Nanoscale spatial organization of the HoxD gene cluster in distinct transcriptional states

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Chromatin condensation plays an important role in the regulation of gene expression. Recently, it was shown that the transcriptional activation of HoxD genes during vertebrate digit development involves modifications in 3D interactions within and around the HoxD gene cluster. This reorganization follows a global transition from one set of regulatory contacts to another, between two topologically associating domains (TADs) located on either side of the HoxD locus. Here, we use 3D DNA FISH to assess the spatial organization of chromatin at and around the HoxD gene cluster and report that although the two TADs are tightly associated, they appear as spatially distinct units. We measured the relative position of genes within the cluster and found that they segregate over long distances, suggesting that a physical elongation of the HoxD cluster can occur. We analyzed this possibility by super-resolution imaging (STORM) and found that tissues with distinct transcriptional activity exhibit differing degrees of elongation. We also observed that the morphological change of the HoxD cluster in developing digits is associated with its position at the boundary between the two TADs. Such variations in the fine-scale architecture of the gene cluster suggest causal links among its spatial configuration, transcriptional activation, and the flanking chromatin context.

In the nuclei of mammalian cells, chromatin is packaged according to several levels of organization (1–3), which can either reflect or affect transcriptional regulation (e.g., ref. 4). By combining DNA FISH and microscopy, it was shown that chromatin decondensation occurs concomitantly with transcriptional activation (5), suggesting that the opening of chromatin makes gene promoters accessible for transcription. Recently, however, studies involving super-resolution microscopy have revealed a more complex relationship, showing that a higher compaction of chromatin can also be associated with an active state of transcription (6). In this latter case, the compaction of local regulatory elements allowed for a stronger enhancer effect, leading to a more robust activation.

Approaches based on chromosome conformation capture [and derivatives thereof (7)] at the mammalian HoxD locus have revealed that interactions between genes and their enhancers can occur not only during an active phase of transcription but also in the absence of transcriptional read out, in cells that do not necessarily express the related target genes (8). Such constitutive contacts covering large regulatory landscapes and their target gene or genes were found to be present in mammals genome-wide (9) and are referred to as topologically associating domains, or TADs (9, 10). TADs have been associated with a variety of regulatory functions (11), either in their implementation (e.g., ref. 12) or in their emergence during vertebrate evolution (13, 14).

Hox gene clusters have been successfully used to study the functional organization of TADs (15–17), as well as the relationship between the progressive decompaction of both genes and enhancers and their transcriptional read-out. Studies of the mouse HoxD cluster have provided insights into the global regulation of its nine consecutive genes during limb development, including the presence of multiple regulatory sequences spanning a 2-megabase large DNA interval (18). Recently, this gene cluster, similar to its HoxA relative (14, 15), was shown to reside at a boundary between two TADs (located ca. between Hoxd11 and Hoxd12), with each TAD containing enhancers required to regulate different subgroups of genes in developing organs or structures (9, 16, 19) such as distal limbs, proximal limbs, genitals, or the eccum. Interestingly, all enhancers sharing a particular specificity are found within the same TAD, and thus far, no cell type or tissue was reported where these two opposite regulatory landscapes would operate concomitantly (8, 16, 17, 19), suggesting a functional switch occurs between these two TADs in their capacity to regulate subsets of target Hox genes.

TADs were originally defined by biochemical approaches (9, 10). The correspondence between the averaging of multiple interactions, some of them of unknown significance, on the one hand (see ref. 20), and a chromatin structure in the nuclear space of single cells, on the other hand, is of great interest and has recently come under discussion (see ref. 21). In this study, we used DNA FISH to show that the two TADs splitting the HoxD locus are distinct chromatin units, which rarely overlap despite their close association in space. Within this well-defined 3D organization, HoxD genes can segregate over large distances. By using super-resolution microscopy, we observe that these large distances result from extensive elongation events over relatively short genomic distances, which appear to be maximal in tissue with high levels of HoxD gene transcription. Our data suggest this elongation of the HoxD cluster is facilitated by its genomic position at the boundary between two TADs.

**Significance**

Ultrastructural chromatin dynamics may play a key role in regulating transcriptional activation. Here we have used super-resolution microscopy to study the folding mechanics of the HoxD cluster, as assayed by following the elongation of chromatin in single cells with different status of Hox gene activation. We observed that the spatial separation of HoxD genes is strongest in those tissues where they are highly expressed. We also document that the opening of chromatin precedes transcription and that the strongest elongations are observed at the location of the boundary between two major topologically associating domains (TADs). These results shed light on how spatial compartmentalization is achieved, likely to accompany efficient chromatin re-organization upon activation of transcriptional switches.


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Data deposition: The RNA-seq data reported in this paper have been deposited in the Gene Expression Omnibus (GEO) database, www.ncbi.nlm.nih.gov/geo (accession number: GSE72865).

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Results

Hoxd Genes Are Localized at the Boundary Between Two TADs. To directly observe the chromatin topology surrounding the HoxD cluster, we used 3D DNA FISH by labeling several consecutive BACs per TAD, such as to cover each TAD with a different color (Fig. 1A and SI Appendix, Table S1), and thus to assess the general chromatin spatial organization surrounding HoxD (Fig. 1B). In both active (forelimb) and inactive (forebrain) cells, we observed that TADs are organized as two dense and distinct units, closely associated in the nuclear space (Fig. 1C). High-resolution images using structured illumination microscopy showed that the TADs did not intermingle, and rarely overlapped in any substantial manner (Fig. 1D). This segregation was much less pronounced when BAC probes belonging to the same TAD were used as controls (SI Appendix, Fig. S1).

We measured the distance between TAD centers, which ranged from 0.20 to 1.40 μm, with an average of about 0.79 μm (σ ≈ 0.100 pairs). Because previous studies using chromosome conformation capture (4C) revealed a dynamic switch in contacts between Hoxd genes and the two TADs in the proximal versus distal parts of developing forelimbs (16), we assessed whether this inter-TAD distance would change between these distinct limb domains, but found no significant difference (Fig. 1B, C, and E). A detailed analysis of chromatin morphology revealed that the TADs are rather compact, with low ellipticity (Fig. 1F). We nevertheless observed a slight variation in ellipticity in cellular domains displaying different Hoxd gene expression patterns. Both TADs indeed appeared slightly less elongated in presumptive digit cells in the distal part of the limb (Fig. 1F), yet their global structure was rather stable in all analyzed situations. Finally, we assessed the position of the HoxD cluster relative to the TADs, using a 39-kb large fosmid probe covering the Hoxd8 to Hoxd12 genes. As expected, the signal was consistently detected at the interface between the two chromatin domains (Fig. 1G). In some cases, the HoxD cluster signal appeared somewhat elongated (arrowheads, Fig. 1G).

Hoxd Genes Are Close to Their Regulatory Islands. The fact that TADs display comparably condensed aspects in both proximal and distal limb cells in vivo, that is, independent from their transcriptional activities, supports the view of a preformed or predesigned background regulatory structure (8); that is, a state of preferential chromatin compaction within a given TAD in which constitutive interactions occur without necessarily eliciting a transcriptional output (9, 11). In a previous study using FISH in ES cells, the Hoxd1 gene localized at a substantial distance from Hoxd13, suggesting that the HoxD cluster could display spatially distinct moieties with its centromeric region (containing Hoxd13) positioned at the TAD boundary (22). To assess whether this apparent decompaction of Hoxd genes is linked to their relative topological position within the cluster and, consequently, whether their inclusion into either one of the TADs or their location at the inter-TAD boundary could explain this tendency to decompact, we examined the spatial relationships between four different parts of the HoxD cluster.

We used fosmid clones either covering the Hoxd1 gene or the Hoxd3 to Hoxd4 region, (i.e., two regions normally included within the telomeric TAD), covering the Hoxd13 gene, and hence entirely positioned within the centromeric TAD, or covering the Hoxd8 to Hoxd12 region located at or around the inter-TAD boundary (16) (Fig. 1H). We measured the distances between combinations of two probes and found that parts of the HoxD cluster could be separated from one another by more than 500 nm (Fig. 1I–K). Of note, the distances measured between the centers of both signals generally correlate with the genomic distance (Fig. 1J and SI Appendix, Table S2). In this respect, we noticed that the telomeric extremity of the HoxD cluster, as monitored by the Hoxd1 probe, was clearly separated from the main part of the cluster, with an average distance between Hoxd1 and Hoxd11 or Hoxd13 of more than 700 nm (Fig. 1J). This apparent separation between Hoxd1 and the rest of the cluster was, however, not constant and followed a broad distribution (Fig. 1K).

Super-Resolution Microscopy of the HoxD Locus. This spatial separation between different parts of the HoxD cluster is reminiscent of a model in which groups of active and inactive genes within HoxD form physically distinct and highly dynamic subdomains (23). To more directly examine the conformation of the HoxD cluster in single cells, we labeled its entire genomic sequence by using a 160-kb large BAC as a DNA probe and carried out DNA FISH. The morphology of the HoxD cluster was subsequently resolved using stochastic optical reconstruction microscopy (STORM), a type of super-resolution imaging based on sequential imaging of single-molecule signals (24–26). As a control, we used a probe of similar size located nearby, within the telomeric gene desert (Fig. 2A).

We initially examined sections of embryonic brain where the HoxD cluster is transcriptionally silent. Both the control and HoxD probes appeared as compact objects (Fig. 2B and E). However, cells from the developing distal forelimb; that is, a tissue with robust transcription of at least the Hoxd13 to Hoxd9 genes (27) (Fig. 3), displayed a higher diversity of morphologies (Fig. 2C and F). Although the total area of the object appeared unchanged, the signal detected in many cells of the forelimb tissues was substantially more elongated (SI Appendix, Fig. S2). We quantified this variation in elongation by the aspect ratio and the circularity of the objects. The major axis of the cluster divided by its minor axis (aspect ratio) was significantly higher in active forelimb cells than in inactive forebrain cells (Fig. 2H), where circularity was also lower (Fig. 2I).

As elongation was maximal in limb cells displaying a high level of HoxD transcription, we assessed whether transcription could cause elongation by evaluating the cluster morphology in ES cells, a cell type in which Hoxd genes have not yet been activated and are labeled by positive and negative chromatin marks, potentially reflective of a poised transcriptional state (28). In these cells, many alleles showed an elongated aspect, with a circularity index close to that scored in forelimb cells, and thus significantly different from the data obtained in brain cells (Fig. 2D, G, and I). From this observation, we conclude that the elongation observed by using STORM microscopy seems to be an important feature of an “open” HoxD cluster, observed either in ES cells where the cluster is poised to be transcribed or in cells where transcription indeed occurs. Finally, we measured the extent of the elongated structure and found that the signals can span more than 500 nm. This high extension rate is consistent with our previous measurements (Fig. 1J) and is in the range predicted by mathematical modeling (29).

Chromatin Elongation at Super Resolution. The mechanisms underly- ing such an elongation process are unclear. Opening and/or relaxation of chromatin may be a consequence of active transcription driven by elongating RNA polymerase II and consequent chromatin remodeling. Alternatively, changes in chromatin morphology could result from states of histone modifications not necessarily associated with different transcriptional activities. For example, in both brain and ES cells, the HoxD cluster is covered by polycomb-associated chromatin marks such as H3K27me3, which may participate in its compaction via the recruitment of the PRC1 complex (22, 28, 30). However, the H3K27me3 coverage observed over HoxD in ES cells is significantly weaker than in brain cells (23), which may be related to the presence of H3K4me3 marks in the former sample. Accordingly, a poised chromatin state may show a level of decompaction comparable to that of transcriptionally active chromatin.

We investigated this issue by using super-resolution microscopy with four smaller DNA probes spanning different sub-regions of the HoxD cluster (Fig. 1H, enlarged in Fig. 3A). These subregions displayed differential coverage by polycomb-associated marks in either distal or proximal limb bud cells (16), and thus were used to evaluate a potential link between the elongation
of the cluster and either its transcriptional activity or the presence of polycomb-associated proteins. For instance, the probe containing Hoxd1 is covered by H3K27me3 marks in distal cells, whereas the Hoxd8 to Hoxd12 fragment shows a moderate coverage. In contrast, the probes encompassing either the Hoxd11 to Hoxd13 or the Hoxd13 to Evx2 DNA intervals are free of H3K27me3 in embryonic day (E) 12.5 distal and active cells, likely as a consequence of high transcriptional activity (Fig. 3A).

Each fosmid clone was labeled with the same Alexa 647 fluorophore and its signal analyzed by STORM microscopy. By comparing the signal morphologies (Fig. 3B), we noticed that the Hoxd1 signals did not appear significantly more compact than the others (Fig. 3B; representative examples). If anything, this area of the HoxD cluster, which is densely covered by H3K27me3 marks, was slightly less circular than images obtained from the fosmids covering regions with lower amounts of H3K27me3 and highly expressed genes (Fig. 3C). To complement this observation, we labeled the two fosmid clones containing either the Hoxd1 or the Hoxd8 to Hoxd12 regions, using Alexa 647 and Alexa 555, respectively, to directly assess their relative morphologies within the same cells. These experiments confirmed that these two DNA regions were decompacted to similar extents in the same cells (SI Appendix, Fig. S2B). The most extensive elongation observed (Hoxd8-d12, Fig. 3B and C) matches the set of genes that was shown to have cell-specific long-range DNA contacts in the developing forelimb (16). To test whether decompaction could change with a modification of long-range contacts, we compared the two tissues in which the pattern of long-range interactions differs drastically: the proximal and distal forelimbs (Fig. 3D). Here we observed similar structures, both of them significantly more dispersed than that which was observed in the forebrain, used as a negative control (Fig. 3D).

We next investigated whether such a decompaction would also be observed in another tissue in which the HoxD cluster is strongly activated, and thus imaged the Hoxd8-d12 region in the developing trunk. We compared forebrain and trunk samples in which the cluster is completely inactive, anterior trunk cells in which transcription occurs from Hoxd1 to Hoxd8 only, and posterior trunk cells in which Hoxd1 to Hoxd12 are activated. We observed the strongest decompaction in those tissue with the highest level of transcription (Fig. 3E), suggesting the existence of a link between transcription and the unfolded structure of Hox genes.

**Effect of the Gene Deserts on Compaction.** 4C studies using microdissection of limb samples have indicated that parts of the HoxD cluster are strongly contacted by enhancer sequences located in both the centromeric and telomeric TADs (8, 16). Such strong enhancer contacts may exert forces upon the HoxD cluster, leading to variations in its global chromatin architecture. We assessed this possibility by analyzing three genetic perturbations in which Hoxd genes were separated from their respective enhancers by large targeted inversions (31, 32). The first allele was an inversion separating Hoxd from its telomeric TAD by displacing the latter by 28 Mb (Inv(antPc-d44); Fig. 3F), which switched off Hoxd-11 in proximal limb cells and decreased their transcription in distal limb cells. The morphologies of signals detected when using this large inversion were as elongated as in noninverted cells, showing that the immediate regulatory neighborhood, even though it contains sequences interacting with the probe, was not critical in the shaping of the cluster, at least in these particular conditions. In both the inverted and wild-type loci, a low circularity index was determined for the Hoxd8 to Hoxd12 DNA region (Fig. 3 G and H), suggesting that the presence of genuine enhancer sequences and constitutive interactions (16) is not necessary for the decompaction of the HoxD cluster.

We also assessed the Inv(Nsi-Jig6) centromeric inversion (SI Appendix, Fig. S3 A and B), which repositions the centromeric regulatory sequences several Mb away from the cluster, thus abrogating all specific enhancer–promoter interactions occurring during digit development and, consequently, turning off Hoxd.

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**Fig. 1.** The HoxD cluster is at the interface between two TADs. (A) Schematic of the centromeric and telomeric TADs (C-TAD; T-TAD) using public data available under ref. 9 with genes as black boxes below and the relative localization of the probes (in red, blue and green) used in the DNA FISH experiments. The red dotted line represents the TAD boundary within HoxD. (B) E12.5 distal forelimb nuclei stained with DAPI (blue) and the centromeric (red) and telomeric (green) TADs. (Scale bar, 1 μm.) (C) Schematic (Left) and distributions (Right) of the various configurations observed in FISH in B. Prox and Dist, proximal and distal limb cells, respectively. (D) An example of structured illumination microscopy showing the absence of overlap between the two TADs. (Scale bar, 500 nm.) (E and F) Quantifications of the parameters observed under B. (E) Distances between the centers of both TADs. n.s., nonsignificant, using a Kruskall-Wallis test followed by Dunn’s multiple comparison posttest. (F) Ellipticity measured for both the telomeric (left) and centromeric (right). *P < 0.0001, using unpaired t tests. (G) Position of the HoxD cluster for two alleles of a representative distal forelimb cell, using a fosmid probe specific for the genes Hoxd8 to Hoxd12 (green). The signal is scored between the centromeric (red) and telomeric (magenta) TADs. (Scale bar, 500 nm.) (Right) Close-ups of both alleles with white arrowheads showing an elongated HoxD cluster. (Scale bar, 200 nm.) (H-K) Quantification of the distribution of interprobe distances between selected pairs of probes. The statistical significance between datasets was tested using a Kruskall-Wallis test followed by Dunn’s multiple comparison posttest. All were significantly different (*P < 0.05) except the ones indicated with n.s. See SI Appendix, Table S2 for details. (K) Frequency distribution of the measurements shown in J.
genes transcription (33). Again, the decompacted aspect of the HoxD cluster was not dramatically modified after STORM imaging (SI Appendix, Fig. S3B). Finally, we used a large inversion, with a breakpoint located between Hoxd10 and Hoxd11, thus splitting the HoxD cluster into two parts, leaving in place only from Hoxd11 to Hoxd13 and their centromeric gene desert (ref. 31; scheme in SI Appendix, Fig. S3). We monitored the aspect ratio of the short remaining cluster, using the fosmid probe covering from Hoxd11 to Hoxd13, and found no difference in elongation (SI Appendix, Fig. S3C), suggesting a full HoxD cluster is not critical to spatially organizing chromatin in its various parts.

Discussion

The organization of chromatin in the nuclear space is a critical parameter for the proper control of transcription, and the extent of chromatin elongation at a given genetic locus may reflect its capacity to be efficiently transcribed (34). We previously determined that the HoxD gene cluster was the target of a bimodal type of global regulation, exerted from either the telomeric or centromeric gene deserts, two regulatory landscapes matching TADs, and separated by the HoxD cluster itself (9, 16). However, chromosome conformation capture-based experiments average the treatment of several million independent cells, and the presence of dense and compact chromatin architectures flanking the HoxD cluster remained to be shown at the cellular level. Here, by concomitantly labeling series of BACs spanning specifically the two TADs, we demonstrate that such configurations indeed exist as dense and separate objects in individual cells, as previously seen on a specific locus at the X-chromosome (10), with a low level of partial overlap consistent with the reduced number of interactions observed in 4C between the two TADs (16, 17). This observation was confirmed by structured illumination microscopy. The distances between TADs were comparable between cells, whether or not they expressed Hoxd genes, supporting the existence of a poised regulatory structure already present in the absence of transcription (8). However, the direct visualization of TADs at this locus is not informative regarding potential fluctuations in contacts within each TAD, from one cell to another (20). It nevertheless demonstrates that such dense structures exist in all cells and on both alleles.

By using super-resolution microscopy (STORM), we scored the most extensively elongated forms in cells from either the developing distal forelimb or the posterior trunk, two tissues in which several Hoxd genes are strongly active. In contrast, dense and compact structures were scored either in brain cells, negative for all Hox gene transcription at this stage, or when using a control probe located outside the gene cluster. These elongated morphologies generally occurred along a major axis and appeared continuous, either when using large probes covering the whole cluster or with smaller probes detecting only one or a few genes. In distal forelimb cells, however, where the Hoxd1 to Hoxd4 region is covered by H3K27me3 chromatin marks (16), we did not observe any significant asymmetric compaction. In contrast, the inactive part of the HoxD cluster was at least as elongated as the transcriptionally active region. One explanation for this unexpected observation may be the robust and nonproductive contacts established by HoxA1 with the telomeric TAD (16), which may lead to tensions elongating the gene cluster (see following).

A significant elongation of the gene cluster was also scored in ES cells, even though the entire HoxD cluster is labeled with both H3K4me3 and H3K27me3 chromatin marks (22, 28, 30) associated with the formation of local chromatin domains (23, 35), in contrast with the idea that Hox clusters lack a “closed” conformation whenever genes are silent (23). In ES cells, however, the 4C interaction patterns are much weaker than in brain cells (35), probably because of the presence of bivalent chromatin marks and the concomitant reduced amount of H3K27me3 modifications [Fig. 1 and figure S1 in ref. 35, the latter being even weaker when ES cells were cultured in 2i medium (36)]. It was also noticed that Hoxd genes established many more interactions with the neighboring gene deserts in ES cells than in brain cells (35), where most interactions involved the gene cluster itself, further suggesting a more decompacted configuration of the locus in ES cells. Therefore, the vast majority of ES cells may display decompacted chromatin at their Hox loci. The presence of some H3K27me3-labeled nucleosomes may induce enough transitory interactions to be translated into read counts after deep sequencing of 4C products.

Altogether, these data suggest that in the developing limbs, the HoxD cluster is decompacted, with an elongated shape including both active and inactive genes. Therefore, the coverage of specific parts of the cluster by polycomb complexes, depending on the proximal versus distal position of the cells (8, 16) and observed in other functional contexts (30, 37, 38), does not seem to significantly influence its level of compaction. The distances apparently associated with these elongated forms correspond to previous distance predictions established by mathematical modeling (29). Stretching events associated with transcription were also reported, which seem to occur within this range of elongation lengths (5), and several recent studies either describe or predict distances ranging from 300 to more than 1,000 nm (39, 40). The elongation of the HoxD cluster in limb cells despite its partial labeling by H3K27me3 may reflect an influence from the flanking TADs in extending the structure in both directions, following either productive or constitutive interactions. We evaluated this possibility by using three large inversions modifying the global relationships between the gene cluster and its two regulatory landscapes. However, none of these rearrangements did affect the capacity of the cluster to elongate. If anything, elongation...
In this context, it is noteworthy that the part of the cluster displaying the highest level of decompaction was the Hoxd8 to Hoxd12 region, which precisely matches with the inter-TADs boundary (9, 16). This particular region of HoxD, located between genes strongly and constitutively interacting with either the centromeric (Hoxd13) or the telomeric (Hoxd1, Hoxd4) TADs, may thus display more flexibility, perhaps reflected by the capacity of Hoxd8 or Hoxd12 to switch their contacts from one TAD to the other in various limb cell-types (16). Interestingly, this region is strongly enriched in binding sites for architectural proteins such as CCCTC-binding factor (CTCF) and cohesin (41, 42), which are known for their capacity to organize genomic boundaries and domains, in particular at TAD boundaries (43). Whether the presence of several sites bound by CTCF in the sequence targeted by the Hoxd8 to Hoxd12 fomspid play a role in the elongation observed by STORM remains to be tested, however, as CTCF-driven interactions would be expected to increase compaction, rather than the opposite. An alternative explanation is that the resolution of our STORM approach, although allowing a clear distinction between fully compacted and decompacted Hox clusters, may not easily detect mixed configurations, in particular when only a small part of the cluster differs in shape from the rest.

In conclusion, these data suggest the structural organization of the HoxD gene cluster may predate transcriptional activation and may subsequently be rather independent from its transcriptional status. Perhaps such an elongated structure is maintained in cells expressing subsets of Hox genes, whereas compaction occurs in those cells where genes are silent, such as the fetal brain. In this latter tissue instead, an elongated structure was never observed. By using 2D and 3D FISH, it was previously reported that a decompaction of the HoxD cluster occurs in ES cells that are differentiating in vitro, and that this process is linked to the progressive reduction in levels of PRC1 components (22). With the resolution of our STORM approach, the HoxD cluster did not appear to adopt a compact structure, which would be released on differentiation. It is possible that the spatial architecture of a decondensed cluster differs only slightly between the active and inactive states, leading to the observation of a range of distances between localized probes used for FISH. The presence of polycomb complexes may, for instance, impose such distinct organizations of the HoxD cluster in space, while having little influence on its general level of compaction. In this context, a detailed analysis of these structures by super-resolution microscopy in ES cells mutant for components of either the PRC1 or PRC2 complexes may be informative.

**Materials and Methods**

All experiments involving animals were performed in agreement with the Swiss law on animal protection, with the appropriate legal authorizations to D.D. Tissues were isolated from E10.5 or E12.5 embryos, either wild-type or mutant for the Inv(attP-cd44) (HoxD RVIII -Cd44) inversion (31). Mouse ES cells were grown in serum with 1,000 U/mL leukemia inhibitory factor under feeder-free conditions and G1-synchronized through mitotic shake off. 3D DNA FISH was as in ref. 5. Structured illumination images were acquired using a Nikon structured illumination microscopy setup (Eclipse T1 microscope fitted with a super-resolution Apochromat 100×–1.49 NA objective IXON3 camera; Andor Technology). For STORM imaging, DNA FISH samples were imaged using a UPlanSaPO 100×/1.40 objective (Olympus), and typically 10,000–15,000 snapshot images with a pixel size of 100 nm and an exposure time of 0.03 s were acquired to create one super-resolution (SR) image. SR images were reconstructed with the Octane software (44), and statistical differences between samples were evaluated with the Kruskall-Wallis test, followed by Dunn’s multiple comparison posttest, with the exception of Fig. 3H (Mann–Whitney U test). RNA-Seq was performed according to the TruSeq Stranded Total internal reflection fluorescence 100×–1.49 NA objective IXON3 camera; Andor Technology). For STORM imaging, DNA FISH samples were imaged using a UPlanSaPO 100×/1.40 objective (Olympus), and typically 10,000–15,000 snapshot images with a pixel size of 100 nm and an exposure time of 0.03 s were acquired to create one super-resolution (SR) image. SR images were reconstructed with the Octane software (44), and statistical differences between samples were evaluated with the Kruskall-Wallis test, followed by Dunn’s multiple comparison posttest, with the exception of Fig. 3H (Mann–Whitney U test). RNA-Seq was performed according to the TruSeq Stranded Illumina protocol, with polyA selection. The reads were mapped to Ensemble Mouse assembly National Center for Biotechnology Information Mouse Assembly 37 (mm9) and translated into reads per gene (RPKM) using the Bioinformatics and Biostatistics Core Facility (BBCF), High-Throughput Sequencing station (available at htsstation.epfl.ch). An extended description of the materials, methods, and data analysis is provided in the SI Appendix.
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Title: Nanoscale spatial organization of the HoxD gene cluster in distinct transcriptional states

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Supplemental material

Content

Full methods
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Full Methods

Animal care, tissue sampling

All experiments involving animals were performed in agreement with the Swiss law on animal protection (LPA) with the appropriate legal authorizations to D.D. Tissue samples were isolated at embryonic day 10.5 (E10.5), with day E0.5 being noon on the day of the vaginal plug. E12.5 specimen homozygous for either the Inv(Nsi-Igα6) and Inv(attP-cd44) or E10.5 specimen homozygous for the Inv(HoxD^{RVIII}-Cd44) alleles (Spitz et al., 2005) were obtained by crossing mice trans-heterozygous for each allele. Control E12.5 or E10.5 forebrains were obtained from wild type littermates.

ES cell culture and synchronization

Mouse H1 ES cells (Dierich and Dolle, 1997) were grown on gelatinized plates under feeder-free conditions in Dulbecco’s modified Eagle’s medium (DMEM, Life Technologies) supplemented with 10% fetal calf serum, non-essential amino acids (Life Technologies), Pen–Strep (Life Technologies), Sodium Pyruvate (Life Technologies), 0.1 mM β-mercaptoethanol and 1000 U/ml Leukemia inhibitory factor (LIF). In order to rule out the possibility that the detection of stretched conformations of the HoxD cluster were associated with DNA replication events, we synchronized the ES cells through a mitotic shake off. ES cells were subsequently plated on 18-mm size 1.5 precision coverslips (Carl Roth GmbH), coated with poly-L-lysine.

DNA-FISH

DNA fluorescent in situ hybridization was conducted as previously described (Vieux-Rochas et al., 2015). E10.5 or E12.5 mouse embryos were fixed in 4% paraformaldehyde, embedded in paraffin blocks and cut at 6 µm. Sections were oriented such that cells belonging either to the distal or proximal parts of the growing limb bud could be unambiguously discriminated. ES cells were fixed with 4% paraformaldehyde for 10 minutes. Probes were prepared by nick-translation with either biotin- or digoxigenin-UTP using fosmid and BAC clones obtained from the BACPAC Resources Center (https://bacpac.chori.org/). 100 ng (fosmid) or 200 ng
(BAC) of DNA were used with 7 µg of Cot1-DNA and 10 µg of sonicated salmon sperm DNA.

For the labeling of entire TADs, several consecutive BACs covering most of the TAD sequences were used concomitantly. They were labeled using either digoxigenin- or biotin-dUTP by nick translation with fluorescent revelations as described previously (Morey et al., 2007). TAD-FISH presented in Fig. 1B-G were labeled using either Alexa 647 or Alexa 568 with dye swap to ensure that the changes in ellipticity presented in Fig. 1F was not due to a specific dye. DNA-FISH shown in Fig. 1I-K were labeled using Alexa 647 and Alexa 568 following the conditions presented in Table S3. STORM signals were detected after labelling using either Alexa 647 or Alexa 555 as fluorophores, due to their higher efficiency to detect single molecules blinking signals required for STORM imaging. Slides were stained with DAPI and mounted in ProLong Gold (Life Technologies).

3D Multi-color DNA-FISH imaging

Images shown in Fig. 1B, 1G and 1I were acquired using a 60X Plan-Apo objective (numerical aperture of 1.42) and a B/W CCD ORCA ER B7W Hamamatsu camera associated with an inverted Olympus IX81 microscope. The image stacks with a 200 nm step were saved as TIFF stacks. Image reconstruction and deconvolution were performed using FIJI (NIH, ImageJ v1.47q; http://fiji.sc/Fiji) and Huygens Remote Manager (Scientific Volume Imaging, version 3.0.3).

Distance measurements between probe signals and their ellipticities were determined using an automated spot/surface detection algorithm followed by visual verification and manual correction using IMARIS version 6.5, Bitplane AG and Matlab 7.5, MathWorks SA. For these measurements in standard widefield microscopy, chromatic shift between the two fluorophores were assessed using tetraspeck beads and shown to be lower than a pixel. Statistical significance analyses of distances were performed using an unpaired two-sample t-test and using the Kruskal-Wallis test followed by Dunn’s post test.

Analysis of conformations and morphological parameters

The analysis of TAD-FISH in Fig. 1C shows the distribution of the different conformations observed (spaced, contact or partial overlap), was conducted by three
different observers, under blind conditions. The analysis shown in figure S1E was also blinded, with only two conformations scored: either segregated or mixed.

The analyses of shapes were defined using three main parameters: aspect ratio, circularity and ellipticity. They were assessed after the replacement of their area selection with the best fit ellipse. Aspect ratio and circularity were quantified using Fiji (NIH, ImageJ v1.47q). Circularity is defined as

$$\frac{4\pi \times \text{Area}}{\text{Perimeter}^2}$$

with the maximal value (1) indicating a perfect circle. As the value approaches 0.0, it indicates an increasingly elongated shape. The aspect ratio was calculated as a ratio between major and minor axes of ellipse fitted to the shape of the cluster. It was is defines as

$$\frac{\text{Major axis}}{\text{Minor axis}}$$

The ellipticity was defined as

$$\frac{\text{Major axis- minor axis}}{\text{Major axis}}$$

with the maximal value, 1.0, indicating a perfect ellipse. It was scored using the IMARIS (v6.5, Bitplane AG) software and quantified automatically. The dispersion ratio shown in figure 3D is the measure of the number of objects observed for each cluster with the lowest value 1.0, indicates a single object.

**SIM Imaging**

Structured illumination images were acquired using a Nikon SIM setup (Eclipse T1 microscope fitted with a super-resolution Apochromat total internal reflection fluorescence 100×/ 1.49 NA objective and an electron-multiplying charge-coupled device camera (IXON3; Andor Technology).

Imaging was performed in 3D SIM acquisition mode with 3 dyes (DAPI, Alexa 568 and Alexa 647), which were excited with fixed wavelength using coherent Cube at 100mW (for 402nm and 561 excitations) and a coherent Jive at 158mW (for 641nm excitation). Image reconstruction was performed using the NIS-Elements software (Nikon; based on Gustafsson et al., 2008) and reconstruction parameters were as follows: Contrast: 0.70, apodization: 1.00 and Widh3DFilter: 0.20.
STORM imaging

STORM imaging was performed on a modified Olympus IX-71 inverted microscope equipped with an oil objective (Olympus, UPlanSApo 100x, 1.40 NA) and a piezo objective scanner (Physik Instrumente, P-725, PIFOC). Fluorescence was excited by a 641 nm laser (Coherent, Cube 640nm-100C) with an irradiance of 10-20 kW cm⁻². For photoreactivation we also used 405 nm (Coherent, Cube 405nm-100C) laser with an irradiance < 0.05 kW cm⁻². Emitted light was directed through a dichroic mirror (89100BS, Chroma) and emission filter (ET700/75m, Chroma) and imaged by an EMCCD camera (Andor Technology, iXon+).

To induce photoswitching, samples were imaged in GLOX buffer (0.5 mg.ml⁻¹ glucose oxidase (Sigma-Aldrich G0543-10KU)), 40 mg ml⁻¹ catalase (Sigma-Aldrich C3515) and 10% glucose (Sigma-Aldrich G8270). For each imaging session, GLOX buffer was freshly prepared, then after 30 min of equilibration, β-mercaptoethanol (Sigma-Aldrich M6250) was added for a final concentration of 143 µM. The pH of the solution was adjusted to 8-8.5 using HEPES (Sigma-Aldrich H3662). Stacks of 10000-15000 images with a pixel size of 100 nm with exposure time 0.03 s were acquired to create each STORM image.

STORM Data analysis

Initially, HoxD clusters were identified using standard deviation (STD) images from the raw image stacks. Each pixel value in the STD image was determined by calculating the standard deviation of intensity values across the image stack. Spots with a signal approximately 10x higher than other objects within the nucleus in STD images were considered as labeled clusters; this first segmentation step enabled us to avoid analyzing non-specific staining (Fig. S3A). After this step, we always obtained less than or equal to 4 bright spots per cell, consistent with the maximum expected number of clusters per cell. In addition, we performed several multicolor standard microscopy experiments with probes used for SR imaging. These experiments confirmed the specificity of staining. Probes that were close to each other on genome were also close to each other in physical space in cells (Fig. S3).

Octane software was used to analyze raw image stacks and reconstruct STORM images, which were later used for morphological analysis. During image reconstruction, the single molecule peak detection threshold was set individually for
each sample set, due to variation in image background intensity. The threshold was chosen using the following qualitative criteria. We required that the majority of bright fluorescent spots corresponding to single molecules that we manually detected based of their shape and size in the cluster region, were also detected by the software. This procedure is illustrated in Fig. S3.

To account for molecules that are fluorescent for several frames, we performed grouping. Localizations were grouped as belonging to the same molecule if the distance between them in space was less than 100 nm (grouping radius) and in time less than two frames (grouping gap). These parameters allowed accounting for most of the molecules that are on for several frames. Octane software allows visual inspection of grouped localizations so we checked that grouping was done properly by manually inspecting grouped localizations on test samples for each dataset. The choice of grouping parameters is further discussed (Fig. S4).

Typical precision in position determination of the molecule (localization precision) that we obtained in our experiments was around 40 nm (as measured with Peakselector software). Based on this we set a pixel size of 10 nm in reconstructed STORM images to slightly oversample the data. STORM images were rendered in the following way: each localized molecule was represented as a spot whose width reflected the mean localization precision for the dataset, usually 40 nm. We found that moderately changing these two rendering parameters (pixel size and spot size) didn’t change the visual appearance or measured geometric parameters of cluster (Fig. S5).

The shape of identified clusters was determined using thresholding of rendered STORM images. The thresholding was performed with Otsu’s algorithm integrated into Fiji software (http://fiji.sc/Fiji). The algorithm sets a threshold by minimizing a weighted sum of variances of the background class and the image class pixels. The results of Otsu algorithm are demonstrated (Fig. S6). The resulting thresholded image then was used to determine the aspect ratio of clusters. The aspect ratio was calculated as a ratio between major and minor axes of ellipse fitted to the shape of the cluster. To evaluate statistical differences between samples, we used the two-sample t-test for means.

It should be noted that the ellipticities we measured are an underestimation of the true elongation of the *HoxD* clusters. This is because we performed 2D imaging to avoid distortion of clusters in the axial dimension, an effect that arises due to the poorer localization precision in z. Therefore, our images represent a projection of 3D
clusters, which can be oriented with different angles with respect to the 2D imaging plane (Fig. S8). If we assume that an elongated object is randomly oriented in space we can estimate in the simplest case that the average real 3D elongation is \((\pi/2) - 1.57\) larger than the average measured 2D elongation.

**RNA-Sequencing**

E12.5 distal forelimbs were dissected and isolated using Trizol LS reagent (Life Technologies) to generate total RNA tissue samples. RNA-Seq was performed according to the TruSeq Stranded Illumina protocol, with polyA selection. The strand-specific total RNA-seq libraries were constructed according to the manufacturers instructions (Illumina). Sequencing was done using 100 bp Single end reads on the Illumina HiSeq system according to the manufacturer's specifications. RNA-seq reads were mapped to ENSEMBL Mouse assembly NCBIM37 (mm9) and translated into reads per gene (RPKM) using the RNA-Seq pipeline of the Bioinformatics and Biostatistics Core Facility (BBCF) HTSstation (available at http://htsstation.epfl.ch). RNA-seq data is available from the Gene Expression Omnibus (GEO) repository under accession number GSE72285.

**References**


Supplemental figures and legends

Supplemental Figure S1

Figure S1: Overlapping capacity between probes in and out of the TAD units.
(A) Schematic of the probe positions as shown in Fig. 1 with the combinations (1 to 4) used for the 3D-DNA FISH. The dotted line represents the TAD boundary as defined in (9). (B) Overlaps were assessed using reconstructed 3D image using Imaris software. The picture is an orthogonal view of a slice showing a combination of BACs (condition 1) that appear overlapping in 2D but were scored segregated when visualized in three dimensions. Scale bar: 500nm (C) Each image shows the extent of overlap observed for each combination tested. Scale bar: 200nm. (D) Quantification of the distances between two probes for each of the combination tested. For each combination is shown the average distance in three different embryonic tissue: E12.5 forebrain (dark grey), proximal forelimb (light grey) and distal forelimb (white). N>100 for each pair including data from 2 different experiments (conditions 1 to 3) or one experiment (condition 4). Error bars denote SD. (E) Blinded and qualitative measurement to show the percentage of pairs (70 to 100 pairs) that overlap. As shown in (B) the pairs were not scored as mixed if they were segregated in 3D. The errors bars for conditions 1 to 3 denote SD (n=2 for each). The measurements for condition 4 came from the same animal (n=1).
Supplemental Figure S2

A

<table>
<thead>
<tr>
<th>ES Cells</th>
<th>Forelimb</th>
<th>Forebrain</th>
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B

<table>
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<tr>
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<th>HoxD</th>
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</table>

Figure S2: Diversity of *HoxD* morphologies in different cell types. The images illustrate the variability in the shape of objects observed after DNA-FISH using a full *HoxD* cluster probe, followed by STORM imaging. Variability is particularly obvious when looking at ES cells, even though compact structures are not scored (upper panels). In forelimb cells (middle panels), elongated structures are preponderant whereas forebrain cells (lower panels) generally display a much more compact signal. Scale bar: 200nm. (B) Two-colors DNA-FISH showing various configurations of either compacted (left) or de-compacted (right) part of the *HoxD* cluster (scheme in the left panel).
Figure S3: Impact of the two flanking TADs upon condensation of the HoxD cluster. (A) Schematics of the three large targeted inversions used in this work as well as of the positions of the BAC and fosmid clones used to monitor elongation (below). The Inv(Nsi-Itga6) separates the HoxD cluster from its centromeric TAD, whereas the Inv(attP-cd44) separates the cluster from its telomeric TAD. The Inv(cd44v7-HoxD^RVIII) splits the HoxD cluster into two small clusters of genes (Spitz et al., 2005). (B) STORM-resolved images obtained with the BAC clone covering the HoxD cluster in E12.5 distal forelimbs showed similar de-compacted aspects for both control and inverted alleles in the 3Mb large centromeric (Inv(Nsi-Itga6)) inversion. (C) Both aspect ratio and circularity index also remained unchanged for the signals obtained with the Hoxd11 to Hoxd13 probe when using the 28Mb large Inv(cd44v7-HoxD^RVIII) telomeric inversion in E10.5 posterior trunk cells. n.s.: not significant when using Mann-Whitney U test.
Supplemental Figure S4

Figure S4: Single molecule peak detection. (A) Standard deviation image of forebrain tissue labeled with HoxD cluster probe; bright spots correspond to HoxD clusters (scale bar 1µm). (B) Single STORM imaging frame. (C) Fluorescent image of two typical single molecules and their fluorescence intensity profiles. (D) Threshold was chosen individually for each image by checking that most of single molecules are detected by software (yellow cross indicates that molecule is detected).

Supplemental Figure S5

Figure S5: Grouping procedure details. (A) Some molecules are fluorescent for several frames and therefore a grouping correction is needed. (B) Distribution of distances between subsequent localizations of grouped molecules. We note that by using a grouping radius of 100 nm we obtain a non-truncated distribution of step sizes between frames with a mean value far below the maximum search radius, which validates our choice. (C) Number of molecules (after grouping) as a function of grouping gap decays with time but does not plateau. This makes it difficult to find an optimum grouping gap. On one hand it is desirable to introduce at least a gap of one frame in grouping to account for molecules that have borderline intensity and may be
not detected in some frames. On the other hand this gap should not be too long as it can lead to false connection of different molecules. Based on these considerations we picked grouping gap of one frame. As it is stated in the methods, we systematically checked manually that grouping was performed properly.

**Supplemental Figure S6**

![Supplemental Figure S6](image)

**Figure S6: Different rendering parameters do not affect geometrical parameters of a cluster.** STORM image of a HoxD cluster rendered with different pixel size and different spot size for a single molecule (scale bar 100nm); calculated aspect ratio presented for each rendering condition.

**Supplemental Figure S7**

![Supplemental Figure S7](image)

**Figure S7: Results of Otsu’s algorithm based thresholding.** Original image of a HoxD cluster (left) and thresholded image (right) (scale bar 100nm).
Supplemental Figure S8

Figure S8: Measured elongation is an underestimation of real elongation. Here, a spheroid with major axis $a_{\text{real}}$ and two other axes ($b_{\text{real}} = c_{\text{real}}$) is imaged (imaging plane XY) as an ellipse with major axis $a_{\text{registered}}$ (in this configuration $b_{\text{registered}} = b_{\text{real}}$) so that the average true 3D elongation ($\pi/2$) $\sim 1.57$ larger than the average measured 2D elongation.
Table S1. List of BAC and fosmid clones used for DNA-FISH in the present study. Genomic coordinates are given for mm9.

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Table S2. Statistical significance between the different conditions of interprobe distances presented in Figure 1J. The asterisk * denotes p<0.05, ** for p<0.01 and *** when p<0.001.

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**Table S3.** Combinations of fluorophores used for 3D DNA-FISH shown in Fig. 1H-K

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