Hominoid fission of chromosome 14/15 and the role of segmental duplications

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Ape chromosomes homologous to human chromosomes 14 and 15 were generated by a fission event of an ancestral submetacentric chromosome, where the two chromosomes were joined head-to-tail. The hominoid ancestral chromosome most closely resembles the macaque chromosome 7. In this work, we provide insights into the evolution of human chromosomes 14 and 15, performing a comparative study between macaque boundary region 14/15 and the orthologous human regions. We construct a 1.6-Mb contig of macaque BAC clones in the region orthologous to the ancestral hominoid fission site and use it to define the structural changes that occurred on human 14q pericentromeric and 15q subtelomeric regions. We characterize the novel euchromatin–heterochromatin transition region (~20 Mb) acquired during the neocentromere establishment on chromosome 14, and find it was mainly derived through pericentromeric duplications from ancestral hominoid chromosomes homologous to human 2q14–qter and 10. Further, we show a relationship between evolutionary hotspots and low-copy repeat loci for chromosome 15, revealing a possible role of segmental duplications not only in mediating but also in “stitching” together rearrangement breakpoints.

[Supplemental material is available for this article.]

Results

Evolutionary history of chromosomes 14 and 15

We reconstructed the evolutionary history of chromosomes 14 and 15 in the primate lineage from published data (Yunis et al. 1980; Yunis and Prakash 1982; Locke et al. 2003; Ventura et al. 2003; Zody et al. 2006; Stanyon et al. 2008; Locke et al. 2011) and data from the marmoset genome sequencing project (Marmoset Genome Sequencing Consortium, unpubl.). The synteny of chromosomes 14/15 is ancestral to placental mammals (Wienberg and Stanyon 1995, 1997; Kent et al. 2002; Pontius et al. 2007; Wade et al. 2009; Meyer et al. 2013), as demonstrated by cat chromosome B3, horse chromosome 1, and rabbit...
chromosome 17 (UCSC Genome Browser alignment net, http://genome.ucsc.edu/). In the primate ancestor, chromosomes 14 and 15 were a submetacentric chromosome with a small p arm (Fig. 1). In the marmoset lineage, one fission and two inversions derived marmoset chromosomes 6p and 10, with 6p maintaining the ancestral association of A–F–E markers (Zody et al. 2006). A pericentric inversion in the Catarrhini ancestor generated a submetacentric chromosome with the same organization as the rhesus macaque chromosome 7 (MMU7). In the Hominoidea ancestor the submetacentric chromosome underwent a fission event and generated the acrocentric chromosomes 14 and 15. The ancestral centromere was inactivated and two neocentromeres emerged (Ventura et al. 2003). Finally, a pericentric inversion occurred in the gorilla and chimpanzee lineages, respectively on chromosomes 14 and 15 (Yunis et al. 1980; Yunis and Prakash 1982; Locke et al. 2003; Ventura et al. 2011).

Comparative analysis of the human and macaque genomic regions orthologous to the hominoid 14/15 fission site

MMU7 has a chromosomal organization resembling the ancestral hominoid 14/15 association and shows perfect marker order conservation with respect to the two human-derivative chromosomes (Ventura et al. 2003). To gain insights into the genomic organization of the hominoid fission region, we constructed a 1.6-Mb contig of 61 macaque BAC clones on MMU7 spanning the 14/15

Figure 1. Evolutionary history of chromosomes 14 and 15. Ten segments (A–J) are used to track the chromosomal rearrangements and are defined as in Stanyon et al. (2008). The following abbreviations are used: A in red, ancestral centromere; N in red, evolutionary neocentromere; (HSA) Homo sapiens; (PTR) Pan troglodytes; (GGO) Gorilla gorilla; (PPY) Pongo pygmaeus; (MMU) Macaca mulatta; (CJA) Callithrix jacchus. In Figures 1, 2, and 5, chromosomes 14 and 15 are colored according to the color code of the UCSC Genome Browser (dark blue for chromosome 14 and light blue for chromosome 15). The NOR (nucleolus organizer region) bearing short arm is drawn with a horizontal black/white striped box.
boundary orthologous to human 14q11 and 15q26.3 (Fig. 2; Supplemental Table S1). We screened the macaque genomic library, CHORI-250, using STS derived from the human 15q sub-terminal (Hsa15a-d) and 14q pericentromeric (Hsa14a-c) regions (hg18/build36) together with macaque-specific STS derived from CH250 BAC end sequences (BES). The corresponding region was

**Figure 2.** Contig of CH250 BAC clones on MMU7. The schematic shows features of the macaque genomic region spanned by the contig (chr7:81552000–83132000, rheMac2). In particular, gap and STS locations, BAC clones, FISH mapping data, chained alignments with hg18 reference, WSSD (whole-genome shotgun sequence detection) depth-of-coverage data (the green color corresponds to single copy; the red color to duplication; Marques-Bonet et al. 2009), macaque RefSeq genes, and human RefSeq genes in the orthologous human regions are shown. STS names are color-coded: STS used in radioactive hybridization (orange), STS used in PCR amplification (blue), and STS used in both assays (green). FISH results on MMU7 and human chromosomes 14/15 are shown for representative BAC clones marked by names and arrowheads in red. Seven different FISH patterns are identified: BAC clones of Groups F1–F3 map to the homologous region on chromosome 15 (light blue); BAC clones of Groups G2–G4 map to the homologous region on chromosome 14 (dark blue); and gray-colored BAC clones (Group G1) showed absent or weak signal on human chromosome 14 and correspond to the fission breakpoint. An asterisk in front of the BAC name indicates BAC clones belonging to the second copy of the macaque duplication. A yellow arrowhead marks BES not anchored to the human reference hg18. Additional FISH signals on other human chromosomes due to the presence of interchromosomal SDs (further indicated by the chained alignments) are shown for the BAC clones of Groups F3, G2, and G3. An empty black box marks the human GOLGA2–ITSN2 core duplicon. Both end segments of the contig (Groups F1 and G4) are in single copy and direct orientation in macaque and human.
single copy in macaque, except for a 134-kb segment duplicated in tandem. We FISH-(fluorescence in situ hybridization) mapped all clones on both macaque and human chromosome metaphases and grouped them accordingly by FISH pattern (F1–F3 clones map to human 15q26.3; G1–G4 clones map to human 14q11); most of the clones were single copy in macaque and duplicated intra- and/or interchromosomally in human (Supplemental Note). We generated 91 BES and mapped them on the macaque (rheMac2) and human (hg18/build36) reference genome assemblies. We found consistent placements on rheMac2, except for those partly covering assembly gaps. In the human reference assembly, several BACs were one-end anchored and some end sequences showed multiple matches due to the presence of SDs in the human genome (Supplemental Table S1).

Extreme groups (F1 and G4) are made up of clones spanning single copy and directly oriented segments in human and macaque. For the homologous region on chromosome 15, we identified an LCR15 copy in macaque and human genomes (Group F2) and human subterminal duplications of the last segment preserving the same centromere–telomere orientation except for 1p (Group F3) (Supplemental Note). We detected clones (Groups F3 and G1) covering the evolutionary fission breakpoint as revealed by the BAC end mapping (one-end anchored on human chromosomes 14 or 15) and absence of signal on human metaphases for three clones. For the homologous region on chromosome 14, we detected three human interchromosomal pericentric duplications (15q11, 18p11, and 21q11; Groups G2 and G3) dated to have duplicated prior to human–orangutan divergence (Supplemental Note). Furthermore, we identified a hominoid-specific inversion and macaque tandem duplication with 99.85% identity (Fig. 2) terminally flanked by human and macaque single copy as well as human/macaque direct-oriented segments in human and macaque. This suggests that the hominoid inversion and 14–15 duplication, as well as human/macaque direct-oriented regions. This suggests that the macaque 9q pericentromeric sequence is incomplete in the macaque reference and differs significantly from the human 10q pericentromeric sequence. This finding was confirmed by the illumina sequencing of two clones (CH250-48F1 and CH250-34C6). On the other hand, Group V and VI clones mapped to a nonpericentromeric region (human 10q22.3 orthologous location) and to a different pericentromeric region (human 12p11 orthologous location), respectively. At human 14q11, clones showed signal (Groups II, III, and IV), faint signal (Groups I and V), or no signal (Group VI).

We found a similar pattern by a reciprocal approach using human BAC clones (Supplemental Fig. S8). In human, 14q pericentromeric sequence (tandem duplicated signal at this locus) showed interchromosomal pericentricomeric duplication mainly...
Figure 3. Relationships between the ITSN2/DNM1–GOLGA2 duplications and chromosome 15 evolutionary hotspots. Gap locations, WSSD-positive regions, depth-of-coverage data, and locations (BLAT search; Kent 2002) of the ITSN2, DNM1, and GOLGA2 duplications are shown for the sequence of chromosome 15 (hg19/build37). (Note, only a few LCR15 copies lack one of the three components.) Asterisks indicate the 11 loci analyzed through FISH. The BAC triplets were chosen as follows: one BAC (in red) spanning LCR15 and the other two at the single-copy regions immediately upstream (in green) and downstream (in blue) of the duplication cluster. SDs (hg19) and FISH mapping data on human, macaque, and marmoset chromosomes are shown for seven loci (named A, AB, D1, D2, DE1, DE2, and F, and indicated by a black asterisk). For BAC clones spanning LCR15 sequences (red-colored) only signals co-mapping with the single-copy clones (green and blue colored) are shown. BAC clones designed in single-copy regions of loci AB and F co-hybridized in human and macaque, whereas in marmoset they hybridized on chromosomes 6 and 10, respectively, for green- and blue-colored clones; no signal is on marmoset chromosomes for the red-colored BAC. Locus DE1 single-copy BAC clones mapped to marmoset chromosomes 10 and 6, respectively; the red-colored BAC clone showed signal on both chromosomes. Locus DE2 single-copy BAC clones hybridized to the macaque 7p and 7q pericentromeric regions, respectively.
Figure 4. Primate phylogeny of GOLGA2 and ITSN2 duplicons. Neighbor-joining phylogenetic trees were constructed using a 2.5-kb region of the GOLGA2 module and a 3.3-kb region of the ITSN2 module. Sequences were retrieved from human (hg19), chimpanzee (panTro3), gorilla (gorGor3), orangutan (ponAbe2), macaque (rheMac2), and marmoset (calJac3) reference genomes. Trees are drawn to scale, with branch lengths measured in the number of substitutions per site and bootstrap values (1000 replicates) shown. Monophyletic clades of duplications mapping to ancestral loci (9q34.11 and 2p23.3 for GOLGA2 and ITSN2, respectively), chromosome 15, marmoset duplications, macaque and marmoset pericentromeric region, inversion breakpoints, and the pericentromeric region of ape chromosome 15 are highlighted in color code. The copies at the proximal inversion breakpoint and a pericentromeric region homogenized, especially in the ITSN2 phylogenetic tree, do not group into two different branches.
Human chromosomes 14 and 15 evolution

with chromosome 22. Of note, all clones hybridized to human 2q21 and the orangutan orthologous locus along with the macaque 9q pericentromeric region. In orangutan they showed interchromosomal pericentromeric duplicated signals (in two cases at 10q); in addition, four clones showed a tandem duplicated signal on macaque chromosome 10.

SD data (Bailey et al. 2001, 2002a) for this region (Table 1) reported that (1) 76% of the duplications (93% of the total length) are shared with ancestral hominoid acrocentric chromosomes (13, 14, 15, 21, and 22 together with 2A, 2B, 9, 11, and 18, which are no longer acrocentric in human; Supplemental Note); and (2) 58% of sites (79% of the total length) map to the q arm pericentromeric region of these chromosomes (13q11, 14q11, 15q11, 21q11, and 22q11 together with 2q21, 9p11, and 18p11, which are orthologous to ancestral q arm pericentromeric regions). The 2q21 locus shows the greatest homology in terms of both the number and the sequence length of shared duplications. Among the other chromosomal positions, the highest values of SD are with 9q12 (chromosome 9 heterochromatin) and chromosome 10.

Table 1. 5Ds of the pericentromeric region of human chromosome 14

| Segmental duplications     | N  | Percent | Size (bp) | Percent |
|-----------------------------|----|---------|-----------|---------|
| Total                       | 245 | 100%    | 5,613,798 | 100%    |
| Total on HAC                | 186 | 76%     | 5,241,374 | 93%     |
| Total on q11 of HAC         | 141 | 58%     | 4,418,845 | 79%     |
| 9p11                        | 7  | 3%      | 86,423    | 2%      |
| 13q11                       | 5  | 2%      | 75,655    | 1%      |
| 14q11                       | 16 | 7%      | 619,901   | 11%     |
| 15q11                       | 21 | 9%      | 440,195   | 8%      |
| 18q11                       | 12 | 5%      | 336,842   | 6%      |
| 2q21 (129.6–132.7 Mb)       | 61 | 25%     | 1,613,843 | 29%     |
| 21q11                       | 8  | 3%      | 293,574   | 5%      |
| 22q11                       | 11 | 4%      | 952,412   | 17%     |
| Total (other)               | 97 | 40%     | 1,087,017 | 19%     |
| chr1                        | 3  | 1%      | 12,334    | 0%      |
| 2p11 (other)                | 3  | 1%      | 14,011    | 0%      |
| chr4_random                 | 3  | 1%      | 40,732    | 1%      |
| chr7                        | 4  | 2%      | 6932      | 0%      |
| chr9q12                     | 22 | 9%      | 555,260   | 10%     |
| chr9 (other)                | 3  | 1%      | 49,962    | 1%      |
| chr10                       | 21 | 9%      | 177,995   | 3%      |
| chr11 (other)               | 3  | 1%      | 4448      | 0%      |
| chr12                       | 7  | 3%      | 27,374    | 0%      |
| 13q (other)                 | 7  | 3%      | 90,912    | 2%      |
| chr16                       | 8  | 3%      | 19,390    | 0%      |
| chr17 and chr17_random      | 10 | 4%      | 55,255    | 1%      |
| chrY                        | 3  | 1%      | 32,412    | 1%      |
| Unclassified                |    |         |           |         |
| chr21_random                | 6  | 2%      | 104,873   | 2%      |
| chr22_random                | 1  | 0%      | 3063      | 0%      |

Number and total length of SDs classified according to the chromosomal location of the duplicate copy. Those mapping to chromosomes orthologous to ancestral hominoid acrocentric chromosomes (HAC, chromosomes 2A, 2B, 9, 11, 13, 14, 15, 18, 20, 21, and 22) and to regions orthologous to their q arm pericentromeric region are distinguished from those mapping to other chromosomal locations. Two different locations, q11 of the ancestral acrocentric chromosome and “other”, were considered for these chromosomes. Regarding chromosome 9, the location 9q12 was also distinguished. Chr21_random and chr22_random are unclassified because the location (21q11 or chr21 [other], 22q11 or chr22 [other]) for these duplications could not be assigned.

Discussion

Reshuffling of the hominoid OR gene repertoire after the 14/15 fission

An ancestral hominoid fission event gave rise to human chromosomes 14 and 15 and converted a single-copy euchromatic region from an interstitial to a terminal location. Since the genomic organization of the macaque region orthologous to the fission site closely resembles the ancestral hominoid one, we constructed a 1.6-Mb macaque BAC clone contig and compared the content and organization of human and macaque. Sequences from the novel 15q subtelomeric and 14q pericentromeric regions were extensively and specifically duplicated to other subtelomeric and pericentromeric regions, respectively, probably as a direct consequence of the evolutionary change in chromosomal location and consistent with the SD enrichments observed for these regions in the human genome (Bailey et al. 2001; Linardopoulou et al. 2005; Bailey and Eichler 2006). Notably, we found that 14q pericentromeric sequences were duplicated to the q arm pericentromeric region of other acrocentrics (15 and 21) or ancestrally acrocentric (18) chromosomes and subsequently experienced duplication exchanges mainly with acrocentric or ancestrally acrocentric chromosomes (Table 1). These observations suggest that, first, the pericentromeric region of acrocentric chromosomes is rather preferentially duplicated with other acrocentric chromosomes and, second, these exchanges occur primarily between the q arms possibly because the presence of the rDNA and other repetitive sequences prevent the “invasion” of the p arm, as previously reported in chimpanzee and gorilla telomeric ends (Ventura et al. 2012).

Clones positive to the STS 265F1417 showed weak FISH signals and single-end sequence placement to human 15q26.3 or 14q11 (Fig. 2; Supplemental Table S1), consistent with the occurrence of the fission event and sequence loss at this site, as previously shown for clone CH250–246C20 (Rudd et al. 2009). The fission and consequent chromosomal rearrangements shuffled the OR gene repertoire of hominoids. In particular, the fission caused the loss of OR gene copies at the breakpoint (Rudd et al. 2009) while the consequent duplication events created additional copies in other subterminal and pericentromeric regions. We observed a correlation between the presence of OR genes and the involvement in chromosomal rearrangements (deletion, duplication, and inversion)—all regions that underwent rearrangement in human and in both human and macaque genomes embedded OR genes (besides the ITSN2/DNM1–GOLGA2 duplication) whereas the flanking single-copy regions contained few OR genes and genes other than OR. OR genes form the largest mammalian gene family and have been subjected to extensive pseudogenization particularly in the primate lineage. Not surprisingly, they have been continuously reshuffled during mammalian evolution by frequent duplication and deletion events (Niimura and Nei 2007). In this regard, their presence at the novel subtelomeric and pericentromeric regions and subsequent duplications unlikely affected the species’ survival, and the fission event and its consequences were likely neutral.

ITSN2/DNM1–GOLGA2 duplication clusters map to evolutionary hotspots

Multiple studies demonstrated the association of SDs and evolutionary rearrangements in primates (Armengol et al. 2003; Locke
et al. 2003; Bailey et al. 2004; Murphy et al. 2005). In this study, we further prove this association, specifically between LCR15 and regions involved in evolutionary rearrangements. We show that LCR15 sequences cluster around (1) the evolutionary new centromere (15q11.2, ~3 Mb away from the centromere); (2) the breakpoints of the pericentric inversion that occurred in the Catarrhini ancestor (15q13 and 15q26.3); and (3) the ancestral centromere (15q24–q25, <1 Mb from the ancestral centromere locus). Interestingly, golgin repeats were also found at the breakpoint of pathogenic chromosome 15 deletions (F Antonacci, MY Dennis, J Huddleston, PH Sudmant, K Meltz, J Graves, L Vives, M Malig, CT Amemiya, A Stuart, et al., unpublished), emphasizing and confirming the dual effect of SD genomic instability in both evolution and disease.

LCR15 copies are present on the marmoset and marmoset orthologous regions of 15q11.2 and, therefore, were likely present in the Haplorhini ancestor, but expanded in human, possibly as a result of the neocentromere emergence. LCR15 sequences are present on human and marmoset yet absent from marmoset regions orthologous to the inversion breakpoints that occurred in the Catarrhini ancestral submetacentric chromosome (loci AB and F), revealing these copies may have emerged as a result of the inversion or may have driven the event.

There is additional evidence of LCR15 evolutionary instability. The same site at 15q11 was shown to be involved in the pericentric inversion of chimpanzee 15p and in the insertion of a novel SD within the human lineage (Locke et al. 2003). Here, we found the occurrence of two independent chromosomal rearrangements in proximity to one site (human 15q26.3) after its acquisition of an LCR15 copy—the chromosomal fission in the hominoid lineage and the duplicative insertion from chromosome 13 in the macaque lineage. This suggests that LCR15 may not only serve as preferential sites of chromosomal breakage and rearrangement, like the chimpanzee inversion, but they may be subject to further expansion and duplication associated with recurrent events, such as the duplicate copies mapping to the inversion breakpoints in the Catarrhini ancestor.

In FISH experiments, LCR15 signals observed on marmoset metaphases mapped mainly to the pericentric regions of chromosomes 6p and 10p—particularly 10p—orthologous to human 15q24–q25 (loci D1, D2, DE1, and DE2), suggesting that the ancestral pericentric region was probably the seed of LCR15 and the first chimeric ITSIN2/DNM1–GOLGA2 unit likely formed at this locus in the Haplorhini ancestor. Moreover, since no copy is present in the marmoset region orthologous to human chromosome 10, this copy emerged in the Catarrhini ancestor; therefore, it was unlikely the site where the first combined unit formed as previously suggested (Zody et al. 2006).

The chromosome 14 pericentric region consists of SDs from the pericentric region of ancestral hominoid chromosomes 2B and 10

After the fission event, human chromosome 14 acquired ~19 Mb of sequence over the last 25 million years. It is largely composed of three blocks: (1) a 1.3-Mb transition region between the q arm euchromatin and the centromere; (2) a 3-Mb centromere composed of alphoid DNA; and (3) a 15-Mb heterochromatic short arm containing arrays of ribosomal RNA genes, satellite sequences, and other repetitive elements (Kehrer-Sawatzki et al. 1998). We analyzed the first of these three blocks and showed that it was mainly formed through pericentric duplicative transpositions from ancestral hominoid chromosomes 2B and 10 (homologous to human 2q14–qter and 10, respectively). We note that the centromere of chromosome 2B inactivated in both the macaque (Ventura et al. 2007) and human lineages—the latter after a fusion event occurred 3–5 million years ago and mapping to 2q21 (Ijdo et al. 1991). The pericentric sequences on chromosome 10 were lost in human and African ape, and evidence of their ancestral location on this chromosome still exists in orangutan and macaque. Notably, chromosomes 2B and 10 have been partners in another process during great ape evolution—the formation of subterminal heterochromatin in the chimpanzee and gorilla lineages (Ventura et al. 2012) but the genomic regions involved in the process were different.

STS derived from the new human chromosome 14 pericentric sequence and probed against the macaque genomic library retrieved clones mapping to either pericentric or euchromatic regions. Among the latter, the euchromatic SD-rich region at human 10q22.3 is associated with chromosomal rearrangement and human disease (Melberg et al. 1999; Quintero-Del-Rio et al. 2002; Fallin et al. 2003; Faraone et al. 2006; Hofmann et al. 2008; Kuhl et al. 2008; Venken et al. 2008; O’Meara et al. 2012); in the gorilla genome, one breakpoint of the inversion of chromosome 10q is 1 Mb proximal to this locus (Carbone et al. 2002; Ventura et al. 2011).

In summary, we propose a model that describes the evolution of ape chromosomes 14 and 15 (fig. 5). After the fission, the resulting two new chromosomal ends achieved different functional roles with respect to the chromosome. The chromosome 14 portion acquired the centromeric functionality necessary for daughter cell segregation, resulting in a telocentric chromosome, while the chromosome 15 end became a subtelomere participating in interchromosomal DNA exchanges with other subtelomeric regions. Both novel chromosome ends earned telomeric repetitive sequences (TTAGGG) to be protected from degradation and to prevent chromosomal end-to-end fusions. The original centromere, located on chromosome 15, became inactive and a neo-centromere was seeded at the chromosome tip not derived by the fission, also making chromosome 15 a telocentric chromosome. The neo-centromeres rapidly acquired the complex organization characteristic of primate centromeres—a large core of alphoid sequences surrounded by SDs (Eichler et al. 1999). SDs mainly from the long arm pericentric region of the acrocentric chromosome 2B and from the pericentric region of chromosome 10 contributed to the formation of the 14q pericentric region. SDs involving this genomic region continue to accumulate in the human lineage, mainly within the long arm pericentric region of chromosome 22 (Bailey et al. 2002b). It is known that the juxtaposition of constitutive heterochromatin and transcriptionally active genes induces position effects, effectively silencing transcription (Dillon and Festenstein 2002). The SDs served the potentially important role of buffering chromosome 14 euchromatin and the novel centromeric heterochromatin, possibly protecting and maintaining the transcription regulatory landscape of the 14q11 genes suddenly found in a pericentric region. Chromosome 15 experienced further LCR15 duplications that generated new copies, particularly within the novel pericentric region. In the human karyotype, the neo-centromere of chromosome 14 acquired alphoid DNA of the suprachromosomal families 2, 4, and 5; the one of chromosome 15 acquired alphoid DNA of the suprachromosomal families 2 and 4 (Alexandrov et al. 2001). Finally, duplications from the p arm of acrocentric chromosomes created the heterochromatic p arm of ribosomal RNA.
genes, turning chromosomes 14 and 15 from telocentric to acrocentric chromosomes. We note that in human, chimpanzee, and orangutan, only acrocentric chromosomes bear the rDNA on the p arm, implying that the acquisition of the rDNA short arm was an obligated fate for chromosomes 14 and 15. Yet, in the gorilla genome only two acrocentric chromosomes, 21 and 22, carry the rDNA while the short arms of the remaining acrocentric chromosomes are composed mainly of satellite III sequences (Ventura et al. 2012).

This work highlights the dynamic and intimate relationship between duplication and rearrangement in the evolution of human chromosomes 14 and 15. It raises the possibility that not only duplication triggers chromosomal rearrangement through NAHR but also that chromosomal breakage gives rise to duplication events (Kehrer-Sawatzki and Cooper 2008; Girirajan et al. 2009). We propose a possible additional role of SDs in genome evolution where the occurrence of duplication events in response to chromosomal breakage might represent a possible “feedback mechanism” that produces additional DNA material to recover the damage caused by the breakage.

Methods

Hybridization of high-density filters

Radioactive genomic hybridization of the sixfold coverage CHORI-250 (rhesus macaque) segment 1 BAC library was carried out according to the protocol available at CHORI BACPAC resources (http://bacpac.chori.org/highdensity.htm). Probes were obtained by means of PCR amplification of macaque genomic DNA (Supplemental Table S4). The number of positive clones provided estimation of the copy number in the macaque genome. BES were repeatmasked (Smit and Hubley 2010) and mapped on macaque (rheMac2) and human (hg18) reference genomes using the BLAT tool at the UCSC Genome Browser (http://genome.ucsc.edu/cgi-bin/hgBlat).

Fluorescence in situ hybridization (FISH)

Metaphase spreads and interphase nuclei were prepared from lymphoblastoid or fibroblast cell lines of Homo sapiens (HSA), Pan troglodytes (PTR), Gorilla gorilla (GGO), Pongo pygmaeus (PPY), Macaca mulatta (MMU), Cercopithecus aethiops (CAE), Trachypithecus cristatus (TCR), and Callithrix jacchus (CJA). FISH experiments were performed using BAC clones (300 ng) directly labeled by nick-translation with Cy3-dUTP, Cy5-dUTP, and fluorescein-dUTP as previously described (Lichter et al. 1990) with minor modifications. Hybridization was performed at 37°C in 2× SSC, 50% (v/v) formamide, 3 μg Cot-1 DNA, and 3 μg sonicated salmon sperm DNA, in a volume of 10 μL. Post-hybridization washing was at high stringency (60°C in 0.1× SSC, three times) or at low stringency for cross-species hybridization (37°C in 2× SSC, 50% formamide, three times, and 42°C in 2× SSC, three times). Nuclei and chromosome metaphases were DAPI-stained. Digital images were obtained using a Leica epifluorescence microscope equipped with a cooled CCD camera. Fluorescence signals detected with Cy3, Fluorescein, and Cy5 filters and chromosomes and nuclei images detected with DAPI filter were recorded separately as grayscale images. Pseudocoloring and merging of images were performed using Adobe Photoshop software.

Phylogenetic analysis

Duplicate copies of a 3-kb region from ITS2 and a 2.5-kb region from GOLGA2 were retrieved from the hg19, panTro3, gorGor3, ponAbe2, rheMac2, and calJac3 reference genomes through BLAT...
search (http://genome.ucsc.edu/cgi-bin/hgBlat) (Kent 2002). The hg19 reference was used instead of hg18 because in the most recent release of the human genome (hg19), all LCR15 duplcions are placed on the chromosome 15 sequence, and there are no chr15 random locations. Multiple sequence alignments were performed using Clustal W (Thompson et al. 1994) (match 5, mismatch −2, match with N 0, gap opening 25, gap extension 0.05 for the IINS2 region; match 4, mismatch −2, match with N 0, gap opening 15, gap extension 0.05 for the GOLGA2 region). Neighbour-joining phylogenetic trees based on the Kimura 2-parameter model (Kimura 1980) (pairwise deletion option) were constructed in MEGA5 (Tamura et al. 2011). A bootstrap test with 1000 replicates was conducted to evaluate the statistical significance of each node (Felsenstein 1985).

Sanger sequencing

Ten STS regions were PCR-amplified from positive BAC clones and PCR products were sequenced using PCR primers (Supplemental Table S4). DNA from BAC clones were extracted using the BAC Prep Protocol (Schein et al. 2004), and BAC ends were sequenced (if not available in the NCBI Trace Archive) using SP6 and T7 primers. Sequencing was performed on an ABI PRISM 3100 Genetic Analyzer using the BigDye Terminator v3.1 Chemistry (Applied Biosystems) according to manufacturer instructions.

Illumina sequencing and analysis of BAC clones

BAC clone DNA from the CH250 macaque library was isolated by a modified alkaline lysis miniprep procedure and paired-end sequencing on an Illumina HiSeq 2000 (Supplemental Note). Short Illumina reads (50 bp) were mapped to the hard-masked references hg18 and rhesMac2 with mrFAST (Ihach et al. 2010) using an edit distance of three and single-end mapping mode. We determined the best location for each clone using the coverage of reads across each genome. For each clone we calculated the mean and standard deviation of its coverage and selected regions for which the coverage was greater than the mean plus two standard deviations and the total size of the region was >1 kbp. We manually investigated each of the resulting regions in the UCSC Genome Browser to determine the best location and whether any regions should be merged. We determined the repeat content of short reads by aligning reads to all known RepeatMasker and Tandem Repeat Finder (TRF) sequences in hg18 and rhesMac2. Repetitive coordinates were obtained from UCSC Genome Browser tracks (rmsk.txt.gz for RepeatMasker and chromTrf.tar.gz for TRF) and corresponding sequences were extracted from each reference to create a “repeat” reference. We mapped short reads to these repeat references using BWA (Burrows-Wheeler Alignment) in single-end mapping mode and counted the total number of reads that mapped to each reference with at least a quality of 1. To determine the amount of vector sequence or bacteria contamination present in the BAC short reads, we aligned these reads to a reference consisting of the vector pTARBAC2.1 and the Escherichia coli strain DH10B using mrFAST with an edit distance of three and single-end mapping mode. We counted the number of reads that mapped to the vector and the E. coli genome with a quality of at least 1 to determine the proportion of vector or E. coli reads in the complete set of reads.

Data access

BAC end sequences are available in the Genome Survey Sequence division of GenBank (http://www.ncbi.nlm.nih.gov/nuccore) under accession numbers KG088791–KG088955. CHORI-250 BAC clone sequence data are available in the Sequence Read Archive (SRA; http://www.ncbi.nlm.nih.gov/sra) under accession number SRP028268.

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Author contributions: G.G. and M.V. designed the study. G.G. and M.P. performed BAC library hybridizations and FISH experiments. G.G. performed PCR product sequencing and phylogenetic analysis. F.A. constructed shotgun sequencing libraries and J.H. analyzed sequencing reads. M.M. and L.V. performed BAC end sequencing. G.G. and M.V. analyzed the data. G.G., E.E.E., and M.V. wrote the manuscript.

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