

# DNA sequence of human chromosome 17 and analysis of rearrangement in the human lineage

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**Chromosome 17 is unusual among the human chromosomes in many respects. It is the largest human autosome with orthology to only a single mouse chromosome<sup>1</sup>, mapping entirely to the distal half of mouse chromosome 11. Chromosome 17 is rich in protein-coding genes, having the second highest gene density in the genome<sup>2,3</sup>. It is also enriched in segmental duplications, ranking third in density among the autosomes<sup>4</sup>. Here we report a finished sequence for human chromosome 17, as well as a structural comparison with the finished sequence for mouse chromosome 11, the first finished mouse chromosome. Comparison of the orthologous regions reveals striking differences. In contrast to the typical pattern seen in mammalian evolution<sup>5,6</sup>, the human sequence has undergone extensive intrachromosomal rearrangement, whereas the mouse sequence has been remarkably stable. Moreover, although the human sequence has a high density of segmental duplication, the mouse sequence has a very low density. Notably, these segmental duplications correspond closely to the sites of structural rearrangement, demonstrating a link between duplication and rearrangement. Examination of the main classes of duplicated segments provides insight into the dynamics underlying expansion of chromosome-specific, low-copy repeats in the human genome.**

Human chromosome 17 is implicated in a wide range of human genetic diseases. It is home to genes involved in early-onset breast cancer (*BRCA1*), neurofibromatosis (*NF1*) and the DNA damage response (*TP53* encoding the p53 protein). The complex rearrangement and duplication structure of chromosome 17 predisposes it to non-allelic homologous recombination (NAHR), resulting in DNA rearrangements that cause several well-studied microdeletion disorders. These include hereditary neuropathy with pressure palsies

(HNPP)<sup>7</sup> at 17p12, and Smith–Magenis syndrome (SMS) deletions at 17p11.2 (refs 8, 9). Microduplication counterparts for both of these conditions are also known, famously in the case of Charcot–Marie–Tooth disease type 1A (CMT1A) at 17p12 (refs 10, 11), the most common inherited peripheral neuropathy (see ref. 12 for a review), and more recently for SMS<sup>13</sup>. Further, the complex architecture in this region, consisting of ~50-kb subunits of direct and inverted repeats able to form cruciform structures, is also responsible for susceptibility to one of the most common somatic rearrangement events characterized, isodicentric 17q, which is associated with several cancers and signifies poor prognosis<sup>14</sup>. Finally, all cases of Miller–Dieker syndrome<sup>12</sup> and many cases of isolated lissencephaly sequence<sup>12</sup> are associated with haploinsufficiency effects of deletions at 17p13.3.

As part of the Human Genome Project<sup>2</sup>, we generated a finished sequence of human chromosome 17, which comprises ~2.8% of the euchromatic genome. The finished sequence contains 78,839,971 bases and is interrupted by nine euchromatic gaps and one gap containing the centromere region (Fig. 1). The total size of the euchromatic gaps is estimated at 854 kb (see Methods and Supplementary Table S1). The frequency and size of gaps is slightly higher than the human genome average<sup>2</sup>. (It is also significantly higher than for mouse chromosome 11; see below.) This is largely due to segmentally duplicated or unclonable regions on human chromosome 17 (see Supplementary Information). A clone list for the most recent version of the sequence is provided in Supplementary Table S2.

An overview of chromosome 17 is shown in Fig. 1. The chromosome has features characteristic of gene-rich regions<sup>2</sup>, including a high average G+C content (45.5%) and a collection of transposable

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element fossils (45.3% overall) with a relative excess of short interspersed elements (SINEs, 22.3%) and a deficit of long interspersed elements (LINEs, 14.4%). It also has a large excess of segmental duplications (defined as having >90% identity and being >1 kb in length<sup>15</sup>). Such segmental duplications constitute 8.6% (7 Mb) of the finished sequence, in contrast to a median of 3.9% for human chromosomes and an average of 4.3% across autosomal sequences. As with other duplication-rich chromosomes, chromosome 17 has a large proportion of intrachromosomal duplication: 62.3% of the duplicated sequence is strictly intrachromosomal, 17.4% is both intra- and inter-chromosomal, and 20.3% is solely interchromosomal. We elaborate below on the segmental duplication content of the chromosome.

We have produced a manually curated catalogue of protein-coding genes on chromosome 17 (see Methods), annotating 1,266 gene loci and 274 pseudogene loci (Supplementary Table S3). This catalogue includes all of the 1,079 genes on the chromosome in the RefSeq database (see Methods and Supplementary Information). The gene density (16.2 genes per Mb) is substantially higher than the genome average<sup>2</sup> and ranks only behind chromosome 19 (26.2 genes per Mb; ref. 3). There is evidence of extensive alternative splicing, with gene loci having an average of five distinct transcripts, and 76.6% having at least two transcripts, a proportion comparable to recent reports<sup>3,16–19</sup>. Of the 274 pseudogenes on the chromosome, 73.0% are processed

(intronless) pseudogenes arising from retroposition. In addition to protein-coding genes, we identified 42 transfer RNA genes on the chromosome (Supplementary Table S4).

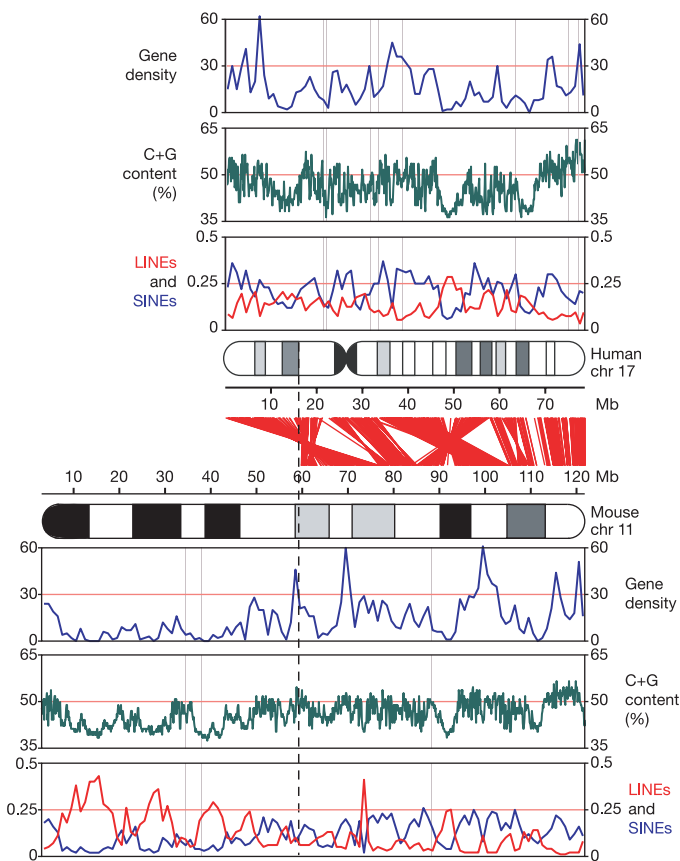
As part of the Mouse Genome Sequencing Consortium (MGSC), a finished sequence of mouse chromosome 11 has been produced (Supplementary Table S5). The sequence was derived from the C57BL/6J strain and has a finished euchromatic length of 118,816,080 bases, representing ~4.6% of the mouse genome. The finished sequence is interrupted by only two euchromatic gaps, with an estimated total size of 97 kb, and one additional gap containing the acrocentric heterochromatin (Fig. 1, Supplementary Table S1 and Methods). See Supplementary Table S5 for a clone list of the most up-to-date version of mouse chromosome 11.

Mouse chromosome 11 can be divided into two parts on the basis of conserved synteny with human chromosomes (Fig. 1). Although the proximal 59 Mb shows conserved synteny with five different human chromosomes (1, 2, 5, 7 and 22), the distal ~60 Mb has conserved synteny exclusively with human chromosome 17 (see Fig. 1 and Supplementary Fig. S1). The distal portion of mouse chromosome 11 is similar to human chromosome 17 in various respects, including having a high G+C content (45.5%), high gene density (18.7 genes per Mb), an excess of SINEs (14.4%), and a deficit of LINEs (8.2%). The ratio of SINEs to LINEs is similar to that in human, although the absolute density is lower, consistent with the fraction of detectable transposon sequence in mouse. In contrast, the proximal portion of mouse chromosome 11 has a lower G+C content (41.9%), low gene density (8.2 genes per Mb), and an opposite ratio of SINEs and LINEs (7.8% versus 19.1%). Although it is similar to human chromosome 17 in most other respects, the distal region of mouse chromosome 11 differs sharply with respect to segmental duplications: such sequences constitute only 1.1% of the distal region. Indeed, mouse chromosome 11 has the lowest rate of segmental duplication (1.4%) among all mouse chromosomes.

The block of undirected synteny (corresponding regions of orthology between the species, in which order may be disrupted by internal rearrangements) between human chromosome 17 and mouse chromosome 11 is the second largest such autosomal block between human and mouse (see Supplementary Information). Within this block, there are 23 segments of directed (collinear) synteny of >100 kb, indicating frequent inversion. To better understand this relationship, we reconstructed the chromosomal structure of the primate–rodent ancestor of human chromosome 17 (Supplementary Fig. S2) on the basis of genome sequences from human, mouse, dog and opossum (see Supplementary Information). The primate–rodent ancestor could be resolved except for one small segment, and also represents the state of the boreoeutherian ancestor except in two regions of conserved directed synteny between human and mouse that differ from dog and opossum (see Supplementary Information and Supplementary Fig. S2).

Comparison of the modern chromosomes with our ancestral reconstruction (Fig. 2a) shows that the mouse structure is virtually unchanged, whereas significant rearrangements have occurred in human. There are 20 rearrangement breakpoints in human but only three in mouse. (The terminal break and one internal break are used for different events in both species, leading to only 22 observed differences in the map of direct human–mouse conserved synteny; see Supplementary Fig S1.) This is contrary to the general pattern for the human and mouse genomes, which typically shows many more rearrangements along the rodent lineage<sup>5,6</sup> (see Supplementary Information). The mouse-specific rearrangements are confined to a single region orthologous to the Smith–Magenis region in human, which shows an even more complex pattern of rearrangements at finer scale (Fig. 2b).

What accounts for the extreme rearrangement of human chromosome 17? The answer seems to be related to the extensive segmental duplication on the human chromosome. The human-specific breaks



**Figure 1 | Landscapes of human chromosome 17 and mouse chromosome 11.** Approximate alignments of ideograms of the two chromosomes are shown in the centre of the figure, with red lines showing the relationships between orthologous genes. From top to bottom for each organism, tracks show gene density (in genes per Mb), G+C content (on a scale from 35–65%), and densities of LINEs (red) and SINEs (blue) (fraction of bases). The vertical dashed line represents the approximate boundary between the distal region of mouse chromosome 11, which has shared synteny with human chromosome 17, and the proximal region, which does not. Chr, chromosome.

in conserved synteny are highly correlated with regions of intrachromosomal duplication (Fig. 3). Roughly 74% of the duplicated bases reside in the breakpoint regions between stretches of conserved synteny, which comprise only 7.3% of the bases in the chromosome sequence.

To further explore the nature of these duplicated sequences, we clustered the duplcons into classes on the basis of shared sequence (see Methods). After clustering, most of the duplicated sequences fall into one of three classes (1–3), which account for 51.6%, 19.9% and 3.1%, respectively, of the intrachromosomal duplications (Fig. 3 and Table 1). For each duplication class, we then sought to define a 'core element', corresponding to the maximum length of sequence occurring in the largest number of duplcons (see Methods). These core elements can serve as useful markers for identifying class members and may provide insight into the origins of the duplications<sup>20</sup>.

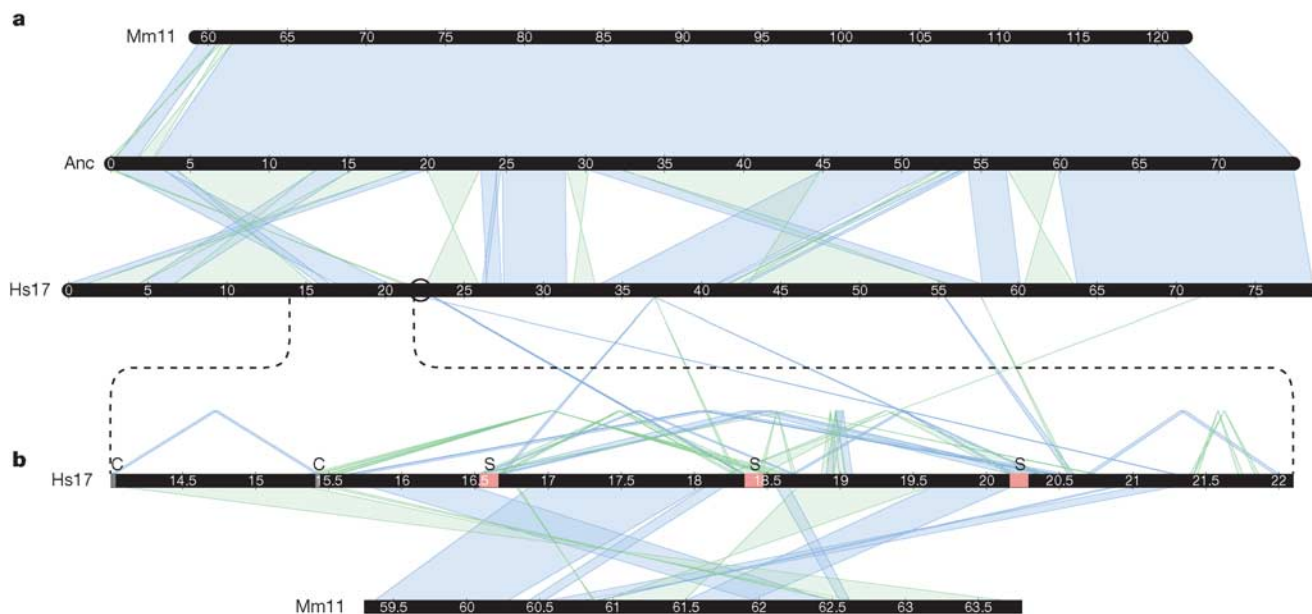
The class 1 core element (~11 kb in length) contains a TBC1 domain and is found in 12 of the 19 human-specific breakpoint regions. Several of the class 1 core elements are actively transcribed, including six very recently duplicated genes from the TBC1D3 family (at 31–33 Mb) and the *USP6* oncogene (also known as *TRE-2*), which is a fusion of a TBC1D3 gene and the *USP32* gene<sup>21</sup>. The TBC1D3 family genes encode proteins with TBC1 domains, and are believed to be GTPase-activating proteins specific to *RAB5*, a RAS-related gene involved in vesicle trafficking. The TBC1D3 gene family has no known homologues outside the primates, and we could find no similar sequence in dog or mouse, making it impossible to identify an ancestral copy. However, the element certainly pre-dates the primate clade: human copies show substantial divergence (~34%) (Supplementary Fig. S3) and copies can be identified in ring-tailed lemur by polymerase chain reaction (PCR) (see Methods). The element is also duplicated in macaque, and a phylogeny of the elements in human and macaque (Supplementary Fig. S3) shows that at least some of these duplications pre-date the divergence of Old World monkeys, as some macaque copies have a human copy as their nearest neighbour.

The class 2 core element (~4 kb in length) is found in 9 of the 19

human-specific breakpoint regions (Fig. 3). The ancestral copy is probably the copy immediately upstream of the *NSF* gene at 41.95 Mb, as this is the only copy with an orthologous sequence in mouse. The class 2 duplcons occur primarily in three regions of chromosome 17q, at 26–27 Mb, 40–42 Mb and 60–63 Mb. These regions are all sites of extensive, small local inversion events that have occurred since the primate–rodent ancestor. A phylogenetic tree of the core elements (Supplementary Fig. S4) shows two main clusters, one for the copies at 26–27 Mb and another for those at 40–42 and 60–63 Mb. Notably, the reconstruction of the ancestral chromosome order (Supplementary Fig. S2) places these latter regions nearly adjacent to one another, suggesting that the core elements dispersed in a local region that was then disrupted by an inversion with breakpoints at 33 and 57 Mb. This ancestral chromosomal order seems to correspond to the modern karyotype of orangutan<sup>22</sup>, which has a paracentric inversion with respect to the human karyotype on 17q, with nearly coincident breakpoints. This would imply that the inversion in the human lineage is a fairly recent event. Notably, two class 2 core elements (the ancestral copy and a copy at 40.97 Mb) flank the boundaries of a 900-kb polymorphic inversion in the human population<sup>23</sup> and also the boundaries of a human rearrangement with respect to the ancestral chromosome (Fig. 2 and Supplementary Fig. S2), indicating a reuse of these breakpoints within the human lineage (see Supplementary Information).

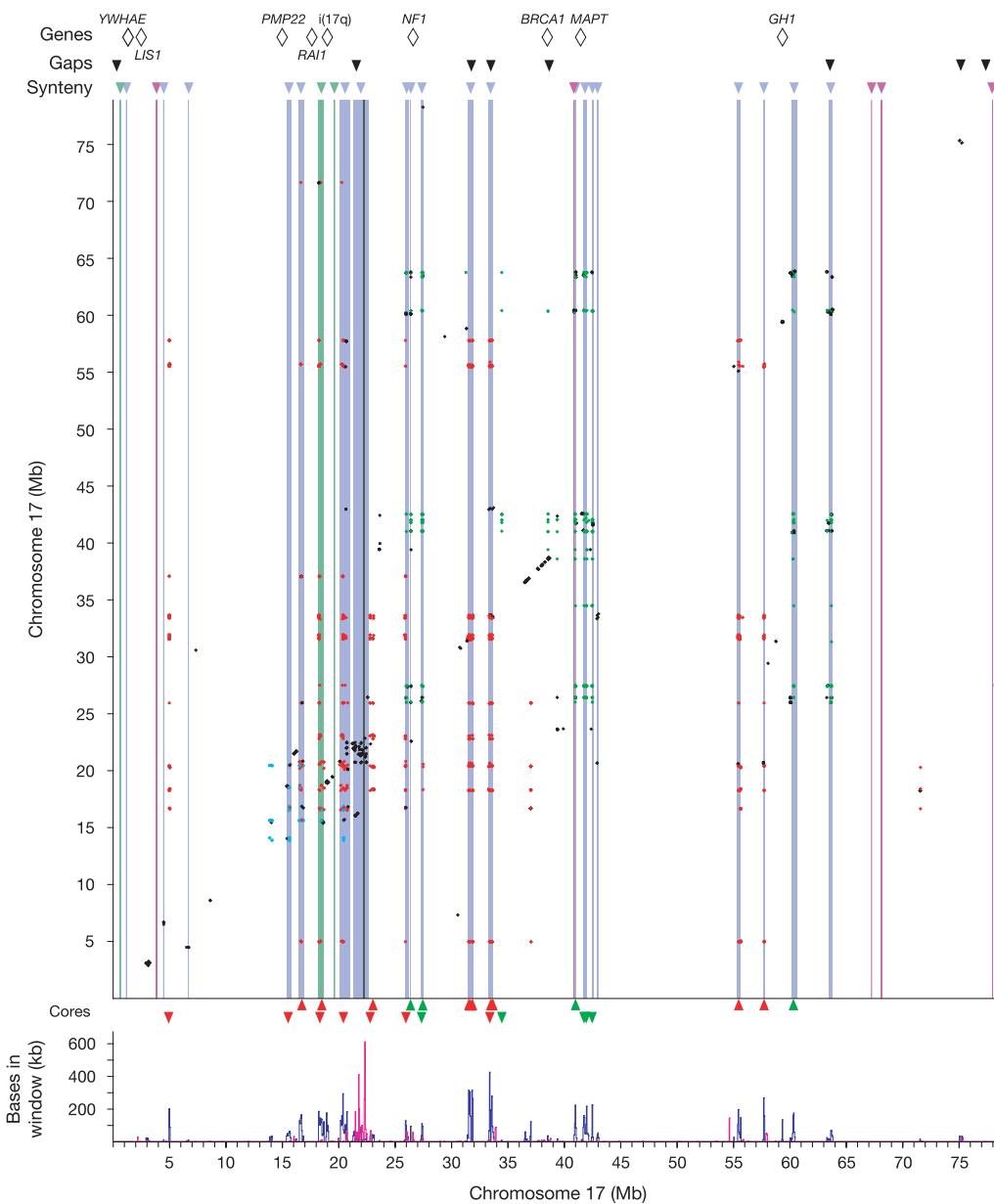
The class 3 duplcons all reside in the CMT1A/SMS region (13.8–20.4 Mb), which seems to be a region of remarkable genomic fragility. The class is too small to define a meaningful core element, but examination of the longest copy shows that it is a previously identified low-copy repeat, LCRA1<sup>10</sup>. This element is the result of a fusion of two distinct 3'-UTR elements from elsewhere in the human genome. Although at least one of the individual sub-elements comprising the fusion is found in mouse and dog, the fusion product is not seen in these species, but only in primates. One of the two sub-elements is actively transcribed (see Supplementary Information).

Previous studies<sup>8,10</sup> of the CMT1A/SMS region have identified large, highly similar local repeat blocks. The blocks are themselves



**Figure 2 | Syntenic relationship between mouse, human and the ancestral chromosome.** **a**, Conserved syntenic blocks of 100 kb or more between human chromosome 17 (Hs17), mouse chromosome 11 (Mm11), and a most parsimonious primate–rodent ancestor (Anc) reconstructed using dog and opossum as outgroups. **b**, An enlargement of the CMT1A/SMS region and the mouse orthologous sequence (human 14–22 Mb, mouse 59–64 Mb), including conserved syntenic blocks of 10 kb or more between human and mouse, and human segmental duplications of 20 kb or more. Segmental

duplications are shown above the human CMT1A/SMS line. Duplications with copies outside the enlarged region connect to 17q in **a**. Coloured blocks marked 'C' represent CMT1A duplicons, and those marked 'S' represent SMS duplcons. In both sections, direct (reference strand) blocks of conserved synteny or segmental duplication are shown in blue; inverted blocks are shown in green. Chromosome blocks are labelled in megabases. The black circle indicates the location of the human chromosome 17 centromere.



**Figure 3 | Duplication landscape of chromosome 17 and its association with breaks in conserved synteny.**

The top line shows genes/breakpoints of interest for rearrangements mentioned in the text. Black triangles show the locations of the nine remaining gaps in the build. Below that, coloured triangles show breaks in synteny (blue for human, green for mouse, pink for dog). Intrachromosomal duplications are shown as an all-versus-all dot plot, coloured by class (class 1 in red, class 2 in green, class 3 in blue, all other classes in black; see Methods). Breaks in conserved synteny are represented as vertical bands coloured as above. The centromere is shown as a vertical dark grey band. Triangles at the bottom show core element locations (class 1 core elements in red, class 2 core elements in green), with direction showing orientation (up, forward strand; down, reverse strand). The panel at the bottom shows a density plot of intrachromosomal (blue) and interchromosomal (red) duplication, using 50-kb non-overlapping sliding windows.

fusions of multiple duplons (including class 1 core elements) that do not occur together elsewhere in the genome. None of these duplicated sequences has a mouse orthologue within the region (Fig. 2b), having all come from the q arm of 17, indicating that this region may be a site of multiple inversion. Indeed, cytogenetic mapping of primate rearrangements<sup>22</sup> shows the CMT1A/SMS region to be the site of an inversion breakpoint in orangutan and of a translocation between chromosomes 17 and 5 in gorilla<sup>24</sup>.

Certain common characteristics emerge from the study of segmental duplication on chromosome 17, as well as from our recent study of segmental duplications on chromosome 15 (ref. 20). First,

the vast majority of duplons in these classes fall in breaks of conserved synteny. Second, segmentally duplicated sequence seems to spread readily within local regions. Local clustering is evident for all duplton classes studied on chromosomes 15 and 17. Third, segmentally duplicated sequence then seems to spread over larger distances by chromosomal inversions. In two cases (class 2 on chromosome 17 and the main class on chromosome 15; ref. 20), such dispersal is confirmed on the basis of both sequence phylogeny and reconstruction of ancestral chromosome structure. With additional comparative data on lower primates, it should be possible to reconstruct the dispersal of most of the segmentally duplicated

**Table 1 | Duplication classes on human chromosome 17**

Class	No. of duplications	No. of duplications containing core element	No. of duplications in syntenic break regions	Core element size (bp)	Total bases in duplons	Duplton bases in syntenic break regions	Total duplton bases in syntenic break regions (%)
Class 1	529	225	509	10,734	1,837,890	1,581,557	86.05
Class 2	204	65	202	4,185	979,083	735,413	75.11
Class 3	32	0	30	NA	125,758	105,741	84.08
Other (132)*	260	0	196	NA	2,449,945	1,574,952	64.29

NA, not applicable.

\* Each of these 132 duplton classes has fewer than 15 events.

sequences in the human genome. Finally, the core elements underlying the main classes of segmentally duplicated sequence on chromosomes 15 and 17 all show active transcription from a large number of the duplicons<sup>10,25,26</sup>. This observation suggests a link between active transcription, duplication and rearrangement. It is possible that the nature of transcription, the chromatin structure or the transcripts at these loci may render the regions more susceptible to breakage, duplication or non-allelic recombination.

Our comparison of human chromosome 17 and distal mouse chromosome 11 shows that a genomic region can undergo strikingly different rearrangements in different lineages. There has been remarkably little interchromosomal rearrangement in the region orthologous to human chromosome 17 across all mammals, which may perhaps reflect an inherent stability of the region or a selective constraint on contiguity. In contrast, the region shows extensive internal rearrangement in the human lineage (20 breakpoints relative to the ancestral genome) but little rearrangement in mouse (only three rearrangements relative to the ancestor). The occurrence of the human breakpoints near segmental duplications supports a mechanistic link between these two features that may explain the unusual history of this genomic region in the primate lineage. Together with our recent analysis of chromosome 15, this analysis sheds light on the forces that shape the large-scale structure of genomes and, in the process, create both evolutionary variation and the chromosomal fragility underlying human disease.

## METHODS

The finished sequence of human chromosome 17 was generated almost entirely (>99%) at the Broad Institute of MIT and Harvard (formerly the Whitehead Institute/MIT Center for Genome Research) (Supplementary Table S6). Chromosomal positions in this paper refer to NCBI human build 35. The finished sequence of mouse chromosome 11 was generated almost entirely (>99%) at the Sanger Institute (Supplementary Table S6). Chromosomal positions in this paper refer to NCBI mouse build 34. Methods for clone mapping, sequencing and tiling-path validation are described in the Supplementary Information. The gene catalogue and annotation for human chromosome 17 were produced as previously published<sup>16</sup>.

**Syntenic maps and segmental duplications.** Syntenic maps and segmental duplications were constructed as previously described<sup>15,16,20</sup>.

**Duplication class clustering.** Pairwise duplications were clustered by physical overlap of  $\geq 150$  bp and at least 5% of the smaller element. We extended this by transitive closure to build maximally linked sets (see Supplementary Information).

**Identification of core elements of duplicons.** For each duplicon class, aligned coverage of each base in the duplicated region (the union of all duplicons) was recorded. Core elements were chosen by visual inspection to identify the longest contiguous sequence in the most deeply covered region. For each duplicon class, the longest core element sequence was aligned to the entire genome with PatternHunter<sup>27</sup>, and all full-length alignments were identified as core element copies.

**Construction of core element phylogeny.** Construction of core element phylogeny was performed as previously described<sup>20</sup>.

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Supplementary Information is linked to the online version of the paper at [www.nature.com/nature](http://www.nature.com/nature).

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**Author Information** Accession numbers for all clones contributing to the finished sequence of human chromosome 17 can be found in Supplementary Table S2, and for mouse chromosome 11 in Supplementary Table S5. The updated human chromosome 17 sequence can be accessed through GenBank accession number NC\_000017. The updated mouse chromosome 11 sequence can be accessed through the accession numbers listed in Supplementary Table S5. Reprints and permissions information is available at [npg.nature.com/reprintsandpermissions](http://npg.nature.com/reprintsandpermissions). The authors declare no competing financial interests. Correspondence and requests for materials should be addressed to M.C.Z. ([mzcody@broad.mit.edu](mailto:mzcody@broad.mit.edu)) or C.N. ([chad@broad.mit.edu](mailto:chad@broad.mit.edu)).