Under the genomic radar: The Stealth model of *Alu* amplification

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Alu elements are the most successful SINEs (Short INterspersed Elements) in primate genomes and have reached more than 1,000,000 copies in the human genome. The amplification of most *Alu* elements is thought to occur through a limited number of hyperactive “master” genes that produce a high number of copies during long evolutionary periods of time. However, the existence of long-lived, low-activity *Alu* lineages in the human genome suggests a more complex propagation mechanism. Using both computational and wet-bench approaches, we reconstructed the evolutionary history of the *AluYb* lineage, one of the most active *Alu* lineages in the human genome. We show that the major *AluYb* lineage expansion in humans is a species-specific event, as nonhuman primates possess only a handful of *AluYb* elements. However, the oldest existing *AluYb* element resided in an orthologous position in all hominoid primate genomes examined, demonstrating that the *AluYb* lineage originated 18–25 million years ago. Thus, the history of the *AluYb* lineage is characterized by ~20 million years of retrotranspositional quiescence preceding a major expansion in the human genome within the past few million years. We suggest that the evolutionary success of the *Alu* family may be driven at least in part by “stealth-driver” elements that maintain low retrotranspositional activity over extended periods of time and occasionally produce short-lived hyperactive copies responsible for the formation and remarkable expansion of *Alu* elements within the genome.

[The sequence data from this study have been submitted to GenBank under accession nos. AY791249–AY791290.]

As the most successful SINEs (Short INterspersed Elements) in primate genomes, *Alu* elements have enjoyed remarkable proliferation during the primate radiation and have expanded to more than 1,000,000 copies in the human genome (Lander et al. 2001; Batzer and Deininger 2002). Despite considerable progress in the understanding of their biology and distribution throughout primate taxa (Salem et al. 2003b; Singer et al. 2003; Carter et al. 2004; Hedges et al. 2004; Otieno et al. 2004; Roos et al. 2004; Ray et al. 2005), a great deal of uncertainty still remains concerning their strategy for survival. It has been generally accepted that the amplification of most *Alu* elements occurs through a small number of long-lived high activity “master” genes (Deininger et al. 1992; Deininger and Batzer 1993), although there is considerable debate as to the details of this amplification strategy (Matera et al. 1990; Schmid 1993; Batzer et al. 1995; Cordaux et al. 2004; Price et al. 2004). In this model, the mutations accumulated in the master genes are inherited by the copies they produced, and consequently, a series of hierarchical *Alu* subfamilies that share the novel diagnostic mutation(s) are generated (Slagel et al. 1987; Willard et al. 1987; Britten et al. 1988; Deininger and Slagel 1988; Jurka and Smith 1988; Deininger et al. 1992; Batzer and Deininger 2002). On the other hand, the evolutionary history of the *AluYa5* lineage, one of the most active human *Alu* lineages, suggests that the founder gene of this *Alu* lineage existed long before the major expansion of the lineage within the human genome

(Leefflang et al. 1993). Contrary to the prediction of the master gene model, it has maintained low retrotranspositional activity, and this founder gene itself may not be directly responsible for the propagation of the recent human *AluYa5* elements (Shaikh and Deininger 1996). Thus, these studies suggest that the expansion of *Alu* elements may follow a more complex propagation mechanism. Unfortunately, aside from the *AluYa5* lineage, little data exist concerning the evolutionary origin of other *Alu* subfamilies in humans, making it difficult to assess the evolutionary significance of the results reported in the original *AluYa5* lineage studies.

To gain additional insight into *Alu* subfamily propagation, we reconstructed the evolutionary history of the *AluYb* lineage, one of the largest and most active *Alu* lineages in the human genome (Jurka 1993; Carter et al. 2004) that composes ~40% of the human-specific *Alu* elements (Hedges et al. 2004). This lineage, originally termed Sb2, is characterized by a 7-nt duplication involving positions 246 through 252 of the *AluY* consensus sequence (Jurka 1993; Batzer et al. 1996). The *AluYb* lineage in the human genome is subdivided into three major subfamilies as follows: *AluYb7*, *AluYb8*, and *AluYb9* (Batzer et al. 1996; Roy-Engel et al. 2001; Jurka et al. 2002; Carter et al. 2004) based on diagnostic mutations following the standardized nomenclature for *Alu* repeats (Batzer et al. 1996). The majority of the human *AluYb* elements integrated into the genome during the last 3 to 4 million years (Myr) and reached a total copy number of about 2000 elements (Carter et al. 2004). The human diseases caused by de novo *AluYb8* insertions suggest that this subfamily is currently actively retrotransposing, and a comprehensive analysis of the *Yb* lineage indicated that about 20% of *AluYb* elements are poly-

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Article and publication are at <http://www.genome.org/cgi/doi/10.1101/gr.3492605>.

morphic in human genome (Muratani et al. 1991; Oldridge et al. 1999; Carter et al. 2004). Previous studies suggested that the evolutionary history of the *AluYb* lineage is much older than its period of major expansion in the human genome, and *AluYb* elements have also been identified in other nonhuman primates (Zietkiewicz et al. 1994; Gibbons et al. 2004). Nevertheless, the extent to which these elements are distributed among nonhuman primate species remains undetermined.

Using both a computational approach and polymerase chain reaction (PCR) display methodology, we have determined the distribution of *AluYb* elements in different hominoid genomes. We find that the long-term evolutionary history of the *AluYb* lineage exhibits a pattern that is remarkably similar to that of the *AluYa* lineage. Thus, the evolution of the *AluYa* and *AluYb* lineages illustrate a common strategy for *Alu* element proliferation. We propose a model for the expansion and survival of *Alu* elements in the primate order.

Results

AluYb elements in the common chimpanzee genome

To determine the evolutionary history of the *AluYb* lineage, we first computationally retrieved all of the *AluYb* elements from the first draft of the common chimpanzee (*Pan troglodytes*) genomic sequence (panTro1 Nov. 2003 assembly). A total of 12 *AluYb* elements were identified and subjected to PCR amplification on a common chimpanzee population panel and a separate primate panel composed of human and eight additional nonhuman primates (see Methods). The number of *AluYb* elements recovered from the chimpanzee draft sequence is in good agreement with a previous study (Gibbons et al. 2004). Detailed information on each locus, including primer sequences, annealing temperature, PCR product sizes, and chromosomal locations are shown in Table 1. Among the 12 elements identified, 10 belong to the *AluYb8* subfamily, while the other two are non-*AluYb7/8/9* elements (Table 2). Of the 10 *AluYb8* elements, four loci were spe-

cific to the common chimpanzee lineage, including three elements that were polymorphic within the chimpanzee population panel (Figs. 1, 2B). Five elements were shared between the pygmy chimpanzee and common chimpanzee lineage (Fig. 2C) and one was present within human, chimpanzee, and gorilla genomes. These results suggest that the initial expansion of the *AluYb8* subfamily predates the divergence of gorillas and humans/chimpanzees, which is thought to have occurred ~7 million years ago (Mya) (Goodman et al. 1998). In addition, two non-*AluYb7/8/9* elements displayed PCR amplicon sizes consistent with the presence of an *Alu* element in all of the hominoid primates we tested.

To confirm that the PCR products were derived from authentic *AluYb* elements, all of the filled amplicons were cloned and sequenced. This sequencing effort revealed additional insight into the evolutionary history of the following four loci: Pan1, Pan2, Pan3, and Pan4. At the Pan1 locus, an *AluYb8* element was inserted into the chimpanzee lineage (pygmy and common) after the divergence of humans and chimps. However, the pygmy chimpanzee and common chimpanzee showed both filled and empty size amplicons in their PCR amplifications, suggesting the presence of an *Alu* insertion polymorphism in both species (Fig. 3A). Sequence analysis of the locus showed that the *AluYb8* element inserted immediately upstream of an *AluSx* element in the same orientation, and that the integration site is partially shared (three copies of direct repeats). More recently, a nonhomologous recombination or intrachromosomal recombination event in the common chimpanzee genome generated a hybrid *Alu* element that is composed of the first half of the older *AluSx* element and the second half of the newly inserted *AluYb8* element. In contrast, the smaller allele amplified in other primates appeared to be the preintegration site of the *AluYb8* element (Fig. 3B).

DNA sequence analysis of the Pan2 and Pan3 loci resulted in the recovery of two gene conversion events. The Pan2 locus appears to be the oldest *Alu* element that we recovered. PCR analysis of the locus showed the presence of an *Alu* element or filled

Table 1. *AluYb* loci identified in this study

Name ^a	Forward primer sequence (5'–3')	Reverse primer sequence (5'–3')	Chr. loc. ^b in human	Direct repeat	A. T. ^c	Product size	
						Filled	Empty
Pan1*	ACCAAATGCAGGTCTCTTGT	CTTTTCTTTATGCGCATTTCT	5:171753413	AAAACATCCTC	55	892	561
Pan2*	CCTGGCCTATTGATGTTTTCT	GCCTCAGAAGGAGTTTTGTTGT	12:42243314	AAAACCTCCCTCTGAG	58.6	481	129
Pan3*	CCGATATCATGCAATTTCCATA	TGGCAAGAAAGACATGATTGAA	3:155790154	AAAAATAAATACCA	60	728	412
Pan4*	AGGGCAATTACTATGTTTCAGGA	TTTTTCACGTTCTTACAATAGAACA	18:66254955	ACAATAGAACATTCCCT	55	403	52
Pan5*	GCTTCATTCTGCTGCTTATT	TCTGCAAAATTAACCTCAAACC	10:97871078	AAAAACAGCAAGT	55	504	181
Pan6*	GGCATTTTAAGCTCTTTGATAGC	CATGCTAGACATGAGAACAAACA	2:183429337	GAAATAGTTCTCTGCT	60	749	257
Pan7*	GCAGCTGCTTTCTGTCTCTGTA	TCAGCAACAATAAGGAACGAAG	3:49401325	AAAAGAAACCAAGTCAC	55	502	175
Pan8*	CCCAGATTGATTCTTCCCTTTA	ATGCCAGTTCCATTATTTCCAC	8:101149947	AAGAGAGAAGAAA	55	834	506
Pan9*	CCAGGACCCAGAGCTTAGTAGT	TCCAAAGGAATATGATGTCACAG	14:52808462	AAAGAAATTGATT	55	434	111
Pan10*	TGTGGCAGATTTTATTGTAGACTT	TCCATGCTCTTGGAGTAAATGA	4:72958275	AAAAAATTCATCTG	55	500	174
Pan11*	AGGAGCTCTTCTAGAAAATCCA	AAAGATCCACGACTAGGCACAT	2:214354697	AAAGAAAAGGAAAAAAC	60	521	193
Pan12*	CGTTAGCTCTGGATTTTCATGT	TATTCTCCACCATGACCAAGTG	X:93270745	AAAAATTAGCCGGGCA	60	871	548
Gorilla7 [#]	TGAAGAATTGGGAGGAAAAAGAA	ACTGCAAGATTGCTGTTCACAT	13:32666822	AAAAATGACTAACAG	55	583	261
Gorilla8 [#]	GCACATCCGTACCTGACTTA	TCAGAGCATCTCTTCTGTCCA	2:229645741	AAAAAAGGAGGCAA	60	540	219
Gorilla16 [#]	AATCTTTGGGGTAGGTGGAAT	CACAACGTACACCTAAATGG	X:24004038	AAGAGTTGCAGTTGCTCA	55	531	202
Gorilla19 [#]	CAAGTTGTGTTATGTGAGGTTTGA	GGAGCCCTAATGTATAGCATGG	7:10262937	AAGAGAGTAGATCTT	59.5	597	279
Gorilla21 [#]	AACAAGAGATGCTAGAAAGCCAAT	CGGAGTTGGACACATTTCTTTT	1:51666132	AAAGAAAGGAGGA	55	749	434
Pygmy4 [#]	GCAGTTTTTCCCATTTGCTCTA	TTGAGTCTTTTCTGGGCTTTC	2:130058820	AAGACTTACTATA	55	600	281

^a(*) Retrieved from chimpanzee genome draft sequence; ([#]) identified using PCR display methodology.

^bChromosomal location.

^cAnnealing temperature.

Table 2. Primate phylogenetic PCR panel results for chimpanzee *AluYb* elements

Name	Chimpanzee panel	Human (HeLa)	Pygmy chimpanzee	Gorilla	Orangutan	Gibbon	Siamang	Green monkey	Owl monkey
Pan1	Fixed present (<i>AluYb8</i>)	—	Polymorphic (<i>AluYb8</i>)	—	—	—	—	x	x
Pan2	Fixed present (<i>AluYb6</i>)	+ (<i>AluYb6</i>)	+ (<i>AluYb6</i>)	+ (<i>AluYb6</i>)	+ (<i>AluYb6</i>)	+ (<i>AluYb6</i>)	+ (<i>AluYb6</i>)	+ (<i>AluSg</i>)	x
Pan3	Fixed present (<i>AluYb</i>)	+ (<i>AluYb</i>)	+ (<i>AluYb</i>)	+ (<i>AluYb</i>)	+ (<i>AluY</i>)	+ (<i>AluY</i>)	+ (<i>AluY</i>)	—	x
Pan4	Fixed present (<i>AluYb8</i>)	+ (<i>AluYb8</i>)	+ (<i>AluYb8</i>)	+ (<i>AluYb8</i>)	—	—	—	—	—
Pan5	Polymorphic (<i>AluYb8</i>)	—	—	—	—	—	—	—	x
Pan6	Fixed present (<i>AluYb8</i>)	—	+ (<i>AluYb8</i>)	—	—	—	—	—	—
Pan7	Fixed present (<i>AluYb8</i>)	—	+ (<i>AluYb8</i>)	—	—	—	—	—	—
Pan8	Fixed present (<i>AluYb8</i>)	—	—	—	—	—	—	—	—
Pan9	Fixed present (<i>AluYb8</i>)	—	+ (<i>AluYb8</i>)	—	—	—	—	—	—
Pan10	Fixed present (<i>AluYb8</i>)	—	+ (<i>AluYb8</i>)	—	—	—	—	—	—
Pan11	Polymorphic (<i>AluYb8</i>)	—	—	—	—	—	—	—	x
Pan12	Polymorphic (<i>AluYb8</i>)	—	—	—	—	—	—	—	—

(+/-) Presence/Absence of *Alu* element; (x) no amplification.

allele in all of the hominoid primates we examined. The green monkey also showed a filled size amplicon, but the amplicon was slightly smaller than the predicted size (Fig. 4A). DNA sequence analysis showed that all hominoid primates possessed an *AluYb* element at the Pan2 locus with the *AluYb* lineage diagnostic duplication and five of the eight additional mutations that charac-

terize the *AluYb8* consensus sequence. However, an *AluSg* element was found in the orthologous locus in green monkey. This result suggests that the *AluYb6* element (missing two diagnostic mutations compared with the *AluYb8* consensus) at the Pan2 locus was introduced into the genome via a gene conversion event after the divergence of Old World monkeys and hominoids, but before the radiation of the hominoid primates. Thus, the *AluYb6* element at the Pan2 locus is 18–25 Myr old (Goodman et al. 1998). Given the sequence identity between the *AluYb* and the *AluSg* consensus sequences, we estimate that the starting point of the gene conversion event was located within the first 75 bp of the *Alu* element, and that the 3' terminus was located between positions #267 and #310.

The Pan3 locus also contains a gene conversion event (Fig. 4B). Siamang, gibbon, and orangutan possess an *Alu* element that does not contain the *AluYb* lineage diagnostic duplication and has the highest sequence identity to the *AluY* consensus. However, the *Alu* elements in the gorilla, chimpanzee, and human genomes belong to the *AluYb* lineage, as indicated by the presence of the *AluYb* diagnostic duplication in these elements. This indicates that a gene conversion event occurred after the divergence of gorillas and orangutans, but prior to the divergence of gorillas, humans, and chimpanzees. During this process, a small proportion of the ancestral *AluY* element has been converted to an *AluYb*-like sequence, including the *AluYb* diagnostic duplication and another adjacent tightly linked diagnostic mutation. The gene-converted region could be as small as 8 bp (251–259) or as long as 30 bp (237–267) in this case. Thus, the overall gene conversion rate (2/12) we observed here seems to be much higher than in previous studies (Maeda et al. 1988; Kass et al. 1995; Roy-Engel et al. 2002; Salem et al. 2003a). However, this is not surprising, since the difference may be due to the small sample size in the current study or reflect the longer evolutionary time period that each element may have been subjected to gene conversion.

Figure 1. Polymorphic *AluYb8* elements in the common chimpanzee genome. Gel chromatographs of two loci are shown. The locus name is shown at right. The product sizes for filled and empty alleles (pre-integration size) are indicated at left. The DNA panel is composed of 12 unrelated common chimpanzee individuals and other primate species. The template used in each reaction is listed at top.

Figure 2. Lineage-specific *AluYb8* insertions. Four examples of gel chromatographs are shown. The locus designation is shown at right, while the product sizes for filled and empty alleles (pre-integration size) are indicated at left. The DNA template used in each reaction is listed at top. (A) Pygmy chimpanzee-specific *AluYb8* insertion. (B) Common chimpanzee-specific *AluYb8* insertion. (C) Chimpanzee lineage-specific *AluYb8* insertion. (D) Gorilla-specific *AluYb8* insertion.

The Pan4 locus contains the oldest *AluYb8* element we identified. Interestingly, the Pan4 *AluYb8* element in the human genome only has one CpG mutation (G→A) at position #5, compared with the *AluYb8* consensus sequence, while the chimpanzee and gorilla *AluYb8* elements at the orthologous Pan4 locus each have accumulated five species-specific mutations compared with the *AluYb8* consensus sequence. Since no mutations are shared by the human, chimpanzee, and gorilla *AluYb8* elements at the Pan4 locus, we believe that the ancestral *AluYb8* element at the Pan4 locus was a canonical *AluYb8* element. Using the BLAT program (<http://genome.ucsc.edu/cgi-bin/hgBlat?command=start>), we identified three additional *AluYb8* elements in the human genome with 100% sequence identity to the Pan4 element. This result suggests that the Pan4 element may still be retrotranspositionally active in the human genome, although we cannot

completely rule out the possibility that these four elements independently mutated at position #5. Multiple alignments of the Pan1, Pan2, Pan3, and Pan4 loci are available on our Web site under publications (<http://batzerlab.lsu.edu>).

AluYb insertions in other primate genomes

To further investigate the propagation of the *AluYb* lineage, we next searched for the presence of *AluYb* elements in additional nonhuman primate genomes. Since no complete draft genomic sequences are available other than human and common chimpanzee, we used a modified PCR display method to identify *AluYb* elements from other primate genomes (see Methods). The display procedure was performed for four hominoid primates: pygmy chimpanzee, gorilla, orangutan, and gibbon; three Old World monkeys: green monkey, rhesus monkey, and pig-tailed macaque. In addition, this approach was also applied to the common chimpanzee to identify additional *AluYb* elements absent from the common chimpanzee genomic sequence. Two restriction enzymes were used for every template, and a minimum of 72 colonies were sequenced for each species. For the common chimpanzee, a total of seven *AluYb* elements were retrieved, all of which had previously been identified from the draft sequence. This suggested that our method involving two restriction enzymes yielded a ~60% coverage (7/12) of the *AluYb* elements in the genome. One pygmy chimpanzee-specific and five gorilla-specific *AluYb8* insertions were identified and confirmed by a PCR analysis of nonhuman primates (Fig. 2A,D) and DNA sequencing. No *AluYb* element was recovered from the orangutan genome using the PCR display approach. The only locus identified within the gibbon genome was Pan2, which had been previously identified in the chimpanzee genome using the computational approach. Despite at least two trials for each species, no *AluYb* elements were identified in the three Old World monkeys (green monkey, rhesus monkey, and pig-tailed macaque) examined. All of the new *AluYb* loci identified by PCR display are listed in Table 2.

Age estimates for the *AluYb8* insertions in chimpanzee and gorilla

To estimate the average age of *AluYb8* elements in the chimpanzee genome, CpG and non-CpG mutation densities were calculated for 10 chimpanzee-specific *AluYb8* elements as reported previously (Xing et al. 2004) using the chimpanzee *AluYb8* consensus. The 10 elements contained a total of nine non-CpG mutations of 2420 nucleotides and 10 CpG mutations of 460 CpG nucleotides. The mutation densities were $0.37\% \pm 0.30\%$ (average $\pm$ standard deviation) and $2.17\% \pm 2.29\%$ for the non-CpG nucleotides and CpG nucleotides, respectively. Using a neutral mutation rate of 0.0015/site/Myr for non-CpG sites and a mutation rate of 0.0090/site/Myr for CpG sites (Xing et al. 2004), the average non-CpG and CpG mutation densities yield age estimates of 2.48 ± 2.03 and 2.42 ± 2.55 Myr, respectively. For the gorilla-specific *AluYb8* elements, a total of five elements were analyzed. The mutation densities were $0.17\% \pm 0.23\%$ and $3.48\% \pm 3.30\%$ for the non-CpG and CpG sites, yielding age estimates of 1.10 ± 1.51 and 3.86 ± 3.66 Myr, respectively.

Phylogenetic analysis of *AluYb* elements in the human genome

The presence of *AluYb8* elements in different primate genomes suggests that the origin of the *AluYb8* subfamily may be much older than its major expansion in humans. In addition, the pres-

bers of the *AluYb8* subfamily, while the other two are non-Yb7/8/9 elements. The presence of an *AluYb* element at the Pan2 locus within siamang and gibbon genomes suggested the *AluYb* lineage originated before the divergence of all hominoid primates. However, no *AluYb* elements have been identified in multiple old-world monkey genomes; thus, the origin of the *AluYb* lineage was after the divergence of Old World monkeys and apes. These results place the origin of the *AluYb* lineage at the early stage of the

The origin of the *AluYb* lineage

The *AluYb* lineage is one of the most active *Alu* lineages in the human genome, with an estimated copy number of ~2000 (Carter et al. 2004). To obtain further insight into the origin of the *AluYb* lineage in the primate order, we analyzed the draft sequence of the common chimpanzee genome and identified 12 *AluYb* insertions, 10 of which are mem-

Figure 4. Gene conversion of *AluYb* elements. Gel chromatographs of PCR products derived from a phylogenetic analysis of the Pan2 locus (A) and Pan3 locus (B) are shown at left. The DNA template used in each lane is shown at top. The product sizes for filled and empty alleles (pre-integration size) are indicated at left. The schematic diagrams depict the potential evolutionary scenarios on the right. Flanking sequences are shown as green boxes; target site direct repeats are shown in red boxes. *Alu* elements are shown as arrows, and the direction of the arrow indicates the orientation (5'→3'), with the head of the arrow denoting the end of the *Alu* elements.

hominoid evolution, about 18–25 Mya (Goodman et al. 1998) It is worth noting that although the *AluYb6* insertion at the Pan2 locus is the oldest *AluYb* element we identified, it was most likely generated via a gene conversion event. Therefore, it may not be the founder gene of the *AluYb* lineage, but rather an early offspring of the *AluYb* founder gene, which was subsequently lost in extant primates.

The Pan4 *AluYb* element appears to have been fixed in the human, chimpanzee, and gorilla genomes, and contains all eight diagnostic mutations that characterize the *AluYb8* consensus sequence. This suggests that within the 10 Myr or so after the *AluYb* lineage initially arose, it was not very active in terms of retrotransposition, if at all. However, this lineage retained its retrotranspositional potential during this extended period of time. Furthermore, our results suggest that in gorilla and chimpanzee genomes, the copy number of *AluYb* elements is two orders of magnitude lower than that in human. Therefore, over several million years following the insertion of the *AluYb8* at the Pan4 locus, the *AluYb* lineage still retained a very low retrotransposition activity (~1.5 fixed copies/Myr in the chimpanzee genome) until the major expansion of the *AluYb* lineage in the human genome within the past 3–4 Myr (Carter et al. 2004; Xing et al. 2004).

The “stealth driver” model of *Alu* evolution

The long evolutionary history of *AluYb* lineage leads to the conclusion that the *AluYb* lineage has remained in the genome with little or no retrotransposition activity for an extended period of time, while retaining the ability to generate an appreciable num-

ber of new copies later in a species-specific manner (Fig. 6). This scenario is different from the classic “master” gene model, in which a master gene is defined as an element that is highly active over long periods of time (Deininger et al. 1992). In general, the amplification dynamics of the *AluYb* lineage show a striking similarity to that of the *AluYa5* lineage (Leeftang et al. 1993; Shaikh and Deininger 1996). Although the existence of low-activity *Alu* source genes has previously been suggested for the *AluYa5* subfamily (Shaikh and Deininger 1996), here we provide evidence that low retrotransposition activity *Alu* source genes should be recognized as a major factor driving *Alu* expansion and evolution.

We propose a model of *Alu* evolution and retrotransposition, in which the low-activity *Alu* elements are termed “stealth drivers.” In contrast to master genes, stealth drivers are not responsible for generating the majority of new *Alu* copies, but rather for maintaining genomic retrotransposition capacity over extended periods of time. By generating new *Alu* copies at a slow rate, a stealth driver may generate some daughter elements that are capable of much higher retrotransposition rates. These hyperactive daughter elements may act as master genes for the amplification of *Alu* subfamilies and are responsible for producing the majority of the subfamily members. In addition, the new master genes may also give birth to additional retrotranspositionally active *Alu* copies that serve as secondary master genes or sprouts and also contribute to the expansion of the subfamily (Deininger and Batzer 1993). This view is supported by the fact that recently integrated human *Alu* subfamilies typically contain ~15% of such secondary master genes, in addition to the primary

Figure 5. Median-joining network of the human-specific *AluYb* elements. The network of the *AluYb* lineage was reconstructed using the 36 non-Yb7/8/9 elements and the consensus sequences of the Yb7, Yb8, and Yb9 subfamilies as representatives of these three subfamilies. Black circles denote sequence types. Reconstructed nodes are identified as empty circles. The size of circles indicates the number of *Alu* loci with this sequence type, while arbitrary sizes were chosen for the Yb7/8/9 nodes to represent the relative sizes of the three subfamilies. Lines denote substitution steps, with a one-step distance being indicated in the bottom-right corner. Broken lines indicate that the length of the branch is not drawn to scale.

subfamily master gene (Cordaux et al. 2004). While the highly active master genes that are produced from the stealth driver would be deleterious and generally be subject to negative selection, the low-activity stealth driver itself will not be subject to such selection and would allow the lineage to persist for extended periods of time. Furthermore, both stealth driver and its daughter elements will also generate new elements that exhibit low levels of retrotransposition activity, effectively becoming new stealth-driver elements. In fact, it is possible that the continuation of a lineage over extended periods of time may depend on the production of new driver elements.

But the question remains as to why, after persisting relatively quietly for millions of years, a lineage can show a sudden increase in numbers, as appears to be the case for *AluYa* and *AluYb* lineages. Under the stealth-driver model, the master or stealth locus does not need to be “turned on,” as one or more such drivers has been active during the entire history of the lineage. There are multiple scenarios that may account for the sudden expansion of the *Alu* lineage. In the scenario that we favor, periods of rapid expansion may be related to the ability of highly active daughter elements to escape selection at the population level and, consequently, produce more progeny. The change in the efficiency of natural selection in weeding out overactive elements may be related to population bottlenecks or other demographic factors (Hedges et al. 2004). Alternatively, periodic increases in element numbers may simply be due to the stochastic nature by which active daughter elements are produced. Yet another possibility is that, contingent on the relative abundance of available L1 retrotransposition machinery, there may be fluctuation in *Alu* expansion rates (Dewannieux et al. 2003). Less likely is the possibility that molecular host defense mechanisms, which were previously suppressing the activity of *Alu* elements, failed for some reason during these periods.

In the *AluYb* lineage, multiple lines of evidence suggest that

the *AluYb8* element at the Pan4 locus may be a recent stealth-driver gene in human-specific *Alu* retrotransposition. First, the Pan4 element is the oldest *AluYb8* element we identified; second, it only accumulated one point mutation over the last 7 Myr; third, the presence of three identical human *AluYb8* elements is consistent with recent low levels of retrotransposition of this element. On the other hand, since the Pan4 *AluYb8* element has accumulated five mutations in the chimpanzee and gorilla genome, and there are no other *AluYb* elements in these genomes with the same mutations, the *AluYb8* element at Pan4 locus is unlikely to be the current driver gene in the chimpanzee and gorilla genomes. The reasons why the Pan4 *AluYb8* element may be a stealth driver in human, but not chimpanzee and gorilla, are unclear. However, there is a striking correlation between sequence similarity to the *AluYb8* consensus sequence and total number of *AluYb* copies in these species, raising the possibility of mutational inactivation of the Pan4 *AluYb8* element in chimpanzee and gorilla, but not in the human lineage.

Previous studies have shown that the amplification of the *Alu* family reached its peak about 30 Mya and subsequently underwent retrotranspositional quiescence (Shen et al. 1991; Britten 1994). Although we still do not know the underlying mechanisms for the retrotranspositional quiescence, the stealth-driver model may explain why the *Alu* lineage has been subjected to periods of retrotranspositional quiescence interspersed with episodic bursts of amplification, as suggested by the accumulation of at least 5000 human-specific *Alu* elements since the human-chimpanzee divergence (Carroll et al. 2001; Batzer and Deininger 2002; Xing et al. 2003; Carter et al. 2004; Otieno et al. 2004). The *AluYa* and *AluYb* lineages that comprise >60% of the human-specific *Alu* elements may be just two successful examples of this strategy. Similar patterns of amplification have also been observed in the retrotransposition of rodent SINE family, ID, (Kim et al. 1994) and the rodent LINE (Long Interspersed Element) family, Lx (Pascale et al. 1993). In addition to these evolutionary observations, there is also experimental evidence that indicates

Figure 6. Putative evolutionary scenario for the *AluYb* lineage. A schematic diagram of the hominid primates is shown with the approximate time scale shown on the bottom in million years. The blue lines indicate the expansion of the *AluYb* lineage, and the thickness of the lines represents its relative retrotransposition activity. The estimated copy number of *AluYb* elements in various primates is shown after their names. The blue triangle represents the estimated integration time period of *AluYb* founder gene, and the star represents the estimated integration time period of the oldest known *AluYb8* element (Pan4).

that some varieties of mobile elements evolve strategies to attenuate their own activity. In *Alu*, the acquisition of a second monomer to form its dimeric structure has been linked to decreased retrotranspositional activity (Li and Schmid 2004). In vitro modifications to currently active L1 elements can produce orders of magnitude increases in the L1 amplification rate (Han and Boeke 2004). In addition, cryptic polyadenylation sites throughout the L1 sequence may serve to quell the number of full-length, retrotransposition-competent L1 copies generated (Perepelitsa-Belancio and Deininger 2003). Taken together, this evidence suggests that the stealth-driver model may not be a unique feature of the *Alu* family itself, but rather may be one variant of a common survival strategy for SINE and LINE elements. More generally, the ability of mobile elements to maintain low to moderate levels of amplification activity, rather than more rapid duplication rates, may be a common feature of long-lived, successful families of transposons.

Methods

Computational identification of *AluYb* elements

A 31-bp (TGCGCCACTGCAGTCCGCAGTCCGGCCTGGG) oligonucleotide that included the *AluYb* lineage-specific duplication was used to screen the common chimpanzee genome draft sequences (panTro1 Nov. 2003 assembly, <http://www.ncbi.nlm.nih.gov>; R. Waterston, pers. comm.) using the Basic Local Alignment Search Tool (BLAST) program available at <http://www.ensembl.org/multi/blastview> (Altschul et al. 1990). All *Alu* elements that have the diagnostic 7-bp duplication were selected and extracted along with 1000 bp of unique DNA sequence adjacent to both ends of the elements. The program RepeatMasker (<http://repeatmasker.genome.washington.edu/cgi-bin/RepeatMasker>) was then used to annotate all known repeat elements within the DNA sequence. Flanking oligonucleotide primers for the PCR amplification of each *Alu* element were then designed using Primer3 (http://www-genome.wi.mit.edu/cgi-bin/primer/primer3_www.cgi). The primers were subsequently screened against the GenBank NR database using BLAST queries to determine whether they resided in unique DNA sequences and would only amplify the *Alu* elements of interest.

Identification of *AluYb* elements with PCR display

The *Alu* element PCR display methodology has been reported previously (Ray et al. 2005). Using this approach, 500 ng of genomic DNA was partially digested using restriction endonucleases *NdeI* or *MseI* as recommended by the manufacturer (New England Biolabs) in 120 μ L reactions. Digestion products were then ligated with double-stranded linkers and amplified and "displayed" using the primer LNP (5'-GAATTCGTC AACATAG CATTCT-3') and an *AluYb*-specific primer (5'-GGCCGGA CTGCGGACT-3') to acquire partial *Alu* sequences and the accompanying flanking unique sequences from each template. Since the *Alu* sequence in the amplicon is about 300 bp long, the PCR products were then purified by BD CHROMAS SPIN-400 columns (BD Biosciences) to select the fragments larger than 400 bp, so that enough unique flanking sequence can be obtained to locate the orthologous sequences in the draft sequence of the human genome. A second-round amplification was performed using the LNP oligonucleotide and a second nested *AluYb*-specific primer (5'-AATCTCGGCTCACTGCAAGCTCCGCT-3') to increase the specificity of the amplicons. The second-round PCR products were separated on a 2% agarose gel, and fragments

larger than 400 bp were excised and extracted from the gel using the Wizard gel-purification kit (Promega). The purified products were then cloned into the TOPO-TA cloning vector (Invitrogen). At least 72 clones were randomly isolated from each template, and DNA sequences were determined from both strands using chain-termination sequencing (Sanger et al. 1977) on an ABI 3100 automated DNA sequencer.

After obtaining the sequences, the BLAST-Like Alignment Tool (BLAT) program (<http://genome.ucsc.edu/cgi-bin/hgBlat?command=start>) (Kent 2002) was used to compare the resulting partial *AluYb* element and its adjacent flanking sequence with human draft sequences to identify orthologous sequences. Using a combination of the unique flanking sequence and the orthologous human DNA sequences, oligonucleotide primers were designed around each newly identified *AluYb* element as outlined above.

PCR analysis of *AluYb* elements

All of the *AluYb* loci were screened on a panel composed of human genomic DNA (HeLa cell line ATCC CCL-2) and DNA samples from the following nonhuman primate species: *Pan troglodytes* (common chimpanzee), *Pan paniscus* (bonobo or pygmy chimpanzee), *Gorilla gorilla* (lowland gorilla), *Pongo pygmaeus* (orangutan), *Hylobates syndactylus* (siamang), *Hylobates lar* (white handed gibbon), *Chlorocebus aethiops sabaues* (green monkey), and *Aotus trivirgatus* (owl monkey). The nonhuman primate DNA is available as a primate phylogenetic panel (PRP00001) from the Coriell Institute for Medical Research. The chimpanzee-specific *AluYb* loci were also screened for insertion presence/absence using a common chimpanzee population panel composed of 12 unrelated individuals of unknown geographic origin, which was provided by the Southwest Foundation for Biomedical Research.

PCR amplification of each locus was performed in 25- μ L reactions using 10–50 ng of target DNA, 200 nM of each oligonucleotide primer, 200 μ M dNTP's in 50 mM KCl, 1.5 mM $MgCl_2$, 10 mM Tris-HCl (pH 8.4), and 2.5 units *Taq* DNA polymerase. Each sample was subjected to an initial denaturation step of 94°C for 150 sec, followed by 32 cycles of 1 min of denaturation at 94°C, 1 min of annealing at optimal annealing temperature, 1 min of extension at 72°C, followed by a final extension step at 72°C for 10 min. Resulting PCR products were fractionated on a 2% agarose gel with 0.25 μ g of ethidium bromide and visualized using UV fluorescence.

DNA sequence analysis

To confirm the presence of *AluYb* elements, all PCR products suggesting the presence of an *Alu* element were gel purified using the Wizard gel-purification kit (Promega). Purified PCR products were then cloned into vectors using the TOPO TA cloning kit (Invitrogen) and sequenced using chain-termination sequencing (Sanger et al. 1977) on an Applied Biosystems 3100 automated DNA sequencer. All clones were sequenced in both directions to confirm the sequence. The DNA sequences generated in this study are available in the GenBank under accession numbers AY791249–AY791290.

Acknowledgments

We thank Dr. David Ray for critical reading and suggestions during the preparation of this manuscript. We also thank Anthony Carter for providing the human *AluYb* sequences. This research was supported by the Louisiana Board of Regents Millennium Trust Health Excellence Fund HEF (2000-05)-01 (M.A.B.), Na-

tional Science Foundation BCS-0218338 (M.A.B.) and EPS-0346411 (M.A.B.), National Institutes of Health RO1 GM59290 (M.A.B.), and the State of Louisiana Board of Regents Support Fund (M.A.B.).

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Received November 19, 2004; accepted in revised form January 28, 2005.

Under the genomic radar: The Stealth model of *Alu* amplification

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Genome Res. 2005 15: 655-664

Access the most recent version at doi:[10.1101/gr.3492605](https://doi.org/10.1101/gr.3492605)

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