. 2002) and MAF locus repositioning to the periphery with its repression (Hewitt et al. 2004). Contrasting differentiation, lymphocyte activation is much more rapid and dynamic and involves significant genome restructuring (Drings and Sonnemann 1974). Compacted peripheral chromatin dissipates upon activation concomitant with large-scale gene activation (Pompidou et al. 1984). Our previous finding of changes in nuclear membrane protein composition during
lymphocyte activation (Korfali et al. 2010) led us to investigate whether there are also specifically-regulated gene exchanges at the nuclear envelope.

**Results**

**Mapping gene expression and repositioning changes during T-cell activation**

The extensive electron-dense peripheral heterochromatin of resting T-cells dissipates during activation concomitantly with the induction of immunogenic genes (Hirschhorn et al. 1971; Pompidou et al. 1984; Manteifel et al. 1992; Rawlings et al. 2011). However, it is unclear whether this reorganization is whole-scale or there is also more specific reorganization for immune activation. We predicted that loss of peripheral heterochromatin would correlate with release of T-cell activation-associated genes from the periphery. Accordingly, we investigated coordinated gene expression and genome organization changes during T-cell activation using microarrays and DamID (Fig. 1). Jurkat cells were incubated with Raji B-cells that had been pre-conjugated with staphylococcal enterotoxin E (SEE) resulting in the formation of antigen-independent immunological synapses (Fig. 1A) (Gonzalez-Granado et al. 2014). This yielded >95% activation of Jurkat cells by fluorescence activated cell sorting (FACS) (Fig. 1B). To determine associated gene expression changes, RNA was extracted at 0, 8, 24 and 48 h post-SEE stimulation and analyzed by microarray. Upon activation, 1,111 genes were upregulated at least 1.4-fold at any of the three time points and these were significantly enriched in Gene Ontology (GO)-terms positively supporting T-cell activation and early effector function (Figure 1C, Supplemental Fig. S1). Similarly, 1,016 genes were repressed at least 1.4-fold that were enriched in GO-terms inhibiting mitosis and cell division, presumably permitting accelerated proliferation of activated cells.

The same cell populations were subjected to lamin B1-DamID to globally map changes in genome-nuclear envelope associations. A GFP reporter confirmed efficient lentiviral transduction of Jurkat T-cells (Fig. 1D), but DamID signal was not observed until ~48 h post-transduction (Fig. 1E).
Hence, as no carry over DamID signal from the resting state would be observed, Jurkat T-cells were transduced with DamID constructs 24 h before antigen presentation so that measured changes reflect genome organization between 24 and 48 h into activation (Fig. 1A). Bacterial Deoxyadenosine methylase (Dam)-lamin B1 fusions were expressed to uniquely methylate DNA proximal to the lamin polymer (Vogel et al. 2007). To control for local variation in chromatin accessibility, parallel experiments expressed Dam methylase alone (Dam-only). The methylated DNA was then isolated and identified by sequencing.

Log$_2$(Lamin B1 Dam/Dam-only) ratios were generated and used to identify LADs in resting and activated Jurkat T-cells based on a circular binary segmentation algorithm slightly modified from that previously described (Harr et al. 2015; see methods). LADs were mapped onto the hg19 assembly because hg19 was used for earlier datasets used for comparison, but all regions examined by fluorescence in situ hybridization (FISH; see below) were confirmed to be unaltered in the newer hg38 assembly. Interestingly the magnitude of changes were similar between this rapid activation system and previous differentiation DamID studies (Peric-Hupkes et al. 2010; Robson et al. 2016) with 96% of LAD coverage shared between resting and activated cells and the remaining 4% roughly equally distributed between lost and newly formed LADs (Figures 2A and 3A). The 18.6% of genes found within LADs were less transcriptionally active than the 81.3% of genes in non-LADs (Fig. 2B), consistent with the periphery being repressive (Pickersgill et al. 2006; Pindyurin et al. 2007). Similar gene expression differences between LADs and non-LADs were observed comparing the Jurkat DamID data with published expression data from primary CD4$^+$ cells (Kundaje et al, 2015; Supplemental Fig. S2A). Hence, the immortalized T-cell lymphoblastic leukemia Jurkat and naïve CD4$^+$ T-cells display a broad correspondence between their gene expression state and lamina-association.

**Gene expression and function reflects lamina-association frequency**
On a genome-wide scale lamina-association frequency relates to gene expression levels (Fig. 2B) (Guelen et al. 2008). However, as LADs are enriched in developmentally-regulated loci (Peric-Hupkes et al. 2010), we tested if this genome-wide relationship applied to functional gene classes. 10 classes of genes (Supplemental Fig. S2B) that displayed distinct transcriptional dynamics in a published CD8^+ T-cell activation time course (Best et al. 2013) were analyzed for lamina-association frequency compared with their expression in resting Jurkat T-cells (Fig. 2C). This revealed that 5 gene classes involved in early stages of T-cell activation, such as cell cycle progression and early/short term effector function, were significantly depleted from LADs relative to the whole genome while displaying increased gene expression. In contrast, 4 gene classes not needed until later in T-cell responses, such as late effector and memory T-cell function, were found in LADs as frequently as the whole genome and were mostly transcriptionally repressed. These relationships are largely maintained when instead considering published expression data from naïve CD4^+ T-cells (Supplemental Fig. S2C) (Ni et al. 2016). However, 2 of the 5 gene classes depleted from LADs and expressed in resting Jurkat T-cells, active cell division and short term effector functions, were not expressed in naïve CD4^+ T-cells (Supplemental Fig. S2C). This likely reflects Jurkat cell origin as an actively dividing cancer cell line compared to the strong suppression of cell division found in primary resting naïve T-cells. Also, one class associated with naïve memory precursor function displayed elevated transcriptional activity in the naïve CD4^+ T-cells, which was absent in resting Jurkat T-cells.

All genes displaying induction and repression in Jurkat T-cell activation were similarly analyzed. As expected, genes becoming repressed during lymphocyte activation are active and depleted from LADs in resting cells, but surprisingly many induced genes were similarly depleted from LADs (Fig. 2D). However, although displaying elevated expression upon Jurkat T-cell activation, the induced genes also displayed elevated basal expression in the resting state relative to the whole genome (Fig. 2E) and this was recapitulated with naïve CD4^+ T-cells (Supplemental Fig. S2D). Depletion from LADs of basally expressed genes was also observed in myogenic
differentiation (Robson et al. 2016) (Supplemental Fig. S3A and B). Hence, LAD depletion invariably corresponds to increased gene expression for all gene classes tested.

**Important T-cell genes are released from the periphery and induced during T-cell activation**

We next sought to determine how gain or loss of LADs during T-cell activation correlates with gene expression changes. To ensure repositioning events to or from the periphery were identified with high confidence, we employed a stringent binary LAD or non-LAD definition. Whereas ~40% of the genome was in LADs in both resting and activated Jurkat T-cells, only 1.9% of the genome entered LADs in activated versus resting cells (interior-to-periphery regions, or IP) constituting 2,251 regions containing 554 genes. The opposite tendency (periphery-to-interior, or PI) was observed for 2.5% of the genome in 2,753 regions containing 792 genes (Fig. 3A). Size parameters of all LADs and differential IP/PI regions are summarized in Figure 3A. An example of log$_2$(B1/Dam-only) signal intensity PI shifts is shown in Figure 3B. Consistent with peripheral heterochromatin dissipation, decreased associations with the nuclear periphery dominated over increased associations during T-cell activation.

Accordingly, released PI regions contained significantly more induced than repressed genes (Fig. 3C). In the opposite direction, IP regions favored repression (Fig. 3C). Loci remaining at the periphery (PP) similarly favored repression. Nonetheless, the majority of genes moving to or from the nuclear periphery displayed no change in expression (Supplemental Fig. S3D). The trend towards the periphery being repressive was also observed in a recent myogenic study (Robson et al. 2016), but the association between repositioning and altered gene expression was stronger than for Jurkat activation (Supplemental Fig. S3C-E). DamID and microarray data for all genes is in Supplemental Table S1.

The subset of genes undergoing both repositioning and expression changes tended to be important for T-cell regulation based on the aforementioned lymphocyte activation functional
classes and GO-terms. To confirm their repositioning inferred from DamID, five genes were directly tested by FISH: guanylate binding protein (GBP) gene cluster, interleukin 2 (IL2), Cbl proto-oncogene B (CBLB), CD200 molecule (CD200), and B and T lymphocyte associated (BTLA). These were chosen for their being induced throughout the Jurkat T-cell activation time course (Fig. 3D, Expr.) and presence in PI regions (Fig. 3D, log₂(B1/Dam) and PI Regions). GBP1 regulates the cytoskeleton during T-cell activation (Sharma et al. 2011) while IL-2 is a potent inducer of T-cells (Boyman and Sprent 2012). The other three genes regulate or attenuate immune responses, encoding respectively a ubiquitin ligase, receptor and inhibitory ligand (Watanabe et al. 2003; Rygiel et al. 2009; Wallner et al. 2012). The absolute distance between the peak fluorescence intensity of the gene and co-stained lamin B2 was measured in the mid-plane from deconvolved wide-field images and plotted as a cumulative frequency plot. The FISH confirmed activation-induced release for all genes tested with differences statistically significant by Kolmogorov-Smirnov test (Fig. 3E,F). By contrast, an activated PP gene important for T-cell-vascular endothelium associations, vascular cell adhesion molecule 1 (VCAM1), was not repositioned following T-cell activation (Supplemental Fig. S4A-B), confirming that repositioning from the periphery is not whole-scale. As an additional control, peripheral distance measurements for IL2RA (also known as CD25), encoding the high-affinity IL-2 receptor subunit alpha that was in a non-LAD in both resting and activated cells, confirmed the DamID data (Supplemental Fig. S4C,D). The absence of a change in measured distance from the periphery for IL2RA also serves as a control that the increases measured for other loci represent bona fide gene repositioning rather than reflecting a global increase to nuclear volume.

**Peripherally sequestered enhancers are released from the periphery upon their induction**

Enhancers, cis-regulatory elements that activate target genes by physical proximity, become transcribed prior to their functional activation (Arner et al. 2015). We predicted that enhancers in
LADs would, like genes, reposition away from the periphery to be activated. To test this, a list of enhancer clusters/super-enhancers identified as functioning during T-cell activation by H3K27ac accumulation (Hnisz et al. 2013) were compared to Jurkat activation LADs. Roughly half of enhancer clusters were induced only in activated T-cells (Fig. 4A). Of these, only 27 occurred in LADs in the resting Jurkat T-cells, but over half of these were released from the periphery in the activated Jurkat T-cells (Fig. 4B).

FISH confirmed the PI repositioning upon activation of an enhancer cluster devoid of genes which accumulated H3K27ac and H3K4me1 upon activation of human CD4+ T-cells (Bernstein et al. 2010) (Fig. 4C,D). Hence, similar to genes, peripherally positioned enhancers are released concomitantly with their activation and acquisition of H3K27ac during T-cell activation.

Topologically Associated Domains form the unit of gene repositioning

LADs are reported to significantly overlap with TADs, chromosome regions displaying preferential local interactions (Dixon et al. 2012; Fraser et al. 2015; Kind et al. 2015; Jabbari and Bernardi 2017). As TADs are largely invariant between cell types (Rao et al. 2014; Dixon et al. 2015; Fraser et al. 2015), we hypothesized that regions with altered peripheral localization would be delimited to single TADs rather than spanning multiple TADs.

To investigate this we contrasted Jurkat activation DamID maps to the exact CD coordinates from a published GM12878 lymphoblastoid cell Hi-C dataset (Rao et al. 2014). Global analysis of their CD and our LAD distributions revealed that both resting and activated Jurkat LADs are frequently entirely contained within CDs. (Supplemental Fig. S5A). 500 randomly selected LADs were tested for their complete envelopment within CDs. As a control, frequency of complete envelopment was measured for the same 500 LADs after random shuffling along their chromosomes with non-mappable regions (e.g. centromeres) excluded from the shuffling. This process was repeated over 1,000 iterations to ensure all LADs in the genome were analyzed.
multiple times and frequencies of LAD envelopment within CDs plotted (Supplemental Fig. S5). All LADs occurred in their entirety within CDs significantly more frequently than for shuffled genome iterations. No statistically significant difference was observed between resting and activated LADs. We infer from this that a CD overlapping with a LAD repositions as a unit. A similar analysis using the GenometriCorr package, a tool that performs multiple analyses testing the spatial independence of domains along chromosomes (Favorov et al. 2012), confirmed the CD-LAD correspondence, uniform distance separation, and increased overlap frequency compared to chance (Supplemental Fig. S5B).

To determine if CD structure is similar in GM12878 and Jurkat cells, we performed pairwise FISH measurements of 5 fosmid probes containing sequences that were separated by 149 kbp from each other in series over the VCAM1 locus (Fig. 5A). Probes i and v were in neighboring CDs upstream and downstream of VCAM1. Probes ii and iv were within the VCAM1 CD boundary but close to the edges and probe iii in the center. The distance between different equally spaced probe pairs, two crossing a CD boundary (pair 1:i+iii and pair 3:iii+v) and one within the same CD (pair 2:ii+iv), was determined in 2D from max projection deconvolved images. Supporting similar GM12878 and resting Jurkat T-cell CD structure, probes within the same CD were significantly closer (0.15 µm) than probes between adjacent CDs (~0.30 µm) (Fig. 5B).

To test whether a CD is the unit of repositioning or if its structure changes during lymphocyte activation-induced gene repositioning, FISH was performed on resting and activated Jurkat T-cells using a similarly generated set of 5 probes (except 174 kbp apart) covering the GBP gene cluster (Fig. 5C). Again, the distance was significantly reduced for probe pairs within the same CD versus between adjacent CDs and this organization was maintained in both resting and activated states (Fig. 5D). Hence, TADs form the unit of radial repositioning within which LADs are frequently subsumed.
Changing LADs correlate with specific chromosome compartments

Compartments represent interactions between regions of chromosomes sharing similar functional states (Lieberman-Aiden et al. 2009; Dixon et al. 2012; Nora et al. 2012; Rao et al. 2014). Transcriptionally inactive B compartments can be further subdivided into B1 enriched in Polycomb H3K27me3, B2 and B3 correlating with LADs (Lieberman-Aiden et al. 2009; Gibcus and Dekker 2013; Rao et al. 2014), and B4 sub-compartments (not considered as they are restricted to chromosome 19). A sub-compartments are similar to one another in having active chromatin, but A2 displays increased H3K9me3 (Rao et al. 2014). As we now have mapped regions specifically changing at the periphery between resting and activated Jurkat T-cells, we sought to determine if Jurkat LADs and IP/PI regions correlated with any particular GM12878 sub-compartment.

First, we extracted the Rao et al sub-compartments and separately 1kb upstream and downstream flanking DNA. As compartment size varies, individual compartment regions were plotted with lengths equalized. The average \( \log_2 \) (Lamin B1/Dam-only) ratio from resting Jurkat T-cells was then plotted over these size-normalized compartment regions and their 1 kb sequences (Fig. 6A). As anticipated, in resting Jurkat T-cells the B2 and B3 sub-compartments exhibited a peak of \( \log_2 \) (Lamin B1/Dam-only) signal relative to flanking regions, while the A1 and A2 sub-compartments exhibited depletion. By contrast, polycomb-associated B1 sub-compartments exhibited only a modest increase in \( \log_2 \) (Lamin B1/Dam-only) signal. Comparing this data to a genome with randomly shuffled lamina-association, LADs are significantly enriched in B2 and B3 compartments and depleted in B1 and A1 sub-compartments (Fig. 6B). Moreover, correlating with the degree of diminished lamina-association, A1 sub-compartments were the most highly expressed followed by A2, B1 and B2/B3 sub-compartments (Fig. 6C).

Interestingly, the absolute DamID signal was much higher in the regions flanking the A2 sub-compartment than the regions flanking B1 and A1 sub-compartments (Fig. 6A). This striking difference in flanking lamina-association suggested that these compartments are spatially
segregated with respect to the nuclear periphery. The Log2(Lamin B1/Dam-only) signal within 1 Mbp of all A sub-compartment region boundaries was plotted as a heatmap, revealing that A1 sub-compartments tended to be further from LADs than A2 sub-compartments (Fig. 6D). To quantitate this, we plotted the distances from the midpoint of sub-compartment regions to the nearest LAD (Fig. 6E). This revealed, as expected, that B2 and B3 sub-compartments correspond to LADs (median distance 0 bp to the nearest LAD). The A2 sub-compartments were next closest to LADs (median distance 193 kbp) while B1 and A1 sub-compartments were significantly further away (median distances respectively 0.767 and 1.468 Mbp). Supporting the specificity of these results, A1, A2 and B1 sub-compartments all were further from LADs than expected in a genome with randomly shuffled LADs (Fig. 6E, Obs./Exp.). Statistically significant results were obtained by GenometriCorr analysis (Supplemental Fig. S6A). Therefore, we propose, due to differences in their proximity to LADs along chromosomes, that A1 and A2 sub-compartments represent radially segregated chromatin with respect to the periphery.

It follows that a gene being released from the periphery during lymphocyte activation would be more likely to enter an A2 than an A1 sub-compartment. Unfortunately, the GM12878 Hi-C data did not follow activation, nor was the activation state of their cells established. However, comparing the Jurkat activation DamID data with the GM12878 compartment maps revealed that many of the genes that were released from the periphery upon activation appeared in the A2 compartment. PI genes also displayed elevated expression in GM12878 cells relative to resting Jurkat T-cells (Supplemental Fig. S6B). Therefore we presume that the GM12878 lymphoblastoid cell data reflects a state more similar to that of an activated than resting T-cell. Thus, the presence of PI genes in LADs in the resting Jurkat T-cells likely indicates that they are in a B2/B3 sub-compartment until T-cell activation when they move to the A2 sub-compartment. Indeed, we observe that PI regions are significantly enriched in the A2 sub-compartment in GM12878 cells (Fig. 6F). Similar results were observed using GenometriCorr analysis, except that the B1 sub-
compartment was also enriched in PI regions (Fig S6C). The radial segregation of A1 and A2 sub-compartments was directly confirmed for several loci tested (Fig. 6G).

**Release of a predicted T-cell activation-specific enhancer from the nuclear periphery enables its association with similarly released genes in the interior**

The Jurkat activation system further enabled direct testing of this repositioning between compartments by FISH. Loci released from different B compartment LADs to an A2 compartment during T-cell activation would be predicted to display increased proximity in stimulated T-cells. Therefore, a 16 Mbp region of chromosome 3 was selected where several Jurkat activation PI genes and a predicted PI enhancer corresponded to an A2 compartment identified in GM12878 cells (Fig. 7A). Several of these genes were tested in Figure 3 to confirm release from LADs during Jurkat T-cell activation (CBLB, BTLA and CD200). We additionally tested NFKBIZ, which was induced during T-cell activation and in a GM12878 A2 sub-compartment, but only proximal to and not overlapping a PI region. This locus was close to the periphery in resting cells and became more internal during activation, indicating that genomic context is important to consider in addition to direct lamina-association itself (Fig. 7B).

All these genes were compared to the PI locus containing the predicted enhancer described in Figure 4C-D that occurs in a GM12878 A2 sub-compartment. FISH performed on combinations of the putative enhancer probe with the individual gene probes in resting and activated cells revealed that the distance between the putative enhancer and NFKBIZ, CBLB, CD200, and BTLA was much greater when the loci were at the periphery in resting cells than after their release upon activation (Fig. 7C). This is consistent with their being released from B2/B3 sub-compartments in resting Jurkat T-cells to associate in an A2 sub-compartment. By contrast, the GBP cluster and VCAM1 are similarly spaced along chromosome 1 (11 Mbp apart) and both are similarly in LADs in resting cells, but only GBP gets released upon Jurkat T-cell activation: accordingly, no significant change in spatial distance was observed upon activation (Supplemental Fig. S7) and the loci
display no GM12878 Hi-C interactions. Likewise, *IL2* and *IL2RA* located on separate chromosomes did not exhibit any significant change in distance (Supplemental Fig. S7B-C). Thus, multiple loci released and activated following T-cell activation display increased spatial proximity once in the nuclear interior.

**Release and predicted enhancer-gene association occur in primary lymphocyte activation**

The DamID was necessarily performed in the Jurkat model T-cell activation system due to poor transduction efficiencies of primary lymphocytes and the possibility of activation during lentiviral treatment. Thus, activation state differences for the small subset of activation gene classes (Fig. 2) necessitated validation of these findings in primary human lymphocytes.

Naïve CD4$^+$ T-cells were isolated from fresh human peripheral blood mononuclear cells by polychromatic FACS (Supplemental Fig. S8). Some were directly fixed for FISH and some were activated with Phorbol 12-myristate 13-acetate (PMA) for 24 h. PMA was chosen for activation to better match results to the published ChIP-seq study on naïve CD4$^+$ T-cells (Bernstein et al. 2010). The activated population was split for FISH and determining activation level by CD69$^+$ FACS (78%; Fig. 8A). The same probe pairs used in Figure 7 were used for FISH. In all cases the predicted enhancer became closer to the gene locus upon activation and this was significant for all except *NFKBIZ* (Fig. 8B). Also, as expected, the predicted enhancer, *NFKBIZ*, *CBLB*, *CD200*, and *BTLA* were all at the periphery in the resting primary lymphocytes and all moved away upon activation and repositioning was highly significant (Fig. 8C). Thus, the Jukats accurately reflect genome organization changes that occur upon human lymphocyte activation at least for the genes tested.
Discussion

Genes subject to spatial regulation are important for T-cell activation

Nuclear envelope-regulated genes in a myogenesis study were often required early in differentiation, but were inhibitory at later stages and so needed to be tightly regulated (Robson et al. 2016). Similarly, here, though first responder genes that initiate activation were absent from LADs, those subject to spatial regulation were important for subsequent T-cell function and regulation. For example, *IL2* encodes interleukin 2, that functions in propagating an immune response (Boyman and Sprent 2012), while the *GBP1* product regulates the cytoskeleton to influence T-cell signalling (Sharma et al. 2011). If these activities were not tightly shut down in resting cells they could potentially activate an immune response in the absence of infection. Others function in modulating the activation response. For example, induction of the nuclear IκB family member encoded by *NFKBIZ* is required for the subsequent expression of *IL17* following T-cell stimulation (Okamoto et al. 2010). Similarly, receptor-ligand products of *BTLA* and *CD200* and the *CBLB*-encoded ubiquitin ligase act to attenuate T-cell activation and thus fine tune the immune response (Watanabe et al. 2003; Rygiel et al. 2009; Wallner et al. 2012). Consistent with these important regulatory activities, the dysregulation or loss of function of a number these peripherally-regulated gene products is associated with autoimmunity (Watanabe et al. 2003; Rosenblum et al. 2004; Sharma et al. 2011; Wallner et al. 2012; Johansen et al. 2015; Hussman et al. 2016). Thus in all cases the tighter regulation of these genes in resting lymphocytes from adding spatial control would benefit the organism. Similarly regulating *cis*-regulatory elements such as enhancers is likely beneficial due to their ability to act on multiple target genes (Symmons and Spitz 2013). Importantly, the proximity of the A2 sub-compartment into which they are released from the periphery would potentially enable regulation also in the opposite direction.
The relationship between gene activity and lamina association

Specific gene repositioning to and from the periphery occurs during lymphocyte activation, but what controls tethering or release remains unclear. We find that frequency of lamina-association inversely relates to gene expression levels, consistent with previous studies (Guelen et al. 2008; Peric-Hupkes et al. 2010; Akhtar et al. 2013; Robson et al. 2016). It has been suggested that transcriptional activation drives gene release from the periphery (Tumbar et al. 1999; Tumbar and Belmont 2001; Chuang et al. 2006; Therizols et al. 2014) and the same could apply for enhancers that are frequently transcribed during induction (Arner et al. 2015). However, the relationship between gene position and expression is not absolute as we and others found the activity of most repositioning genes remains unaffected (Robson et al. 2016) (Peric-Hupkes et al. 2010).

Depletion of repressive chromatin modifications enriched in LADs such as H3K9me2/3 results in disrupted lamina association (Zullo et al. 2012; Bian et al. 2013; Demmerle et al. 2013; Kind et al. 2013; Harr et al. 2015), suggesting that the marks not only provide for silencing, but themselves contribute to locus affinity for the periphery. At the same time, specific nuclear envelope proteins are important for peripheral tethering of endogenous developmentally repositioning loci (Zullo et al. 2012; Solovei et al. 2013; Robson et al. 2016). Several studies argue that physical recruitment of a locus alone is insufficient for silencing, but works together with transcriptional regulators and other silencing factors at the periphery (Finlan et al. 2008; Robson et al. 2016). Thus, tethering to the nuclear envelope is a function of the chromatin state together with the milieu of nuclear envelope proteins.

Lamina-association provides a compass for interpreting compartment information
By contrasting Hi-C data with LADs we now place lamina-association and gene release within the context of higher-order chromatin structure. Chromosome compartments are defined based on what is close to each other, but this does not reveal location within the nucleus. Previous studies comparing Hi-C and DamID datasets revealed a tendency for B compartments to be enriched in LADs (Lieberman-Aiden et al. 2009; Gibcus and Dekker 2013; Rao et al. 2014); however, no spatial discrimination of active compartments has been described. Here we found that A1 and A2 sub-compartments segregate with their proximity to LADs, with A2 sub-compartments closer to the periphery. Thus the physical anchoring of distal regions in LADs can determine relative spatial positions for the wider genome. It follows that DamID differences between different cell types and differentiation stages could have wider consequences for the regulation of genes in the nuclear interior.

Our data suggests T-cell activation PI genes are switching from a LAD-associated B sub-compartment to a LAD-proximal A2 sub-compartment. The restriction in distance from the periphery for released loci could contribute to explaining a central question in the spatial genome organization field: how compartments are established and how particular genes find one another within the 3-dimensional space of the nucleus. In the analysis here, once released, loci remained relatively close to the periphery, generally <0.6 µm. This is an important limitation because modelling studies indicate that loci 10 Mbp apart are unlikely to meet in a single cell cycle by diffusion alone (Dekker and Mirny 2016). In contrast, a 0.5-0.8 µm space could be sampled by a locus in 1 h, well within the rapid response timeframe for lymphocyte activation and the distances we measured for released loci. This "constrained release" could thus increase the likelihood of incorporation of a released locus into an A2 sub-compartment (Fig. 8D) while keeping it close enough to the periphery to enable its sequestration again once T-cell activation has abated. Indeed, the activities of released genes in regulating T-cell function suggest such tight control may be necessary. A second functional consequence of constrained release may be to promote proximity between enhancers and distal target genes while at the same time sequestering them.
away from other inappropriate targets. Intriguingly, the putative enhancer described herein displays a long distance Hi-C looping contact with BTLA in GM12878 cells suggestive of such a functional association (Fig. 7A; Rao et al., 2014); however it remains untested if this is functionally significant. Nonetheless, data presented here clearly shows that LADs are intrinsically associated with gene regulation and have significant physical and, possibly, functional consequences for genes located in the nuclear interior.
Methods

Cell Culture, Transduction and Immune Conjugation

Jurkat clone E6-1 T-cells and Raji B-cells were cultured in RPMI 1640 supplemented with 10% FBS, 100 units/ml penicillin and 100 µg/ml streptomycin. VSV-G pseudotyped lentivirus’ encoding DamID, GFP-tagged, or pLKO constructs were generated and transduced as described in Supplemental Methods. Raji-B cells were incubated 30 min at 1 million cells/ml in media containing 0.5 µg/ml SEE (Toxin Technologies, Florida, EP404), followed by pelleting at 1,000 rpm for 5 min, two PBS washes and resuspension in growth medium. Conjugated Raji cells were mixed 1:10 with Jurkat T-cells to a total density of 1 million cells/ml for times indicated.

Isolation and Fluorescence Activated Cell Sorting (FACS) of Peripheral Blood Mononuclear Cells

Buffy coats supplied by the Scottish National Blood Transfusion Service were processed with the approval of the Committee for Governance of Blood & Tissue Samples for Non-Therapeutic Use. Human peripheral blood mononuclear cells (PBMCs) were isolated using Histopaque 1077 (Sigma), as per the manufactures instructions and described in detail in Supplemental Methods. CD3⁺ CD4⁺ CD62L⁺ CD25⁻ CD44⁻ naïve CD4⁺ T-cells were then incubated with selected antibodies and isolated from PMBCs using a 4-laser FACS Aria IIu flow cytometer (Beckton Dickinson). A more detailed description of primary cell FACs isolation in Supplemental Methods.

Fluorescence In Situ Hybridization (FISH)
PBS-washed Jurkat or primary CD4$^+$ T-cells were pelleted, spotted on polylysine-coated coverslips and fixed with 4% paraformaldehyde (PFA) and FISH performed as described previously (Zuleger et al. 2013). Distance measurements were performed using Imaris while co-staining with a lamin B2 antibody before FISH allowed the distance between genes to be determined using Imaris. A minimum of 50 cells were analyzed for each experiment.

**Microarrays**

RNA from Resting and Activated Jurkat T-cells was extracted with TRI-reagent (Invitrogen). The RNA was converted to biotin labelled-cRNA using the TotalPrep RNA Amplification Kit (Ambion, AMIL1791) following the manufacturer's instructions. For each experiment, three biological replicates were hybridized to Illumina whole genome gene expression arrays (HumanHT-12 v4 BeadChip). The data were quantile normalized using R (R Core Team 2010) and the Bioconductor package Limma (Smyth 2005). Differentially expressed transcripts with a log$_2$ ratio above 0.5 in absolute value were selected and adjusted for a false discovery rate of 5% (Benjamini and Hochberg 1995).

**DamID**

DamID was performed as in (Vogel et al. 2007) and is described in further detail in Supplmental Methods. Briefly, Resting Jurkat T-cells were transduced with Dam-Lamin B1- or Dam only-encoding lentiviruses over 24 h. Transduced cells were then incubated with Raji B-cells ± SEE for a further 48 h after which genomic DNA was sequenced (Beijing Genomics). Sequenced reads were mapped to human Hg19 genome and the log$_2$(Lamin B1/Dam) value determined for all genomic DpnI fragments in resting and activated cells, obtaining 2.2 and 2.1-fold genome coverage respectively. LADs were determined using a peak finder function in the BioConductor
package DNAcopy (Seshan VE and Olshen A (2016). **DNAcopy: DNA copy number data analysis.** R package version 1.46.0). DamID analysis results are derived from a single experiment.

**Hi-C Data Visualisation**

The displayed processed contact frequency maps are taken directly from the Rao et al. 2014 paper and displayed using the Juicebox Hi-C viewer (Rao et al. 2014).

**Supplemental information**

Plasmids, antibodies, lentiviruses, bioinformatics, listed FISH probes, external datasets used and more detailed DamID and FISH procedures are described in Supplemental Methods.

8 Supplemental Figures, and Supplemental Methods.

**Data access**

Jurkat T-cell DamID and gene expression datasets are available at GEO (Accession GSE94972).

**Acknowledgements**

We would like to thank Lior Pytowski and Anastasia Sokolidi for assistance with taking images, Shelagh Boyle for assistance with probe labelling, Dr David Kelly for microscopy support, and Louise Evenden at the Wellcome Trust Clinical Research Facility for microarray analysis. MIR was supported by a Wellcome Trust PhD Studentship and funding for this work was provided by Wellcome Trust grants 095209 to ECS and 092076 for the Centre for Cell Biology.
Author contributions

Conceived of study and designed experiments, MIR and ECS; performed experiments, MIR, RC, AS; analyzed data, MIR, JdIH and ARWK; wrote the paper, MIR and ECS.

Disclosure declaration

The authors declare no conflicts of interest.
References


Figure Legends

**Figure 1.** Transcription and genome organizational changes during T-cell activation. (A) Transcriptome and DamID analysis pipeline. (B) FACS plot of CD69 vs CMAC staining for Jurkat T-cells exposed to Raji B-cells lacking/presenting SEE antigen. Raji B-cells were gated from CMAC staining to distinguish them from Jurkat T-cells, revealing 97.5% Jurkat T-cell activation. (C) Heatmap of timecourse gene expression changes with 1.4 fold cutoff highlighting GO-terms positively-associated with T-cell activation or negatively-associated with cell division in the right-most columns (GO-terms in Supplemental Fig. S1). (D) Representative micrograph of GFP-encoding lentivirus-transduced Jurkat T-cells indicating high efficiencies for parallel transduced DamID constructs. (E) PCR amplified Dam-methylated genomic DNA. Amplification is only observed in Resting or Activated cells 72 h post-transduction. T4 DNA ligase null (-Lig) and DpnI null (-DpnI) controls represent background amplification. DamID results are derived from a single experiment while gene expression analysis was performed in triplicate.

**Figure 2.** Lamina-association frequency relates to gene expression and function. (A) Summary of resting LAD parameters. (B) Violin plot comparing the microarray-determined signal between genes within LADs and non-LADs. (C) Fold differences in gene expression and frequency of lamina-association in resting Jurkat T-cells for 10 published categories of genes displaying distinct transcriptional behaviors during T-cell activation (Fig. S2; Best et al. 2013). Fold differences were calculated relative to the whole genome (WG) in resting Jurkat T-cells. Higher expressed genes tend to be depleted from LADs, such as categories associated with naïve cells, early effector function and cell division. LADs associated more with late effector/memory function classes. (D) Percentage of genes in resting Jurkat LADs from different expression categories: genes induced (Ind.) or repressed (Rep.) at least 1.4 fold during T-cell activation, unchanged genes (UC.) and all genes. Both induced and repressed genes are 3-4 times less frequently found in LADs compared
to unchanging genes. (E) Violin plot for expression levels of induced, repressed, and unchanging genes. Induced and repressed categories are both higher expressed than the unchanging genes in resting Jurkat T-cells. For B, C and E, significance of differential gene expression was determined by Dunn test following a significant Kruskal-Wallis test. For C and D lamina-association frequency statistical significance was determined by Fisher tests. \( *P < 0.05, \quad **P < 0.01 \) and \( ***P < 0.001 \). See also Supplemental Figs. S2 and S3 and Table S1.

**Figure 3.** Important genes are released from the periphery during T-cell activation. (A) Summary of LADs, IP (interior-to-periphery) and PI (periphery-to-interior) regions between resting and activated Jurkat T-cells. (B) Genomic region surrounding ANTRX2 showing DamID signal bar plots and heatmaps (HM), LADs, IP and PI regions, subtracted Resting-Activated DamID signal, and expression changes in Jurkat T-cells during activation from the resting state at 8, 24, and 48 h. The key for \( \log_2(\Delta \text{expr.}) \) is given in the bottom right corner. ANTRX2 locus is highlighted by light shading. (C) Ratio of genes induced (Ind.) and repressed (Rep.) during T-cell activation for the whole genome (All), PI, IP, II (genes in the interior in both cell states) and PP (genes peripheral in both states) regions. Statistical significance was determined by Fisher tests. \( *P < 0.05, \quad **P < 0.01 \) and \( ***P < 0.001 \). (D) Genome browser views for the genomic region surrounding the GBP gene cluster and CBLB, CD200, BTLA and IL2 loci covering the same parameters as in B. (E,F) Representative micrographs and quantification of gene positions in resting and activated cells relative to lamin B2. Median points are marked with dashed red lines. For quantification statistics, the position of loci in the activated sample was compared to the resting sample by KS tests. \( *P < 0.05, \quad **P < 0.01 \) and \( ***P < 0.001 \). See also Supplemental Fig. S4.

**Figure 4.** Enhancers are released from the periphery during T-cell activation. (A) Upon stimulation of naïve CD4\(^+\) T-cells with PMA, 513 out of 974 predicted enhancer clusters identified by Hnisz et
of the 513 induced enhancer clusters, 27 were in LADs in resting Jurkat T-cells (black circle). Of these, 17 become released upon T-cell activation (yellow circle). (C) A 66 kbp chromosome 3 region containing a putative enhancer induced during T-cell activation releases from the periphery (PI) in activated Jurkat T-cells. Categories used in Figure 3 are shown along with ChIP-seq tracks from primary naïve and PMA-activated CD4+ T-cells for H3K4me1 and H3K27ac histone modifications marking active enhancers (data from Bernstein et al, 2010). The H3K27ac peaks defined as enhancers (Enh.) and super enhancers (S. Enh.) by Hnisz et al are also displayed as tracks. (D) Quantification of putative enhancer position in resting and activated Jurkat T-cells cells relative to lamin B2, confirming its activation-induced release. For quantification statistics, the position of loci in the activated sample was compared to the resting sample by KS tests. *P < 0.05, **P < 0.01 and ***P < 0.001.

Figure 5. The unit of repositioning is the Contact Domain. (A) Genomic region surrounding VCAM1 showing Hi-C data and associated CDs (yellow lines on contact frequency heatmap and green track) from published GM12878 lymphoblastoid cell data (Rao et al. 2014). Orange track, TADs from published human embryonic stem cell (hESC) data (Dixon et al. 2012). Other tracks, published ChIP-seq CTCF GM12878 cell data (ENCOD Project Consortium 2012), DamID signals, LADs, and probes and probe pairings tested in by FISH in panel B. (B) Quantification and representative max projection images measuring distance between pairs of ~150 kbp spaced FISH probes. Pairs 1 and 3, containing probe pairings from different CDs, are separated significantly further than probes within the same CD used for Pair 2. (C) Similar genomic region view of the GBP gene cluster and probes tested, showing data for resting and activated Jurkat T-cells. (D) Similar analysis as B for pairs of ~175 kbp spaced FISH probes in resting and activated Jurkat T-cells. While pairs crossing CDs are again separated more than those within, the distances between probes in all pairs remain unchanged during T-cell activation, indicating that CD structure is maintained. Quantification statistics were determined by comparing the separation of Pairs 1 and 3.
with Pair 1 in resting (grey) and activated (black) cells by KS tests. *$P < 0.05$, **$P < 0.01$ and ***$P < 0.001$. See also Supplemental Fig. S5.

**Figure 6.** Compartments are distributed non-randomly with respect to LADs. (A) Size-normalized plot displaying the average $\log_2$(Lamin B1/Dam-only) signal from resting Jurkat T-cells across all A1, A2, B1, B2 or B3 GM12878 sub-compartments and 1 kbp of flanking DNA. Compartment data from Rao et al, 2014. For the plot, subcompartment region sizes were normalized to a uniform standard. (B) Fold enrichment analysis of LADs observed (obs) in each compartment compared to that expected in a genome with randomly shuffled LADs (exp). (C) Box plot displaying the expression distribution in resting Jurkat T-cells for each GM12878 sub-compartment. (D) Heatmap of DamID $\log_2$(laminB1/Dam) signal across +/- 1 Mbp surrounding A sub-compartment regions. Regions are ranked to the right by decreasing distance to nearest LADs and their identity as an A1 or A2 sub-compartment region highlighted below in red or blue, respectively. A2 domains are observed more frequently closer to LADs than A1. (E) Box plot displaying the distribution of chromosomal distances between the midpoint of sub-compartment regions and the nearest LAD in resting Jurkat T-cells. Below is the ratio between observed (Obs.) median distance and that expected (Exp.) in a randomly LAD-shuffled genome. Values are color-coded to show increased (red) or decreased (blue) observed distance compared to expected. (F) Fold enrichment analysis of PI regions observed in each compartment compared to that expected in a genome with randomly shuffled PI regions. (G) Quantification of average distance between A1 (thick red line) or A2 loci (thick blue line) and the lamina determined by FISH for activated Jurkat T-cells where PI loci are detached from the lamina. Lighter lines designate individual A2 (GBP, NFKBIZ, CBLB, CD200, BTLA) and A1 (IL2RA, CCND2, TNF) loci. For B and F significance was determined by Fisher tests while for C and E significance was by Dunn test for multiple comparison testing after a significant Kruskal-Wallis test. For G the significance of the difference between A1 and A2
distances to lamina was determined by KS tests. *P < 0.05, **P < 0.01 and ***P < 0.001. See also Supplemental Fig. S6.

**Figure 7.** Genes released from LADs during T-cell activation associate in the interior in activated Jurkat T-cells. (A) Genomic view of a 17 Mbp region of chromosome 3 displaying GM12878 cell Hi-C interactions from published contact maps (Rao et al. 2014). Hi-C data is displayed as a heatmap of the ratio of observed versus expected contact frequencies in a distance-scaled model. GM12878 sub-compartments (A1, A2, B1, B2, B3) are color coded directly beneath the map. Tracks are displayed as in previous figures. The black circle represents a highly significant DNA-DNA contact detected by Rao, et al. Several genes, NFkBIZ, CBLB, CD200 and BTLA, and a T-cell activation-specific enhancer (Enh.), display reduced lamina association during T-cell activation concomitant with transcriptional induction and are present in the GM12878 A2 sub-compartment. (B) Representative micrographs and quantification NFkBIZ positions in resting and activated cells relative to lamin B2. (C) Representative micrographs and violin plot quantification of distances between the enhancer and indicated genes in resting and activated Jurkat T-cells. All enhancer-gene FISH probe combinations display reduced inter-probe distance following activation. For quantification statistics, relative distance measurements between loci or between loci and lamin B2 were compared between the activated and resting cells by KS tests. *P < 0.05, **P < 0.01 and ***P < 0.001. See also Supplemental Fig. S7.

**Figure 8.** LAD release and association of genes in the nuclear interior also occurs in primary CD4+ T-cells. (A). FACS plot of CD69 staining for naïve (resting, black line) and PMA-treated (activated, red line) CD4+ T-cells indicating 78% activation. See also Supplemental Fig. S8 for details of the naïve population. (B) Violin plot quantification of distances between the predicted enhancer and indicated genes in naïve (light grey) and PMA-treated (black) cells. All loci displayed closer
proximity to the predicted enhancer locus upon PMA activation of naïve CD4+ T-cells. (C) Violin plot quantification of distances between indicated loci and lamin B2 in naïve (light grey) and PMA-treated (black) cells. All loci displayed increased distance from the periphery upon PMA activation. For quantification statistics, relative distance measurements between loci or between loci and lamin B2 were compared between the activated and resting cells by KS tests. *P < 0.05, **P < 0.01 and ***P < 0.001. (D) Model of the relationship between LADs, compartments and gene release. In resting T-cells B2 and B3 sub-compartments exist as LADs while A2 sub-compartment regions remain near to the lamina due to their proximity along the chromosome. By contrast the A1 and B1 compartments, being further from LADs along chromosomes, are distributed more distally in the nuclear interior (not shown). Following T-cell activation, regions of LADs containing necessary gene or enhancer loci are released from the periphery where they associate in the A2 compartment proximal to the lamina in a transcriptionally active state.
Figure 1.

A

+ Dam Virus

Gene Expr Sample

DamID Sample

+Raji -SEE (Rest)

t = -24h

+Raji +SEE (Act)

t = -0h

Dam Methylase Activity

B

Jurkat + Raji(-SEE) 24 h

Act. Jurkat - 2.1%

CD69 (Alexa 647)

Act. Raji

0 10^2 10^3 10^4 10^5 10^6

CMAC (Indo-1)

Rest. Jurkat - 97.9%

Rest. Raji

0 10^2 10^3 10^4 10^5 10^6

CMAC (Indo-1)

Jurkat + Raji(+SEE) 24 h

Act. Jurkat - 97.5%

Act. Raji

Rest. Jurkat - 2.5%

Rest. Raji

D

DAPI

GFP-DAPI

10 μm

E

48h Transduction

72h Transduction

8h 24h 48h

Figure 2.

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B

C

D

E
### Figure 3.

**A**

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**B**

- **Chr 4**: 79 Mbp, 83 Mbp, 87 Mbp

**C**

- Ratio Ind./Rep. Genes
- Gene Class

**D**

- **GBP**
  - Chr 1: 89.5 Mbp, 90 Mbp
- **CBLB**
  - Chr 3: 105.5 Mbp, 106 Mbp
- **CD200 and BTLA**
  - Chr 3: 112 Mbp, 12.5 Mbp
- **IL2**
  - Chr 4: 123 Mbp, 123.5 Mbp

**E**

- GBP, CBLB, BTLA, IL2

**F**

- GBP, CBLB, CD200 and BTLA, IL2

---

*P = 0.007**

*P < 0.001"
Figure 4.
Figure 7.

A

GM12878 Hi-C (Rao et al, 2014)
Observed/Expected
Chr 3: 97-114 Mbp

B

NFKBIZ Lamin B2

Rest.  Act.

C

Enh. CBLB Lamin B2

Rest.  Act.

Enh. BTLA Lamin B2

Rest.  Act.

**P < 0.01

Cumulative frequency

Distance to Enhancer (μm)

Distance to Lamina (μm)

0.0  0.5  1.0  1.5  2.0

0.0  0.2  0.4  0.6  0.8  1.0

0.54 μm  0.40 μm  0.47 μm  0.83 μm  0.84 μm

0.96 μm  0.57 μm  0.53 μm

1: Enh.-NFKBIZ  2: Enh.-CBLB  3: Enh.-CD200  4: Enh.-BTLA
Figure 8.