Localization of *mariner* DNA Transposons in the Human Genome by PRINS

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Homologous recombination occurring among misaligned repeated sequences is a significant source of the molecular rearrangements resulting in human genetic disease. Studies of the Charcot-Marie-Tooth disease locus on chromosome 17 have implicated the involvement of an ancient DNA transposon of the *mariner* family (Hsmar2) in the initiation of double-strand break events leading to homologous recombination. In this study, the genomic locations of 109 Hsmar2 elements were determined by primed in situ labeling (PRINS) using primers designed to match the right and left inverted terminal repeats (ITRs) of the transposon. Although the resolution of the PRINS technique is ~400 chromosomal Giemsa bands, the data presented here provide the first large-scale mapping study of these elements, which may be involved in initiation of homologous recombination events in the human genome.

The human genome is composed not only of DNA sequences coding for genes but also of a significant amount of noncoding or "junk" DNA. However, this noncoding DNA is not a static collection of insignificant DNA sequences. There appears, rather, to be a state of genome flux occurring among these sequences even in the span of a single human generation. Genome evolution models involving gene duplication and exon shuffling have been bolstered by the identification of active L1 elements in the human genome that can transpose not only portions of their own open reading frames but also fragments of surrounding coding sequence to new locations (Kzazian and Moran 1998). Short pericentromeric repeats on both human and other primate chromosomes may be involved in the transposition of duplicated exons or even entire genes to new locations (Eichler et al. 1997). In addition to these possibilities, the most extensively documented method of gene sequence rearrangement in the genomes of eukaryotes is homologous recombination. This kind of rearrangement is often mediated by region specific low-copy repeated sequences (Lupski 1998). Analysis of a well-documented hotspot for homologous recombination in the human genome revealed the presence of a *mariner* DNA transposon near the site of strand exchange for this recombination event (Kiyosawa and Chance 1996; Reiter et al. 1996). It was proposed that this element, and possibly copies of it located elsewhere in the genome, could be involved in the stimulation of homologous recombination among

repeated sequences through the introduction of double-strand breaks in DNA near the region of strand exchange (Kiyosawa and Chance 1996; Reiter et al. 1996, 1998).

The presence of *mariner* DNA transposons of the *irritans* subfamily (Hsmar2) in the human genome is a rather recent discovery (Oosumi et al. 1995; Kiyosawa and Chance 1996; Reiter et al. 1996; Robertson and Martos 1997). Multiple copies of Hsmar2 elements have been identified in the EST (dbEST), nonredundant (nr), and high-throughput genomic sequencing (htgs) databases (Robertson and Martos 1997). Many of these copies are deleted for one inverted terminal repeat (ITR) and, therefore, are unlikely to bind a transposase, which appears to require two ITRs in concert for binding and cleavage. Although these elements appear to be inactive remnants of a functional *mariner* that entered the primate lineage some 80 million years ago, studies of homologous recombination at the Charcot-Marie-Tooth disease type 1A locus on chromosome 17p12 suggest that the presence of an Hsmar2 element within two flanking 24-kb repeats (CMT1A-REP) may stimulate unequal crossing-over events between misaligned CMT1A-REP elements (Kiyosawa and Chance 1996; Reiter et al. 1996, 1997, 1998). These unequal crossing-over events can be resolved as either a 1.5-Mb duplication resulting in Charcot-Marie-Tooth disease or a 1.5-Mb deletion resulting in hereditary neuropathy with liability to pressure palsies (Pentao et al. 1992; Chance et al. 1994). Chromosomal duplications and deletions mediated by large region-specific low-copy-number repeats may prove to be a common

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mechanism of chromosomal rearrangement leading to disease in the human genome (Lupski 1998) and have been implicated in the molecular mechanisms of a growing number of human genetic disorders such as Smith-Magenis syndrome (Chen et al. 1997), Prader-Willi/Angelman syndrome (PWS/AS) (Christian et al. 1995; Robinson et al. 1998), and Williams-Bueren syndrome (WS) (Urban et al. 1996; Pérez Jurado et al. 1998). Although a direct role for *mariner* elements in initiating recombination via the stimulation of double-strand breaks remains unclear, we were interested in determining if other Hsmar2 elements containing both a left and right ITR could be found at sites in the human genome where structural changes occur by an unequal crossing-over mechanism. To determine the genomic locations of elements containing both a left and right ITR among the ~1000 copies of Hsmar2 in the human genome we localized these elements to specific chromosomal bands by primed in situ (PRINS) labeling.

RESULTS

Identifying the Locations of Hsmar2 Elements in the Human Genome

To locate Hsmar2 elements in the human genome we designed 50-bp oligonucleotides from the 3' and 5' ends of the consensus sequence for Hsmar2 (GenBank accession no. U49974). These oligonucleotides were used as primers for the fluorescence in situ hybridization (FISH)-related technique known as PRINS (Luke and Shepelsky 1998). The oligonucleotides were annealed to a male human metaphase chromosomal spread and extended with a mixture containing biotin-labeled deoxy-uracil triphosphate (dUTP). The biotin-labeled nucleotides were then detected using a thymid detection system and viewed under the fluorescence microscope (Fig. 1A). Each chromosome was analyzed independently from 15 different chromosomal spreads to determine the number of signals per chromosome and their positions. Figure 1B shows this type of analysis for chromosome 17, which contains the CMT1A-REP region, the location of the ele-

ment we previously called MITE for *mariner* insect transposon-like element. A total of 109 signals were detected using the left ITR oligonucleotide and 108 signals using the right ITR oligonucleotide. These signals are visually summarized in Figure 1C as the culmination of all signals detected on each chromosome. Note that the transposable elements appear to be spread evenly throughout the genome. This is apparent both by visual inspection of integration sites (Fig. 1C) and by correlation of the number of signals per chromosome with chromosome length. However, no elements were detected on the Y chromosome by this method. Table 1 summarizes the cytogenetic locations of all 109 positions of signals detected by PRINS, presumably representing one or more Hsmar2 elements, using both the left and right ITR oligonucleotide primers.

To demonstrate that the PRINS technique allows the detection of both repeated and unique sequences, control experiments were performed using a telomere-

Figure 1 PRINS signals using the left and right ITRs of Hsmar2. (A) Raw PRINS data. This is an unenhanced chromosomal spread of the biotin-labeled left ITR. The green signals represent locations of Hsmar2 elements that contain left ITRs that match the consensus for Hsmar2 with enough homology to promote annealing and priming for the PRINS signal. (B) An example of the analysis of PRINS signals on chromosome 17. Twelve independent PRINS experiments are represented here by the 12 individual chromosome 17 images. The green signals are PRINS labeling using the left ITR. Note that not all chromosomes show all of the signals and that some chromosomes only show a signal for one homolog. This limitation of the technique requires the analysis of multiple chromosomal spreads. The signals indicated by numbers 1–4 are located at 17p12, 17q12, 17q22, and 17q24–25, respectively, at a resolution of 400 chromosomal Giemsa bands. (C) Summary of all left and right ITR PRINS data. This is a karyotype representing all loci that contain Hsmar2 elements by PRINS. The image was constructed by combining the PRINS signals for each individual chromosome into a composite image that was then added to other chromosomal images and arranged into the karyotype.

Table 1. Cytogenetic Locations of Hsmar2 Elements Detected by PRINS

Chromosome	No. of Signals									
1	7	1p36	1p35	1p32	1p22	1q22-23	1q31-32	1q41-42		
2	9	2p24-25	2p16	2p14	2q12	2q21	2q24	2q32	2q34	2q36
3	7	3p24-25	3p21	3p13	3q13	3q21	3q25-26	3q28		
4	8	4p16	4p15	4p12	4q12-13	4q22-24	4q28	4q31	4q34	
5	6	5p15	5p13-14	5q11-12	5q14	5q23-31	5q34			
6	6	6p24-25	6p21	6q12	6q16	6q24	6q26-27			
7	7	7p21	(7p14-15)	7p12	7q11	7q21	7q31	7q32	7q35-36	
8	5	8p23	8p12-21	8q13-21	8q23	8q24				
9	5	9p23-24	9p12-13	9q12	9q22	9q33-34				
10	5	10p12-13	10q21.1	10q21.3	10q24-25	10q26				
11	5	11p15-ter	11p13-14	11q13	11q22	11q24				
12	6	12p13	12p11	12q12	12q15-21	12q21-22	12q24			
13	3	13q14	(13q12-13)	13q21-22	13q33					
14	3	14q12	14q24	14q32						
15	2	15q12-14	15q23-24							
16	3	16p13	16q21	16q23-24						
17	4	17p12	17q12	17q22	17q24-25					
18	4	18p11	18q12-12	18q22	18q22-23					
19	3	19p13	19q13.1-13.2	19q13.3						
20	2	20p13	20q13							
21	2	21q22	21q13							
22	2	22q11	22q13							
X	5	Xp22	Xp11-21	Xq13	Xq25-26	Xq28				
Y	0									
$\Sigma =$		109								

The locations of PRINS signals illustrated in Fig. 1C are summarized, as well as the locations of Hsmar2 elements detected by BLASTN searches of the nonredundant nucleotide database. Loci detected by PRINS alone are plain text, by BLASTN are boldface, and by BLASTN alone are in parentheses. The one PRINS signal that was detected by the left but not the right ITR oligonucleotide is underlined.

specific oligonucleotide and a chromosome X-specific oligonucleotide. The telomere-specific oligonucleotide detected only the telomeric repeat regions of all human chromosomes (Fig. 2A) whereas the X chromosome-specific oligonucleotide only detected a region on the X chromosome (Fig. 2B).

BLAST Searches and Genomic Disorders

As an independent confirmation of the ability to detect complete Hsmar2 elements by the PRINS technique, BLASTN searches were performed using the 50-bp left and right ITR oligonucleotides against the current nonredundant nucleotide database. Thirteen matches were identified in this database by BLASTN (Table 1, bold). These matches contained a left and right ITR and extended at least 22-bp with a perfect match for the last four 3'-end bases. These criteria were presumed to be enough homology to promote priming and extension. Only 2 of the 13 Hsmar2 elements detected by the BLASTN searches were not also detected by PRINS (Table 1, parentheses). Interestingly, within the limits of resolution of the technique (~400 chromosomal Giemsa bands) the locations of PRINS signals co-

incide with the chromosomal map locations of twelve known genomic disorders that result from a homologous recombination event within misaligned repeated

Figure 2 Confirmation of the PRINS technique specificity. (A) Telomere-specific signals. A 50-bp oligonucleotide designed to anneal to the repeated sequences of human telomeres was used for PRINS on a female human metaphase chromosomal spread. The green PRINS signals were detected at the telomeres of all human chromosomes. (B) X chromosome centromere-specific signal. A 50-bp oligonucleotide designed to anneal to a portion of the human X chromosome located at the centromere was used for PRINS on a female human metaphase chromosomal spread. Only two green PRINS signals were detected. These signals were located at the centromeres of the X chromosomes.

DNA sequences. The names of these diseases, chromosomal locations, types of rearrangement, and lengths of the repeating unit are summarized in Table 2.

DISCUSSION

We have demonstrated by PRINS that multiple copies of the *mariner* element Hsmar2 are spread throughout the human genome. At least 108 of these copies appear to contain both a left and right inverted terminal repeat (Table 1). The genomic locations of these elements were easily identified by PRINS prior to the completion of the human genome project. Although there is still no direct evidence for the involvement of *mariner* elements in the stimulation of homologous recombination in humans, we provide here some preliminary evidence that the locations of these elements may coincide with at least 12 locations where homologous recombination is the proposed molecular mechanism responsible for the human disease phenotypes (Table 2). It should be noted, however, that at this resolution the correlation between Hsmar2 elements and genomic disorders is not statistically significant. Further studies of the submicroscopic regions containing the elements identified in this study will be necessary to determine if *mariner* has a role in promoting homologous recombination in humans.

The Hsmar2 element previously called MITE (Reiter et al. 1996) was identified in the CMT1A region (17p12). In addition, an Hsmar2 element was identified in the PWS/AS region (15q13) where large deletions are also a common disease mechanism (Christian et al. 1995; Robinson et al. 1998) and in the WS region (7q11) where flanking repeat sequences have been implicated as a molecular mechanism for disease (Urban et al. 1996; Pérez Jurado et al. 1998). Further studies of these traveling *mariner* genes in the human genome may not only illuminate our understanding of how molecular rearrangements involving repeated DNA se-

quences can cause human disease but also may reveal the manner in which the genome itself continues to evolve.

METHODS

PRINS Labeling and Detection

Primers used for PRINS analysis were designed based on the published consensus sequence for Hsmar2 constructed by Hugh Robertson (accession no. U49974). The left ITR primer begins at the 5' end of Hsmar2 and ends at base 50 of the consensus sequence whereas the right ITR primer was designed from the complementary strand of Hsmar2 beginning at base 1309 and ending 50 bases internal to the element at base 1279. Primers designed to the open reading frame of the Hsmar2 element were not used because this region is highly variable among the elements examined in the databases. Peripheral blood lymphocytes from a healthy male donor were used for PRINS analysis using the left and right ITR oligonucleotides. Lymphocytes from a healthy female donor were used for the X-chromosome and telomere-specific oligonucleotides. Cells in metaphase were denatured on microscope slides in denaturation solution (80 μ l of deionized formamide, 10 μ l of ddH₂O, 10 μ l of 20 $\times$ SSC) covered with a 24 $\times$ 50-mm coverslip and heated to 75°C on a heating plate for 5 min. The coverslip was then removed and the slides were subjected to a series of ethanol washes (70%/90%/100%) and air dried. The 50- μ l PRINS reaction solution was then added to each slide: 3 μ l each of 10 mM dATP, dCTP, and dGTP; 1 μ l of 10 mM dTTP; 3 μ l of 10 mM biotin-dUTP; 1 μ l of 1 pmol/ μ l primer; 1 μ l 10 $\times$ BSA; 2 μ l 25 mM MgCl₂; 0.5 μ l *Thermus aquaticus* (Taq) polymerase; 10 $\times$ Taq polymerase buffer; 27.5 μ l ddH₂O. All reagents were from Pharmacia except for biotin-dUTP and BSA (Boehringer Mannheim). Slides were then sealed with a coverslip and rubber cement by warming to 40°C. Incorporation of labeled dUTP was accomplished by initial denaturation of the DNA for 5 min at 94°C followed by a 50-min annealing/extension period at 58°C for the ITR primers, 55°C for the telomere specific primer, and 60°C for the X-chromosome primer. The reaction was stopped by removing the cover slip and soaking the slides at 60°C in 500 ml of stop solution (50 mM EDTA, 5 mM NaCl in ddH₂O). Detection of the biotin labeled oligonucleotides was performed us-

Table 2. Genomic Disorders That Coincide with the Locations of Hsmar2 Elements Detected by PRINS

Genomic disease	Type of rearrangement	Length of repeat (bp)	Locus
21-Hydroxylase deficiency	deletion	(?) ^a	6p21.3
Williams-Bueren syndrome	deletion	>30,000	7q11.2
Glucocorticoid-remediable aldosteronism	duplication	10,000	8q21
Prader-Willi/Angelman syndrome	deletion	(?)	15q13
CMT1A/HNPP	duplication/deletion	24,000	17p12
Growth hormone deficiency	deletion	2,200	17q22-24
Debrisoquine sensitivity	deletion	2,800	22q13.1
X-Linked ichthyosis	deletion	20,000	Xp22.32
α -Thalassemia	deletion	4,000	Xq13
Color blindness	deletion	39,000	Xq28
Hunter mucopolysaccharidosis	inversion	3,000	Xq28
Hemophilia A	inversion	9,500	Xq28

^a(?) The length of the repeat involved is unknown.

ing the thymid detection system (TSA-Kit; MEL731A, NEN-Dupont) according to the manufacturer's instructions. Metaphase spreads were counterstained with DAPI (diamidinophenylindol) solution, 20 μ l of the antifade Vectashield (Vectorlabs) was added to the slide, and it was covered with a coverslip. An enhanced DAPI banding pattern was derived by 5-bromo-2-deoxyuridine (BrdU) incorporation into the chromosomes during cultivation of the lymphocytes used for PRINS (Heng and Tsui 1993). The DAPI banding pattern indicated the exact subchromosomal localizations of the probes. Evaluation was done on a Zeiss Axioplan fluorescence microscope and pictures were taken using a CCD-camera and the software of MetaSystems (Altlußheim, Germany).

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