Supplementary Material

Supplementary Figure 1

Genomic Southern blot showing the hybridization results for the Chr1 Dig-labeled probe. The blot consists of human genomic DNA digested with *Eco*RI before separation on a 0.8% agarose gel. DNA size based on the marker lane is shown to the right. The unique 7.3kb band corresponding to a single repeat monomer is indicated to the left. The blot was hybridized with the DIG-labeled probe for 3 hours at 60°C in ExpressHyb (Clontech Laboratories Inc., Mountain View, CA, USA), before washing at 60°C for 8 minutes twice with wash one (2x SSC, 0.1% SDS) and twice with wash two (0.2x SSC, 0.1% SDS). The signal was detected using the DIG High Prime DNA Labeling and Detection Starter Kit II according to the manufacturers instructions (Roche Applied Science, Indianapolis, IN, USA).

Chr1 Probe Specificity Check

-12.0kb -8.0kb -6.0kb -4.0kb

-2.0kb

-1.0kb

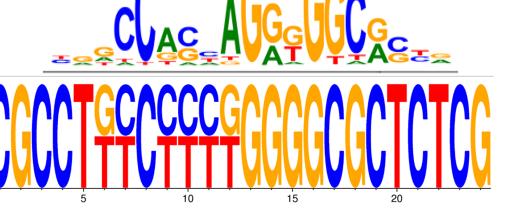
- 0.5kb

Supplementary Figure 2

Comparison of the consensus CTCF binding motif obtained adapted from JASPAR (100), to a probability map of the conserved human and mouse sequence from the tDNA repeat that underlies the human CTCF ChIP-Seq peaks. The probability plot was generated using WebLogo (101), and used the reverse complement of the human and mouse sequences.

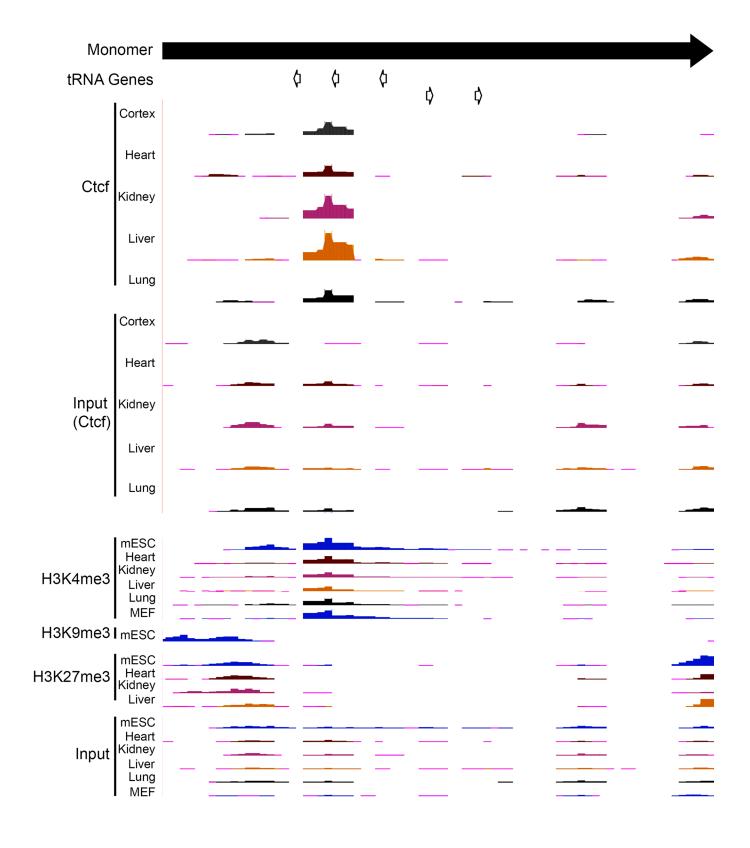
Jaspar Core

Human & Mouse tDNA



Supplementary Figure 3

Chromatin features of the mouse tDNA repeat. The black arrow represents a single monomer at the top of the image, under which the open arrows indicate the location of the tRNA genes. Below this are ChIP-Seq and input control profiles for various mouse cell and tissue types adapted from the UCSC Genome Browser (http://genome.ucsc.edu/index.html).



Supplementary Figure 4

Schematic map showing the relationship of the Human and Mouse tDNA tandem repeats (right-facing black arrows) relative to topological domains, represented by the green and red genomic compartments (93). The image was adapted from the HiC Project at the Ren Lab (http://yuelab.org/hi-c/). White arrows indicate the location of transcripts from the interval, oriented in the direction of transcription and covering the approximate genomic span of the corresponding gene. Human data is shown for human embryonic stem cells (hESC) and lung fibroblast IMR90, and corresponds to 159,520,867-159,880,006 of human chromosome 1 (build hg18). Mouse data is shown for mouse embryonic stem cells (mESC) and mouse cortex, and corresponds to 172,880,561-173,119,754 of mouse chromosome 1 (build mm9).



