

Presence and Phylogenetic Relationships of a Hominoid-Specific Retroposon Family on the Human Y Chromosome

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ABSTRACT—The retroposon SINE-R.C2 was first identified as a human-specific insertion in the complement C2 gene and the insertion of a closely related element has been implicated as a cause of Fukuyama type muscular dystrophy. In a previous study, we found SINE-R-type retroposons, derived from the endogenous retrovirus HERV-K family, have been found to be hominoid specific. In this report, we identified twelve SINE-R-type retroposons from the human Y chromosome by a PCR approach. Compared with other human retroposons (SINE-R.C2, 11, 14, 19, and HS307/HS408), the Y-chromosomal retroposon showed a high degree of sequence homology (86–96%). Phylogenetic analysis using the neighbor-joining method revealed that SINE-R-type retroposons on the Y chromosome were inter-related with hominoid primates, suggesting that SINE-R elements on the Y chromosome have evolved progressively in the hominoid radiation. Two elements SINE-R.Y-2 and Y-7 are more closely related to the human-specific retroposon SINE-R.C2 than other Y-chromosomal retroposons. Such elements thus are likely to be of great potential relevance to recent events in hominoid evolution and to structural change or genetic variation connected to various disease.

INTRODUCTION

It has often been suggested that endogenous retroviruses and retroposons have played a role in influencing the functional organization of the human genome (Baltimore, 1985; Kazazian and Moran, 1998; Sverdlov, 1998). Approximately 1% of the human genome is comprised of such sequences, and most of them have been integrated into the primate lineage thirty-three or more million years ago (Haltmeier *et al.*, 1995; Mager and Freeman, 1995; Steinhuber *et al.*, 1995). The vast majority of these elements were incorporated as proviral DNA which then rapidly became replication-defective by random accumulation of mutations to be vertically transmitted as junk DNA. Therefore, particular interest attaches to families of elements that have changed recently in human and hominoid evolution. By PCR and phylogenetic analyses, Medstrand and Mager (1998) found that some solitary long terminal repeat (LTR) elements of the human endogenous retrovirus K family (HERV-K) were human specific. These elements may still be actively transposing in the human genome. For example, a HERV-K LTR element has been inserted in the antisense orientation between the gag and pol gene of the human endogenous retrovirus S71 (Leib-Mösch *et al.*,

1993). Another solitary element has also been found at the HLA-DQB1 (Seidl *et al.*, 1999). It has been estimated that there are 25,000 copies of such elements in the human genome (Leib-Mösch *et al.*, 1993), and they are potentially capable of affecting the expression of closely located genes (Akopov *et al.*, 1998). Simon *et al.* (1994) found that HERV-K-related LTR elements are expressed in human placenta.

Amongst the derivatives of the HERV-K10 retrovirus family is a family of Short Interspersed Nuclear Elements (SINEs) termed SINE-R for retrovirus-derived (Ono *et al.*, 1987). SINE-R11, 14, 19 were isolated by colony blot hybridization with a HERV-K10 probe, and derive from the 3' long terminal repeat and small upstream flanking regions. These elements entered in the genome of hominoid primates after the split of Old World monkeys in the Oligocene period (Kim *et al.*, 1999a, b). The SINE-R.C2 element was discovered in relation to a variable number of tandem repeat (VNTR) sequence within the third intron of the gene for C2, the second component of complement, located in class III of the major histocompatibility complex on the short arm of human chromosome 6 (Carroll *et al.*, 1984; Zhu *et al.*, 1992). By Southern blot analysis, SINE-R.C2 is confined to *Homo sapiens* (Zhu *et al.*, 1994). Within the block in Xq21.3 with homology to Yp11.2 that has been tentatively linked to schizophrenia and schizo-affective disorder (Laval *et al.*, 1998) we previously reported two retroposons (HS307 and HS408) with 92 to 96% homology to SINE-R.C2 (Kim *et al.*, 1999b). An element almost identical in sequence

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to SINE-R.C2 has recently been reported as the cause of Fukuyama-type muscular dystrophy (Kobayashi *et al.*, 1998), and this appears to be the first association of this class of elements with disease. Influences of SINE-R elements on gene expression have been described, for example, a SINE element that acquired a role in signal transduction in the course of mammalian evolution (Shimamura *et al.*, 1998). Here we report the presence of SINE-R-type retroposons on the human Y chromosome and their phylogenetic analysis with those of other humans and hominoids.

MATERIALS AND METHODS

Sample DNA and PCR

A human monochromosomal somatic cell hybrid DNA panel was obtained from the MRC Human Genome Mapping Project Resource Centre (HGMP-RC, Cambridge, UK, Kelsall *et al.*, 1995). The Y chromosomal DNA sample was subjected to PCR amplification. SINE-R-type retroposons were amplified by the primer pair JS902 (5'-GAGAATGGGCCATGATGAC-3', bases 4–22) and JS903 (5'-GATCATTCTTGGATGTTTCT-3', bases 466–485) from schizophrenic brain S11 cDNA (GenBank, accession no. AA772777) which has a high level of sequence homology with SINE-R.C2, 11, 14, 19, and HS307/HS408 elements (Ono *et al.*, 1987; Kim *et al.*, 1999b). The PCR conditions followed were those of Kim *et al.* (1996) with an annealing temperature of 56°C.

Cloning of PCR product

PCR products were separated on a 1.8% agarose gel, purified with the QIAEX II gel extraction kit (Qiagen) and cloned into the T-khs307 vector (Kim *et al.*, 1998). Plasmid DNA was isolated by the alkali lysis method using a High Pure plasmid isolation kit (Boehringer Mannheim).

DNA sequencing and data analysis

Individual plasmid DNAs were screened for inserts by PCR. Positive samples were subjected to sequence analysis on both strands with T7 and M13 reverse primers using an automated DNA sequencer (Model 373A) and the DyeDeoxy terminator kit (Applied Biosystem). SINE-R.C2 (L09706), SINE-R11 (X07417), SINE-R14 (X07418), SINE-R19 (X07419), FMR-1 (L29074), HS307 (AB016781), HS408 (AB016782), SCHIZO (AA772777), Chi-M (AB018364), Chi-F (AB018365), Gor-M (AB018366), Gor-F (AB018367), Ora-M (AB018368), Ora-F (AB018369), Gib-M (AB018370), and Gib-F (AB018371) sequences were taken from the GenBank database. Sequence analyses were done with the aid of GAP, PILEUP, and PRETTY from the GCG program (University of Oxford). Neighbor-joining phylogenetic analysis (Saitou and Nei, 1987) was performed with the CLUSTAL W program (Thompson *et al.*, 1994). Sequence homology was searched for in the BLAST network server (Altschul *et al.*, 1997).

RESULTS AND DISCUSSION

The nucleotide sequences of the human endogenous retrovirus, HERV-K10, have been found to be present in hominoids and Old World monkeys by PCR analysis using genomic DNA (Zhu *et al.*, 1994). SINE-R-type retroposons, that are clearly derived from the long terminal repeat (LTR) of the HERV-K family (Ono *et al.*, 1987; Zhu *et al.*, 1992), have been found to be hominoid-specific by PCR analysis (Kim *et al.*, 1999b). Comparing nucleotide sequences between SINE-R-type retroposons and HERV-K LTR elements, we showed that the

367 bp of SINE-R-type retroposons are deleted from the HERV-KLTR sequences (Ono *et al.*, 1987; Kim *et al.*, 1999b). By BLAST search, cDNA prepared from the brain of a patient with schizophrenia (GenBank, accession no. AA772777) was found to have sequences containing a high degree of homology with the 3' LTR of HERV-K10 (Ono *et al.*, 1986), SINE-R11, 14, 19 (Ono *et al.*, 1987), SINE-R.C2 (Zhu *et al.*, 1994), and HS307/HS408 (Kim *et al.*, 1999b). Therefore, PCR primers, JS902 and JS903, were chosen from the brain cDNA of a patient with schizophrenia. Using genomic DNA from the human Y chromosome, we identified twelve SINE-R-type retroposons by PCR amplification in the present study although we are aware that there are more SINE-R-type retroposons on the Y chromosome. Ono *et al.* (1987) have estimated that SINE-R elements were present at 4000 to 5000 copies per haploid human genome. So far, limited numbers of SINE-R-type retroposons (SINE-R11, 14, 19, C2, and HS307/HS408) have been reported in the human genome (Ono *et al.*, 1987; Zhu *et al.*, 1994; Kim *et al.*, 1999b). As far as we know, this is the first time that the nucleotide sequences of Y-chromosomal retroposons have been demonstrated. The sequences are presented in alignment with those of the other human retroposon family in Fig. 1. The sequence data from the human Y chromosome was demonstrated as derived from SINE-R-type retroposons by comparison with other human retroposons. The hominoid specificity of the PCR products was also demonstrated by the absence of amplified rodent DNA (data not shown). As shown in Table 1, Y-chromosomal retroposons have a high degree of sequence homology with other retroposons. Sequence similarity ranged from 99.5% between SINE-R.Y-7 and SINE-R.Y-2, also between SINE-R.Y-15 and Y-17, to 85.6% between SINE-R.Y-4 and SINE-R.Y-15. The human-specific retroposon SINE-R.C2 shared 96% sequence homology with both SINE-R.Y-2 and Y-7.

To understand the evolutionary relationships among these families with hominoid retroposons, a phylogenetic tree was constructed by the neighbor-joining method. As shown in Fig. 2, the sequence of the chimpanzee male (Chi-M) is closely related to the sequences from SINE-R.Y-8 and SINE-R.Y-4, while SINE-R.Y-15 and Y-17 cluster with the sequence from the gibbon male (Gib-M) and female (Gib-F), the schizophrenic brain cDNA (SCHIZO), and the orangutan female (Ora-F). Retroposons Y-3, 11, 13, 16, 18, 19 are closely related to SINE-R14. SINE-R.Y-7 and SINE-R.Y-2 are more closely related to each other than either of them is to SINE-R11 and they cluster with SINE-R.C2. The human-specific retroposon SINE-R.C2 is closely related to the sequence of gorilla male (Gor-M) and these sequences cluster with the HS307 retroposon that we previously identified in the Xq21.3 region of homology with the Y chromosome (Kim *et al.*, 1999b). This group with SINE-R.C2 is likely to be of great relevance to recent events in hominoid evolution. Thus the sequences of SINE-R elements on the Y chromosome are intermingled with those of similar elements in hominoid primates, suggesting that several lineages of retroposons have been evolving independently during hominoid evolution.

The non-recombining portion of the human Y chromo-

CONSENSUS	ATGGCGGTTT	TGT-GAATAG	AAAAGGGGGA	AATGTGGGGA	AAAGATAGAG
	1				50
SINE-RC2	-----	---G-----	---G-----	--G-----	-----T--
SINE-R11	-----	---G-----	---G-----	--G-----	-----T--
SINE-R14	-----	---C-----	---G-----	-----	G---GG---
SINE-R19	-----	---C-----	-----T--	-----	-----C--
SCHIZO	-----	---C---A--	-----	-----	-----A--
FMR-1	-----A-C--	---G---G--	---G-C---	--G-----	GG---T---
HS307	-----T---	---G---G--	---G-----	--G-----	-----T---
HS408	-----A---	---C---G--	-G-G-----	-----	GGG---G---
Y-2	-----	---G-----	-----	--G-----	-----T---
Y-3	-----	GCCGA-..-A	G-----	-----	-----A---
Y-4	-----	---G-----	---G-----	--G-----	-----G---
Y-7	-----	---G-----	-----	--G-----	-----T---
Y-8	-----	---G-----	---G-----	--G-----	-----G---
Y-11	G-----	---C---A--	-----	-----	-----A---
Y-13	-----	---C-----	-----	-----	-----A---
Y-15	-----	--CC---A--	-----	-----	-----A---
Y-16	-----	---C-----	-----	-----	-----A---
Y-17	-----	--CC---A--	-----	-----	-----A---
Y-18	-----	---C-----	-----	-----	-----A---
Y-19	-----A---	---GA-----	-----	-----A--	-----A---
CONSENSUS	A-ATCAGATT	GTT-CTGTGT	CTGTGTAGAA	AGAAGTAGAC	ATGGGAGACT
	51				100
SINE-RC2	-A---G---G	---G-C---	-----	---G-----	C-----
SINE-R11	-A---G---G	---G-----	-----	-----	-----
SINE-R14	-G---G-T---	---A-----	-----G---	G--G-----	--TA-----
SINE-R19	-A-----A	---G-----	-----	-----	--A--G--T-
SCHIZO	-G-----	---A---C--	-----	-----	--A-----
FMR-1	-AG-----A	---G-C---	-----G---	G-----	-----
HS307	-A-A-G---G	---G-C---	-----	-----	-----T---
HS408	-GG-----	---G-----	-----G	G-----	--A---T---
Y-2	-A---G---G	---G-----	---A---	-----	-----
Y-3	-G-----	T--A-----	---C---	-----	--A-----A
Y-4	-A-----	---G-----	-----	---C---	-----.
Y-7	-A---G---G	---G-----	---A---	-----	-----
Y-8	-A-----	---G-----	-----	---C---	-----.
Y-11	-A-----	---A-----	-----	-----	--A-----
Y-13	-G-----	---A-----	-----A--	-----	--A--G---
Y-15	-G-----	T--A-----	---C---	-----	--A-----A
Y-16	-G-----	---A-----	-----A--	-----	--A--G---
Y-17	-G-----	T--A-----	---C---	-----	--A-----A
Y-18	-G-----	---A-----	-----A--	-----	--A--G---
Y-19	-G-----	---A-----	-----C-	-----	-----
CONSENSUS	-CACTTTGTT	GTTCTGTACT	AAGAAAAATT	CTTCTGCCTT	GGGATGCTGT
	101				150
SINE-RC2	TTT-A--..--	-----	-----	-----	-----C---
SINE-R11	TTT-A--..--	-----	-----	-----T---	-----C---
SINE-R14	C--T---..	-----	-G-----	-----	-----
SINE-R19	C--T---..	-----	-----	-----	-----
SCHIZO	C--T---..	-----	-----	-----	-----
FMR-1	TTT-A--..--	-----C---	GG--G-----	-C-----	-----C---
HS307	TTT-A--..--	-----	-----	-----G-	-----C---
HS408	C--T---..	-----	-----	-----	-----
Y-2	TTT-A--..--	-----	-----	-----	-----C---
Y-3	C-----..	-----	-----	-----G	-----
Y-4	.TT-A--..--	--C-----C	-G--GGGG--	--G-----	-----
Y-7	TTT-A--..--	-----	-----	-----	-----C---
Y-8	.TT-A--..--	-----	GG--GGGG--	--G-----	-----
Y-11	C--T---..	-----	-----	-----	-----
Y-13	T--T-----	C-----	-----	-----	-----
Y-15	C-----..	-----	-----	-----G	-----
Y-16	T--T-----	C-----	-----	-----	-----
Y-17	C-----..	-----	-----	-----G	-----
Y-18	T--T-----	C-----	-----	-----	-----
Y-19	C--T---..	-----	-----	-----	-----CA-

CONSENSUS	TAATCTATGA	CCTTACCCCC	AACCCTGTGC	TCTCTGAAAC	ATGTGCTGTG
	151				200
SINE-RC2	-G--GG--	-----	-----	-----	-----A
SINE-R11	-G-----	-----	-----	-----	-----
SINE-R14	-----A-	-----	-----C--	-----	-----
SINE-R19	-G-----	-----	-----G--	-G-----	-----
SCHIZO	-----A-	-----	-----CC--	-----	-----
FMR-1	-G--G--	---G----	-----	-----	-----
HS307	GG--G--	-----	-----	-----	-----
HS408	-----A-	-----	-----	-----	-A-----
Y-2	-G-----	-----	-----	-----	-----A--
Y-3	-----A-	--A-G----	-----C--	-----	-----
Y-4	-G--G--	---G----	-G-----	-----GGG-	-----A
Y-7	-G-----	-----	-----	-----	-----
Y-8	-G--G--	---G----	-G-----	-----GGG-	-----
Y-11	-----A-	-----	-----	-----	-----
Y-13	-----	-----	-----C--	-----	-G-----
Y-15	-----A-	--A-----	-----C--	-----	-----
Y-16	-----	-----	-----C--	-----	-G-----
Y-17	-----A-	--A-----	-----C--	-----	-----
Y-18	-----	-----	-----C--	-----	-G-----
Y-19	-----A-	-----	-----CC--T	-----	-----
CONSENSUS	TCCAATCAGG	GTAAATGGA	TTAAGGGCGG	TGCAAGATGT	GCTTTGTTAA
	201				250
SINE-RC2	-----	---G-----	-----A--	-----	-----
SINE-R11	-----	-----	-----	-----	-----
SINE-R14	-----	-----AT--	-----	-----	-----
SINE-R19	-----	-----	-----	-----	-----
SCHIZO	--A-----	-----	-----	-----	-----
FMR-1	-----	-----	-----	-----	-----
HS307	-----	-----	-----	-----	-----
HS408	-----	-----	-----	-----	-----
Y-2	-----	-----	-----	-----	-----
Y-3	--A-----	-----	-----	-----	-----
Y-4	-----	-----	-----T--	-----	-----
Y-7	-----	-----	-----	-----	-----
Y-8	-----	-----	-----T--	-----	-----
Y-11	-----	-----	-----T--	-----	C-----
Y-13	-----	-----	-----	-----C--	-----
Y-15	--A-----	-----	-----	-----	-----
Y-16	-----	-----	-----	-----	-----C--
Y-17	--A-----	-----	-----	-----	-----
Y-18	-----	---G-----	-----	-----	-----
Y-19	-----	-----	-----	-----	-----
CONSENSUS	ACAGATGCTT	GAAGGCAGCA	TGCTCGTTAA	GAGTCATCAC	CACTCCCTAA
	251				300
SINE-RC2	-----	-----	-----C--	-----	-----
SINE-R11	-----	-----	-----	-----	-----
SINE-R14	-----	-----	-----	-----	-----
SINE-R19	-----	-----	---A-T--	-----C--	-----
SCHIZO	-----	---A-----	-----C--	-----	-----
FMR-1	-----	-----	-----	-----	--A-----
HS307	-----	-----	-----C--	-----	-----
HS408	-----	-----	---A-----	-----	-----
Y-2	-----	-----	---A-----	-----	-----
Y-3	-----	-----	---T-----	-----	-----
Y-4	-----C-	-----	-----	-----	-----
Y-7	-----	-----	---A-----	-----G-	-----
Y-8	-----	-----	-----	-----	-----
Y-11	-----	-----	-----G--	-----	-----
Y-13	-----	---TG----	-----	-----	-----
Y-15	-----	A-----	-----	-----	---T-----
Y-16	-----	---TG----	-----	-----	-----
Y-17	-----	A-----	-----	-----	-----
Y-18	-----	---TG----	-----	-----	---T-----
Y-19	-----	-----	---T-----	-----	-----

CONSENSUS	TCTCAAGTAC 301	CCAGGGACAC	AAACACTGCG	GAAGGCCGCA	GGGTCCTCTG 350
SINE-RC2	-----	-----	-----	-----	-----
SINE-R11	-----	-----	-----	-----	-----
SINE-R14	-----A-----	-----	-----T-----	-----	-----
SINE-R19	-----	-----	-----	-----C-----	-----
SCHIZO	-----	-----T-----	-----GG--T-----	-----	A--A-----
FMR-1	-----A T-----	-----	-----T-----	-----	-----
HS307	-----T-----	-----	-----	-----	-----
HS408	-----	-----	-----C-----	-----T-----	-----
Y-2	-----	-----	-----T-----	-----	-----
Y-3	-----	-----	-----C-----	-----A-----	-----
Y-4	-----	-----	-----A-----	-----T-----	-----
Y-7	-----	-----	-----T-----	-----	-----
Y-8	-----	-----	-----A-----	-----T-----	-----
Y-11	-----	-----	-----T-----	-----	-----
Y-13	-----	-----TG-----	-----	-----C-----	-----
Y-15	-----	-----	-----A--C-----	-----	-----AA-----
Y-16	-----	-----TG-----	-----	-----C-----	-----
Y-17	-----	-----	-----A--C-----	-----	-----AA-----
Y-18	-----	-----TG-----	-----	-----C-----	-----
Y-19	-----	-----	-----C-----	-----A-----	-----
CONSENSUS	CCTAGGAAAA 351	CCAGAGACCT	TTGTTCACCT	GTTTATCTGC	TGACCTTCCC 400
SINE-RC2	-----	-----	-----	-----	-----
SINE-R11	-----	-----	-----	-----	-----
SINE-R14	-----	-----	-----A-----	-----	-----
SINE-R19	-----	-----	G-----	-----	-----T-----
SCHIZO	-----	-----	-----TG-----	-----	-----TT-----
FMR-1	-----	-----	-----	-----	-----
HS307	-----	-----T-----	-----	-----	-----C-----
HS408	-----	-----	-----	-----	-----T-----
Y-2	-----	-----	-----	-----	-----T-----
Y-3	-----	-----G-----C-----	-----A-----	A-----	-----
Y-4	-----	A-CAGA-A-----	-----	-----	-----
Y-7	-----	-----	-----	-----	-----T-----
Y-8	-----	-----	-----	-----	-----
Y-11	-----	-----C-----	-----A-----	-----G-----	-----
Y-13	-----	-----	-----TA-----	-----	-----T-----
Y-15	-----	G-----	-----G-----	-----	-----TT-----
Y-16	-----	-----	-----TA-----	-----C-----	-----T-----
Y-17	-----	G-----	-----G-----	-----	-----T-----
Y-18	-----	-----	-----TA-----	-----	-----TT-----
Y-19	-----	-----G-----C-----	-----A-----	A-----	-----
CONSENSUS	TCCACTATTG 401	TCCTATGACC	CTGCCAAATC	CCCCTCTCTG 442	--
SINE-RC2	-----	-----G-----	-----	-----G-----	..
SINE-R11	-----	-----	-----	-----G-----	..
SINE-R14	-----	-----	-----	-----	..
SINE-R19	-----G-----	-----A-----	-----A-----	-----GC-----	..
SCHIZO	-----A-----	-----	-----C-----	-----C-----	..
FMR-1	-----	-----C-----	-----	-----G-----	..
HS307	-----T-----	-----	-----	-----GC-----	..
HS408	-----	-----	-----	-----GC-----	..
Y-2	-----	-----C-----	-----	-----G-----G-----	..
Y-3	-----	-----	-----	-----	..
Y-4	-----T-C-----	-----	-----	-----CTGT-----	GC
Y-7	-----	-----C-----	-----	-----G-----G-----	..
Y-8	-----	-----	-----	-----G-----	..
Y-11	-----A-----	-----	-----T--C--T-----	-----	..
Y-13	-----A-----	-----	-----C-----	-----	..
Y-15	-----A-----	-----	-----C-----	-----	..
Y-16	-----	-----	-----	-----	..
Y-17	-----A-----	-----	-----C-----	-----	..
Y-18	-----	-----	-----	-----	..
Y-19	-----	-----	-----	-----	..

Fig. 1. Nucleotide sequence alignments of the retroposon family. Consensus sequences are shown on the top row. Dashes indicate no change to the consensus sequence and dots indicate gaps. The nucleotide sequence data reported in this paper for Y-chromosomal retroposons will appear in the DDBJ/EMBL/GenBank nucleotide sequence databases with the accession numbers AB025287-AB025298.

Table 1. Percentage similarity of nucleotide sequences in the retroposon family

	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20
1. SINE-RC2	—																			
2. SINE-R11	97.3	—																		
3. SINE-R14	92.2	93.6	—																	
4. SINE-R19	92.4	94.5	91.1	—																
5. SCHIZO	90.2	92.0	92.0	91.1	—															
6. FMR-1	93.2	94.1	90.2	89.9	87.6	—														
7. HS307	95.9	96.4	90.8	92.0	89.7	92.3	—													
8. HS408	91.8	93.4	93.1	92.0	91.5	92.0	91.3	—												
9. Y-2	96.1	98.2	92.7	93.3	90.8	93.2	95.2	92.9	—											
10. Y-3	89.6	91.7	91.7	90.3	92.4	88.0	89.2	90.8	90.8	—										
11. Y-4	90.4	91.3	87.9	87.6	86.0	89.0	89.5	88.3	89.9	86.2	—									
12. Y-7	96.1	98.2	92.7	93.3	90.8	93.2	95.2	92.9	99.5	90.8	89.9	—								
13. Y-8	93.4	94.7	91.1	90.8	89.0	92.9	92.4	91.5	93.3	89.2	96.1	93.3	—							
14. Y-11	91.5	93.8	93.4	91.5	93.4	89.0	90.6	91.5	92.4	92.6	87.9	92.4	91.1	—						
15. Y-13	90.2	92.5	92.9	92.0	94.1	87.7	89.3	91.5	91.1	91.9	87.2	91.1	90.4	93.8	—					
16. Y-15	89.2	91.5	91.5	89.9	94.5	87.0	88.3	90.6	90.4	94.7	85.6	90.4	88.6	92.9	93.1	—				
17. Y-16	90.4	92.7	93.1	92.2	93.4	87.9	89.5	91.8	91.3	92.2	87.4	91.3	90.6	93.1	98.9	92.4	—			
18. Y-17	89.7	92.0	92.0	90.4	95.0	87.4	88.8	91.1	90.8	95.2	86.0	90.8	89.0	93.4	93.6	99.5	92.9	—		
19. Y-18	90.7	92.5	93.0	92.0	93.6	87.7	89.3	91.5	91.1	91.9	87.2	91.1	90.4	92.9	98.6	92.7	98.9	92.7	—	
20. Y-19	91.3	93.3	92.9	91.3	92.9	88.8	90.6	91.5	92.2	95.2	87.4	92.2	90.4	93.6	92.9	91.7	93.1	92.2	92.9	—

some includes relatively few genes and has the capacity to accumulate repetitive and other junk DNA sequences unconstrained by the functional considerations that apply elsewhere. It shows considerably greater variation between hominoid species (Glaser *et al.*, 1997). However, particular interest attaches to regions of homology between the X and Y chromosomes. Such homologies have generally been created by X to Y transpositions and a number of such events have been tracked in the course of mammalian and primate evolution (Lambson *et al.*, 1992; Glaser *et al.*, 1997). Genes within these regions are thus exposed to new evolutionary pressures, and their X homologues are subject to differential patterns of inactivation (Jegalian and Page, 1998). Some evolutionary changes are known to have been influenced by retroposons, eg the LINE-LINE recombination associated with the paracentric inversion on the Y chromosome (Schwartz *et al.*, 1997) and they may be the most recent chromosomal rearrangement in human evolution. Such X-Y homologous regions are relevant to the detection of a role of retroviruses/retrotransposons in recent evolution in two respects. First, examination of homologous sequences on the X and the Y may reveal new insertions and allow them to be dated. Second, investigation of the relationship between genes within these regions and such elements as may be related to them could uncover a role in gene expression since any recent change will be differentially expressed in the two sexes.

The Xq21.3/Yp11.2 region of homology is of particular interest in relation to human evolution, as it was created by a transposition after the separation of the chimpanzee and human lineages, and 3.5 Mb of the transposed region was subject to a subsequent paracentric inversion (Sargent *et al.*, 1996; Mumm *et al.*, 1997). We identified two retrotransposons (HS307 and HS408) within the Xq21.3 region with homology to the SINE-R.C2 element (Kim *et al.*, 1999b). It will be important to examine the homologies of these elements within the

Yp11.2 region to ascertain whether changes in these or other elements (eg SINE-R.Y-2 or Y-7) might be associated with genes which in turn are relevant to the sex differences that have been described in relation to *Homo sapiens* specific cognitive functions (Crow *et al.*, 1998).

SINE-R11, 14 and 19 have a polypurine tract (PPT) that serves as a primer-binding-site for plus-strand DNA synthesis from retrovirally mediated reverse transcription. This indicates that SINE-R-type elements like human endogenous retroviruses have a potential functional role in the genomic DNA of human chromosomes. According to PCR and phylogenetic analyses, integration of several HERV-K LTR elements have been shown to be human specific (Medstrand and Mager, 1998). We also identified a solitary HERV-K LTR element in a recently proliferated class of LTR elements, this was inserted into Xq26 after the separation of the hominid and great ape lineages (Kim *et al.*, submitted). It seems that the LTR elements have been randomly transposed across the chromosome in the course of evolution. Apokov *et al.* (1998) noted that such sequences have the capacity to modify the expression of neighboring genes. These elements may also have contributed to genetic variation associated with various disease. The HERV-K sequences, to which the SINE-R elements are related, have been reported as a candidate autoimmune gene in type I diabetes (Conrad *et al.*, 1997). Two HERV-K LTR elements have been detected in the human major histocompatibility complex locus HLA-DQ (Kambhu *et al.*, 1990). One LTR element (DQ-LTR3) of the HERV-K family at the HLA-DQB1 locus has been associated with the rheumatoid arthritis (Seidl *et al.*, 1999).

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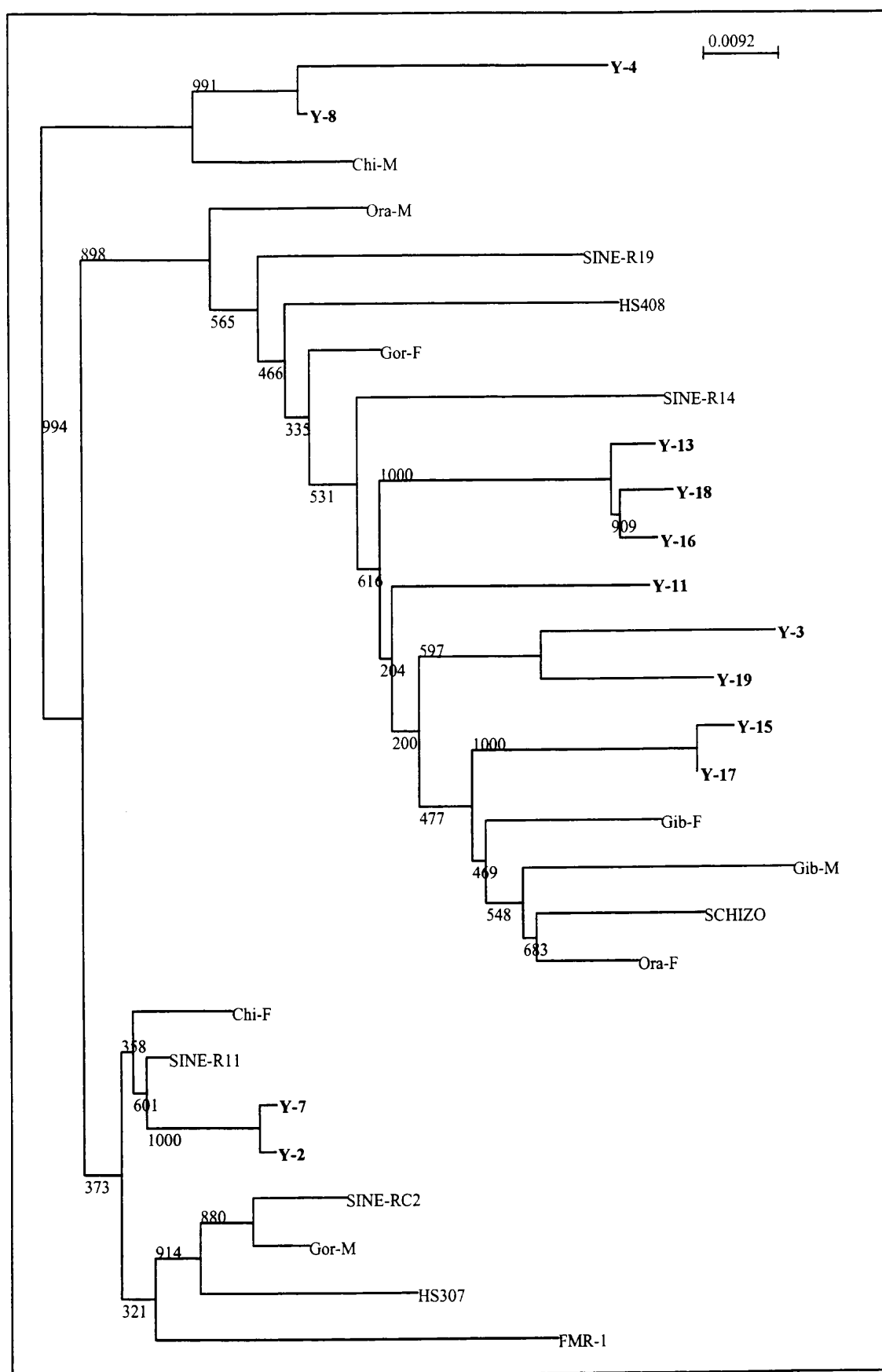


Fig. 2. Phylogenetic tree obtained by neighbor-joining method for the retroposon family. Statistical robustness was tested by bootstrap analysis. The numbers to the left of the nodes of the tree branches represent the number of times out of 1000 trees. The retroposons (Chi-M, Chi-F, Gor-M, Gor-F, Ora-M, Ora-F, Gib-M, Gib-F) from hominoid primates were taken from GenBank database (see Materials and Methods).

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