

Identification of *TSIX*, Encoding an RNA Antisense to Human *XIST*, Reveals Differences from its Murine Counterpart: Implications for X Inactivation

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X inactivation is the mammalian method for X-chromosome dosage compensation, but some features of this developmental process vary among mammals. Such species variations provide insights into the essential components of the pathway. *Tsix* encodes a transcript antisense to the murine *Xist* transcript and is expressed in the mouse embryo only during the initial stages of X inactivation; it has been shown to play a role in imprinted X inactivation in the mouse placenta. We have identified its counterpart within the human X inactivation center (*XIC*). Human *TSIX* produces a >30-kb transcript that is expressed only in cells of fetal origin; it is expressed from human *XIC* transgenes in mouse embryonic stem cells and from human embryoid-body-derived cells, but not from human adult somatic cells. Differences in the structure of human and murine genes indicate that human *TSIX* was truncated during evolution. These differences could explain the fact that X inactivation is not imprinted in human placenta, and they raise questions about the role of *TSIX* in random X inactivation.

Introduction

X inactivation is the mammalian method for equalization of the dosage of X-linked genes in males and females; this is accomplished by down-regulating the transcriptional output of X chromosomes in females, so that only one X is active in diploid somatic cells of both sexes (Lyon 1962). This transcriptional silencing is initiated in female embryonic blastocysts and can be induced in murine embryonic stem (ES) cells. Inactivation is manifested by underacetylated histones, methylated CpG islands, and delayed DNA replication, as reviewed by Heard et al. (1997). The initial events responsible for a single active X in mammalian cells are largely unknown, but some players have been identified. The key control region on the X chromosome, from which inactivation is initiated and signals are spread, has been mapped to Xq13.2 and is referred to as the “X inactivation center” (*XIC* in humans and *Xic* in mouse). X chromosomes can be programmed to inactivate because of the X inactive specific transcript gene (*XIST* in human and *Xist* in mouse [MIM 314670]) located within this region. The expression of *XIST* starts a cascade of events that re-

sults in silencing of the chromosome from which it is transcribed.

Prior to the time that an inactive X is seen in female embryos, *XIST* is expressed at low levels from all X chromosomes, even from the single X in male embryos (Ray et al. 1997). Therefore, it seems likely that an early step in this developmental pathway is one that ensures that the *XIST* locus on one X chromosome is turned off in cells of both sexes; in this way, one and only one X chromosome will be functional. The *XIST* allele that is repressed in female cells is usually chosen randomly; all other X chromosomes in the cell are inactivated by default, via propagation of the *XIST*-induced inactivating signal(s) along the chromosome. The choice is nonrandom in mouse placental tissues, where the *Xist* locus on the maternal X is always repressed; hence, the maternal X is always active. The *XIST* RNA is untranslated and remains in the nucleus, associated with the inactive X (Clemson et al. 1996). The stabilization and accumulation of this RNA around the inactive X is thought to propagate the inactivating signal along the chromosome (Panning et al. 1997).

Clearly, *XIST* is not the only player in this process; if it were, there would be no active X chromosome, since all X chromosomes would be silenced. Something else is needed to repress the *XIST* locus on the active X in both sexes. The X controlling element (*Xce*), a locus downstream of murine *Xist*, affects the randomness of inactivation in a manner not yet known (Simmler et al. 1993). The orthologous locus has not been identified on the human X chromosome. In addition, Lee et

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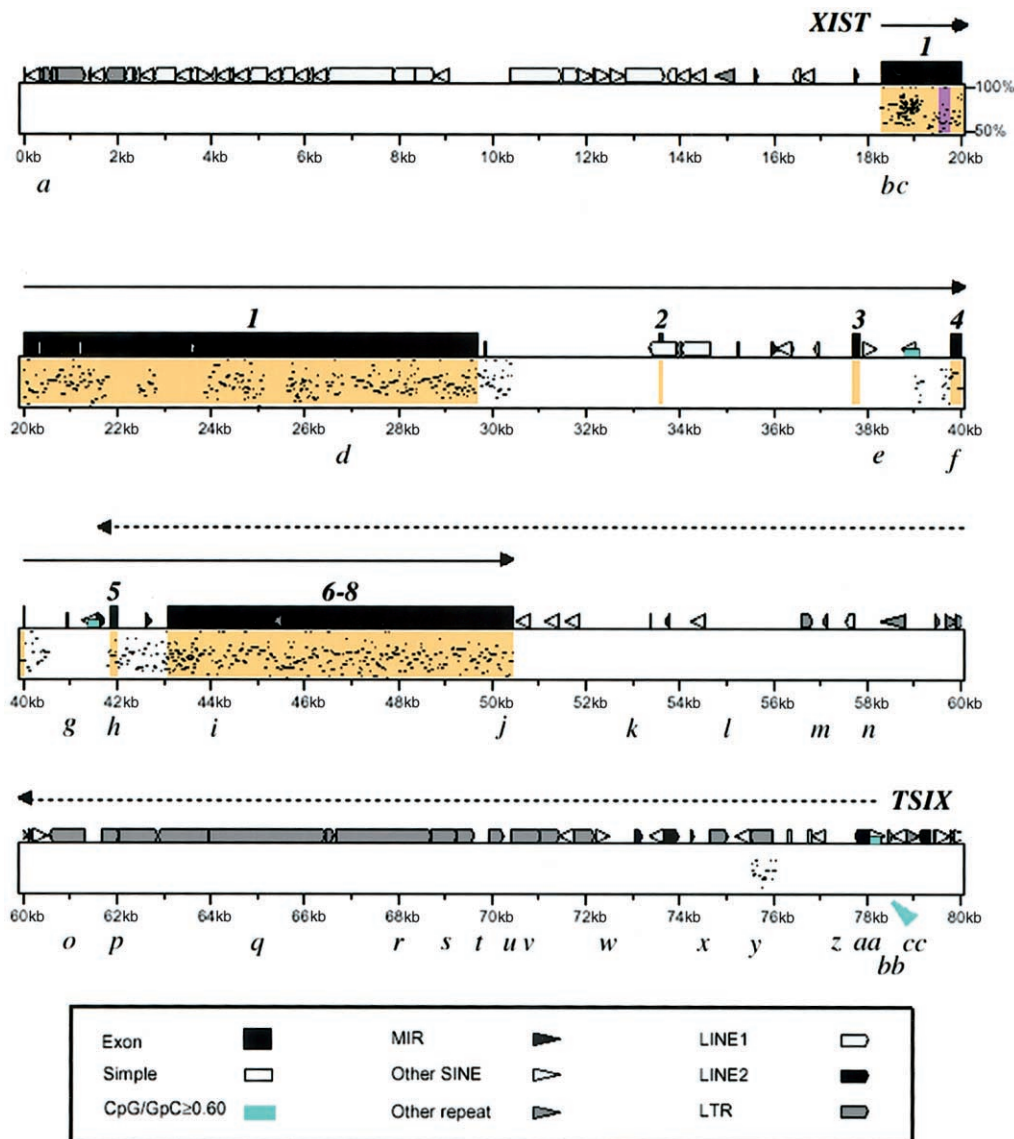


Figure 1 PipMaker percent identity plot of the human XIC region in ES-10, showing the location of a novel transcript and homology to the comparable region in the murine genome. Nucleotides 0–80000 of the U80460 sequence (0–80 kb) are shown. Plotting of *XIST* exons (orange) is based on all known transcribed sequences (Brown et al. 1992; Hong et al. 2000). The black dot pattern shows the percent homology (50%–100%) with the comparable mouse *Xic* region. The *XIST* transcript is shown as a solid line with an arrow. The *TSIX* transcript (dotted line with arrow) is antisense to *XIST* and is initiated from four clustered transcription start sites, the most 5' of which shown (see fig. 3). The purple vertical bar in *XIST* exon 1 represents the single CpG island identified by the Grail program (version 1.3). The small blue boxes indicate a CpG:GpC frequency of ≥ 6 ; the one near the *TSIX* start site is marked by a blue arrow. The locations of RT-PCR primers (a–cc) are shown in lowercase letters. The lack of homology outside of the *XIST* region has also been documented in a PipMaker plot by Nesterova et al. (2001, fig. 5e). MIR = mammalianwide interspersed repeats; SINE = short interspersed elements.

al. (1999) discovered *Tsix* (MIM 300181), the gene encoding an RNA that is, in part, antisense to the murine *Xist* transcript. *Tsix* was initially described as a 40-kb transcript starting downstream of *Xist* and overlapping the entire *Xist* locus. More recently, Sado et al. (2001) have found a processed *Tsix* transcript in murine ES cells. Like *Xist*, *Tsix* has no open reading frames (ORFs), and its RNA is found in the nucleus. However,

unlike *Xist*, the *Tsix* RNA is associated with the future active X. There is some evidence that the *Tsix* transcript may be an antagonist of *Xist*. Studies of mice carrying a *Tsix* deletion show that *Tsix* is required for repression of the maternal *Xist* allele in placental tissues where the paternal X chromosome is preferentially silenced. Lee (2000) and Sado et al. (2001) have proposed that *Tsix* is the maternally expressed factor that protects the ma-

ternal X from imprinted X inactivation in extra-embryonic tissues. The role of *Tsix* in random inactivation remains to be elucidated (this will be discussed below).

We recently showed that it was feasible to study the human *XIC* region through use of human transgenes introduced into murine ES cells, as well as through chimeric mice derived from them. Six tandem copies of one transgene with 480 kb of the human *XIC* region, including the *XIST* locus, inserted into mouse chromosome 11 was able to induce random X inactivation in male chimeric mice, resulting in inactivation of the murine X chromosome in some XY cells carrying the transgene (Migeon et al. 1999). The other, a single-copy transgene with <300 kb of the human *XIC* region carried on mouse chromosome 6, was recognized as an *XIC* in male ES cells but could not induce inactivation in mice (Migeon et al. 2001), presumably because of the low copy number (Heard et al. 1999). Unlike *XIST/Xist*, which is expressed in all cells with an inactive X at any age, murine *Tsix* is expressed only transiently, prior to and during the time that X inactivation is being established. Murine ES cells recapitulate the stages of X inactivation seen in the mouse embryo proper (Lee and Lu 1999) and are an accepted model for studies of this developmental process. Therefore, to determine whether there was a human gene orthologous to *Tsix*, we looked for other human transcripts in the mouse ES cells carrying the human *XIC* transgene. As a result, we have identified a transcript that is antisense to *XIST* and overlaps the *XIST* locus at its 3' end. This transcript is also present in human-embryo-derived stem cells but is absent from adult somatic cells. On the basis of our observations, we suggest that human *TSIX* is not involved in genomic imprinting and may not be an antagonist of human *XIST* expression.

Material and Methods

Transgenes and Cell Lines

Mouse ES cells carrying the ES-10 and ES-5 transgenes were derived from mouse J1 ES cells and have been reported elsewhere (Migeon et al. 1999, 2001). ES-10 contains six copies of the full transgene, and ES-5 contains a single copy of a truncated one. In addition, we studied XY somatic cells from chimeric mouse embryos 949.1 and 850.3, which carry the ES-10 and ES-5 transgenes on autosomes and are hereafter referred to as "ESch10" and "ESch5," respectively. The murine ES cell transfectants were kept undifferentiated on feeders in medium with leukemia inhibitory factor (LIF). For differentiation, they were passaged twice without feeders on gelatinized dishes in the presence of LIF and then were cultured in suspension without LIF, for aggregation into embryoid bodies (EBs).

The human embryonic cells used for analysis were four genetically distinct cell lines originating from primordial germ cells at 5–11 wk postfertilization (pf). These fibroblast cultures were established from human EBs by J. Gearhart and are referred to as human "embryoid-body-derived" (EBD) cells. They are no longer totipotent but do transcribe some genes that are expressed early in development (Shamblott et al. 2001). Cell lines LV (5 wk pf) and SD (11 wk pf) were 46, XX, whereas SL (6 wk pf) and LU2 (7 wk pf) were 46, XY. Our use of the human EBD cells was approved by the university committee concerned with patents and by Geron, Inc., according to the licensing agreement. They were maintained in EGM2MV medium on collagen plates, as described by Shamblott et al. (2001).

RNA Analysis

Primers.—*XIST* is oriented in the opposite direction of the mouse gene, so that the 3' *XIST* is closest to the centromere. Both the ES-10 and ES-5 transgenes include the ~50-kb human *XIST* locus and ~70 kb centromeric (proximal) to the 3' end of *XIST*. They differ in the amount of sequence telomeric (distal) to the 5' end of *XIST* (360 kb for ES-10 and <180 kb for ES-5). Sequence data for both transgenes are provided in PAC 92E23 (GenBank), which has been sequenced in two parts, found under GenBank accession numbers U80459 and U80460. The locations of primers used for RNA analysis are shown in figure 1 and table 1 and are based on the U80460 sequence. Specific primer sequences will be provided on request.

Reverse-Transcriptase PCR.—RNA was obtained from ES cells before and after differentiation into EBs. Total RNA was isolated with TRIZOL (Life Technologies) and treated with DNaseI (at 37°C for 1 h). Poly A+ RNA was isolated using the Nucleotrap mRNA separation kit (Clontech). To screen for transcripts, reverse-transcriptase PCR (RT-PCR) was performed (1 µg RNA, 35 cycles; Migeon et al. 1999). The cDNA was generated using random priming for sequences outside the *XIST* transcription unit, whereas strand-specific priming was used for the portion of *TSIX* that overlaps *XIST*.

For strand-specific RT-PCR, first-strand cDNA was synthesized using either sense or antisense primers, as described by Lee et al. (1999), with minor modifications. RNA (0.5–1 µg) with strand-specific primer (3 pmol) was incubated at 70°C for 5 min and then was equilibrated to 50°C. The cDNA was synthesized with Superscript II reverse transcriptase (200 U, Life Technologies) at 50°C for 1 h. The RT was heat-inactivated at 80°C for 45 min. After treatment with RNaseH (Life Technologies), first-strand cDNA was amplified with *Taq* polymerase for 35 cycles, using paired sense and antisense primers. Controls containing RNA and strand-

Table 1**RT-PCR Analysis of Human *XIST* / *TSIX* Transcripts**

PRIMER (POSITION) ^a	HUMAN SOMATIC CELL (FEMALE)	MOUSE ES CELL TRANSGENES		HUMAN EBD CELLS (FEMALE)	
		ES-10	ES-5	SD	LV
<i>a</i> (.3 kb)	–	–	–	–	–
<i>b</i> (<i>XIST</i> -pro)	–/–	+/-	–/–	–/–	–/–
<i>c</i> (5' <i>XIST</i>)	+/-	+/-	+/-	ND	+/-
<i>d</i> (<i>XIST</i> -ex1)	+/-	+/-	+/-	+/-	+/-
<i>e</i> (<i>XIST</i> -int3)	–	–	–	–	–
<i>f</i> (<i>XIST</i> -ex4)	+/-	+/-	+/-	+/-	+/-
<i>g</i> (<i>XIST</i> -int4)	ND/–	ND/–	ND/–	ND/–	ND/–
<i>h</i> (<i>XIST</i> -ex5)	+/-	+/+	+/+	+/+	+/+
<i>i</i> (<i>XIST</i> -ex6)	+/-	+/+	+/+	+/+	+/+
<i>j</i> (3' <i>XIST</i>)	+/-	+/+	+/+	+/+	+/+
<i>k</i> (53 kb)	–	+	+	+	+
<i>l</i> (55 kb)	–	+	+	ND	ND
<i>m</i> (57 kb)	–/–	–/+	–/+	+	+
<i>n</i> (58 kb)	–	+	+	+	+
<i>o</i> (61 kb)	–	+	+	ND	ND
<i>p</i> (62 kb)	–	+	+	ND	ND
<i>q</i> (65 kb)	–	+	+	+	+
<i>r</i> (68 kb)	–	+	+	ND	ND
<i>s</i> (69 kb)	–	+	+	ND	ND
<i>t</i> (69.8 kb)	–	+	+	+	+
<i>u</i> (70.4 kb)	–	+	ND	ND	ND
<i>v</i> (70.8 kb)	–	+	+	+	+
<i>w</i> (72.6 kb)	–	+	+	+	+
<i>x</i> (74.5 kb)	–	+	ND	–	–
<i>y</i> (75.5 kb)	–	+	+	–	–
<i>z</i> (77.4 kb)	–/–	–/+	–/+	–	–
<i>aa</i> (77.8 kb)	–	+	+	–	–
<i>bb</i> (78.2 kb)	–	–	–	–	–
<i>cc</i> (79 kb)	–	–	–	ND	ND

NOTE.—Controls (human male and mouse J1 ES cells) were negative for each primer set. + = product present; – = product absent. A single symbol indicates a result of random priming of cDNA, and two symbols (i.e., +/+, –/–, or +/-) indicate sense/antisense (*XIST*/*TSIX*), the results of strand-specific priming. ND = not done.

^a Primers are shown in figure 1. Values shown in kilobases indicate the position in U8046. Pro = promotor, ex = exon, int = intron.

specific primer but no reverse transcriptase were processed in parallel.

5' Rapid Amplification of cDNA Ends (RACE).—Total RNA (10 μg) was treated with RNase-free DNaseI (Ambion). 5' RACE (was performed using the GeneRacer kit (Invitrogen), according to manufacturer instructions. First-strand cDNA was primed with a gene-specific primer (U80460 nucleotides (nt) 77426–77443), was extended with Superscript II at 42°C for 50 min and was treated with RNaseH. 5' ends were amplified using the GeneRacer 5' primer and a gene-specific primer (U80460 nt 77565–77588). Nested PCR was performed with the GeneRacer 5' nested primer and two gene-specific primers (U80460 nt 77578–77603 and U80460 nt 77678–77704). PCR products were either gel purified or were

cloned into the pCR4-TOPO vector (Invitrogen) and were sequenced using [³³P]-labeled chain terminators (Thermo Sequenase Cycle Sequencing Kit, USB).

Sequence Analysis

Once identified, the *TSIX* sequence from U80460 (GenBank) was analyzed for CpG islands, using the Grail program (version 1.3; Oak Ridge National Laboratory Web site). Potential TATA boxes were identified using the HCtata program on the WebGene server. Only strong consensus sequences were considered. We assembled the mouse *Xic* sequence (GenBank accession numbers AJ010350, M97167, L04961, U41394, and X999460) through use of Sequencher software and compared it to the human *XIC* sequence (U80460). Both strands were searched for homology by the Advanced PipMaker (Schwartz et al. 2000; PipMaker Web site). The human sequence was masked using the RepeatMasker program (RepeatMasker Web site). Percent identity plot (PIP) and dot plot graphics from the Advanced PipMaker were modified in Adobe PhotoShop, for clarity of presentation.

Results**Search for Transcripts from the Human *XIC* Transgene in Mouse ES Cells**

Both human *XIC* transgenes include nt 1–119,000 of the U80460 sequence; the *XIST* transcription unit extends from nt 18317 to 50425 (Brown et al. 1992; Hong et al. 2000). Initially, we looked for novel transcripts, using RT-PCR primers at 10-kb increments, starting from U80460 nt 1–119000 (70 kb proximal and 18 kb distal to *XIST*). We also used primers at the distal end of U80459 (50 kb upstream of the 5' end of *XIST*), a region also present in both ES-10 and ES-5 cells.

Figure 1 is a PipMaker PIP (Schwartz et al. 2000) showing the U80460 portion of ES-5 and ES-10 transgenes that includes *XIST* and the only other transcript identified in the region. RT-PCR analysis using cDNA obtained by random priming showed that the novel transcript originates ~27 kb from the 3' end of *XIST*. Table 1 shows that it extended as far as *XIST* exon 5; RT-PCR results were positive with primer set *b* but negative with primer set *g*, 600 bp upstream in *XIST* intron 4. The primers that gave negative results in the RNA analysis were positive in the DNA analysis of the same cells. This transcript was not seen in any of the control cells (human male and female or parental J1 ES cells).

Novel Transcript Is Antisense to *XIST*

Strand-specific priming of the cDNA showed that this transcript was present in poly A+ RNA. This transcript was antisense to *XIST*, since it was amplified from the

cDNA primed by the forward primers (relative to *XIST*) and not from that primed by the reverse primers (fig. 2). Strand-specific priming also showed that this antisense transcript overlapped the *XIST* transcript, extending from 3' *XIST* through exon 5 (table 1 and fig. 1). Because this novel transcript is transcribed from the opposite strand as human *XIST* and overlaps *XIST*, we believe that it is the counterpart of murine *Tsix*; therefore, we will subsequently refer to this gene as “*TSIX*.”

Despite numerous attempts, we have obtained little evidence of a processed *TSIX* transcript. Using a large number of closely spaced primers, which in the *TSIX* unique region were 0.1–3 kb apart, with an average interval of 1.15 kb (fig. 1), we found a contiguous transcript. We did identify a small intron (763 bp) at the start of the transcript in ES-10 (fig. 3; discussed below). In addition, no small transcripts were detected on northern blots.

TSIX Has No ORF

The coding potential of the *TSIX* transcript was investigated *in silico*. The sequence from nt 30000 to nt 80000 of U80460 (complementary strand) was analyzed using ORFgene (Rogozin et al. 1996; ORFgene 2 Web site) and GeneBuilder (Milanesi et al. 1999; GeneBuilder Web site) software. ORFgene did not identify any ORFs

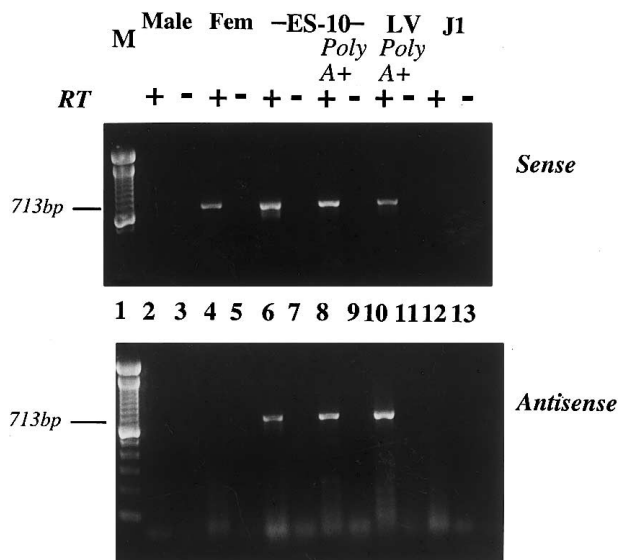


Figure 2 Strand-specific RT-PCR showing polyadenylation and orientation of transcripts. *Top*, Sense (*XIST*) transcript (cDNA synthesized using reverse primer). *Bottom*, Antisense (*TSIX*) transcript (cDNA synthesized using forward primer). RNA samples were total RNA, unless otherwise indicated. Lane 1, 100-bp marker; lanes 2 and 3, male; lanes 4 and 5, female; lanes 6–9, ES-10; lanes 10 and 11, human EBD LV cells; lanes 12 and 13, J1, mouse ES cell control. Samples were assayed using primer set *g* in the presence (+) or absence (–) of reverse transcriptase (RT). Note that ES-10 and LV cells have both sense and antisense transcripts with *XIST* exon 6 primers.

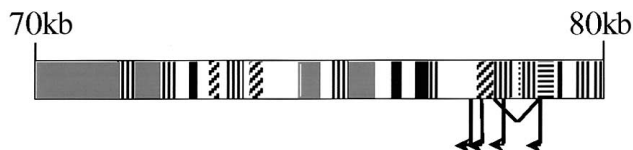


Figure 3 Identification of *TSIX* transcription start sites in ES-10 cells. Transcription start sites (arrows) were determined by 5' RACE using ES-10 as template. The *TSIX* transcript is initiated from four clustered transcription start sites (bases 77657, 77865, 78267, and 78860 in the U80460 sequence). The location of repetitive elements are indicated as follows: black vertical lines denote *Alu* elements, gray boxes denote LTRs, diagonal lines denote LINE elements, and horizontal lines denote MER1 elements.

in this interval. GeneBuilder predicted a series of exons across the interval, which were inconsistent with the results of RT-PCR analysis (table 1 and data not shown). The predicted ORF lacked an initiation codon and showed no homology to any protein in the SWISSPROT database.

TSIX Has Little Homology to Tsix

Figure 1, which is based on the GenBank sequence, shows the homology between the human and mouse DNA sequence in the *XIST-TSIX* region. There are many repeats within this region. Virtually all the significant homology (>50%) lies within the *XIST* locus. The only >50% homology in the region of the *TSIX* transcript is minimal and is limited to a single region at ~76 kb, near the start of the *TSIX* transcript in ES-10; this is most likely the region of human homology to mouse *Tsix* observed by Lee et al. (1999, fig. 6).

Identification of TSIX Transcription Start

5' RACE was used to define the 5' boundary of the *TSIX* transcript in ES-10 cells (fig. 3). Four transcription start sites were identified, at nt 77657, 77865, 78267, and 78860 of U80460. The first three start sites are located within repetitive elements MER58B, *AluY*, and L2, respectively. Transcription from positions 78860 and 77865 may be directed by TATA box sequences within the repeats. An additional upstream exon was identified in the 78860 transcript. The alternate first exon is 23 bases in length, followed by an intron of 763 bases—perhaps the remnant of the small intron between murine *Tsix* exons 2 and 3 (Sado et al. 2001). Splice sites conform to consensus sequences (Shapiro and Senapathy 1987).

Evidence that Human TSIX is Truncated Relative to Tsix

On the basis of the results of 5' RACE studies and homology searches, the start sites in the human gene are at what may be analogous to *Tsix* exons 2 and 3 in the

mouse (Sado et al. 2001) (see region with homology in fig. 1). Unlike murine *Tsix*, which has a CpG island at its 5' end (Lee and Lu 1999), the human *TSIX* gene lacks a proper CpG island (fig. 1). A search of the U80460 sequence by the Grail program (version 1.3) revealed only a single CpG island—the one that overlaps the 5' end of *XIST* (nt 19545–19756; see figs. 1 and 4). Only a small CpG-rich region remains near the *TSIX* start site (fig. 1, arrowed blue box).

To insure that the absence of a CpG island from the 5' end of *TSIX* is not an artifact, the U80460 sequence was compared to that obtained from a whole-genome shotgun assembly of genomic DNA from five unrelated individuals (Venter et al. 2001; Celera Web site). The sequences obtained from the two data sets are colinear over the entire length of U80460, exhibiting 99.96% identity, with the largest discrepancy being a 10-bp insertion/deletion within a pentanucleotide repeat sequence.

The lack of the expected CpG island at the start of *TSIX* suggested an evolutionary breakpoint. Evidence of this comes from the analysis of a neighboring gene. The X-linked testes specific gene (*Tsx*), which resides within 7 kb of murine *Tsix* (Sado et al. 2001), is expressed in rodent testes but has no counterpart expressed in human testes (Simmler et al. 1996). In fact, there is no human ortholog, since only homologous fragments of mouse *Tsx* are present in the human genome. Human *TSX* sequences are scattered in the human X chromosome: exon 1 is in Xq23 and exon 6 is in Xq21. Remnants of exon 5 and exon 4 are present within U80460, but these fragments are 13–20 kb and >50 kb from the 5' end of *TSIX*, respectively (areas 2 and 3, respectively, of fig. 4).

TSIX is Transcribed in Human Embryonic Cells

Because transcripts expressed from our human transgene could have been artifacts of transfection, we were encouraged by the preliminary evidence of a low-level antisense transcript originating “downstream of *XIST* exon 8 and extending only into the 3' end of *XIST*” in the NT2 human embryonic carcinoma line derived from a testicular teratocarcinoma (Chow et al. 2000). To find the *TSIX* transcript that was not present in adult human cells, we chose to examine fetal cells derived from human EBs (Shamblott et al. 2001). To determine the X inactivation status of these cells, we assayed them for *XIST* transcripts. The two female human EBD cell lines (SD and LV) expressed *XIST*, but the male human EBD lines (SL and LU2) did not. Since *XIST* is expressed in male blastocysts prior to the time of X inactivation (Ray et al. 1997), SL and LU2 have passed the developmental stage when *XIST* is expressed. Therefore, both male and female EBD cell lines were at a stage subsequent to the onset of X inactivation.

Since X inactivation had already occurred in these

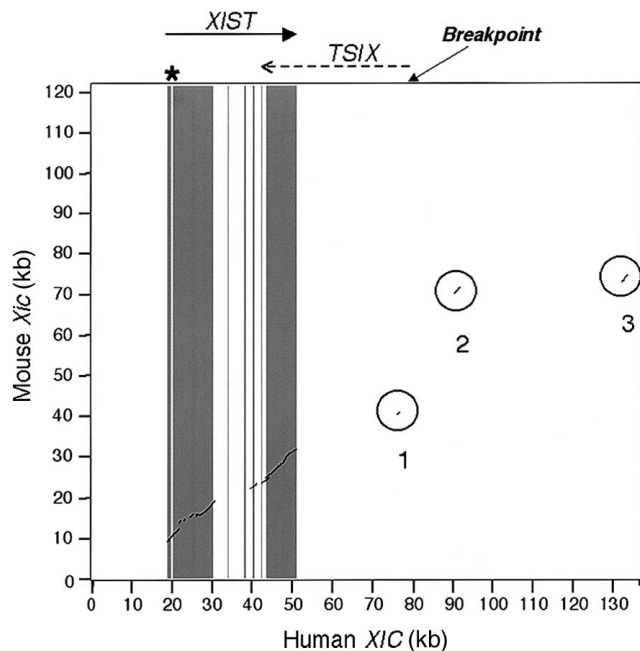


Figure 4 Dot plot showing the evolutionary breakpoint within the human *XIC*. The region shown is an extension of that in figure 1 and includes the region from 0–136 kb of U80460. The dark gray bars represent the *XIST* exons; the light bar shows the CpG island in *XIST* exon 1 (asterisk). The effective evolutionary breakpoint is shown as occurring at the start of the *TSIX* transcript; the actual breakpoint is difficult to determine. The areas circled and numbered 1–3 show the only conserved regions outside of the *XIST* gene. Region 1 is homologous to murine *Tsix*, and regions 2 and 3 are homologous to the murine testis gene (*Tsx*) exons 5 and 4, respectively, which are separated in the human genome by ~40 kb. This figure, a PipMaker dot plot comparing human and mouse *XIC* sequences, is similar to the traditional dot plot of this region, shown by Lee et al. (1999, fig. 6); however, we now know that regions 2 and 3 are homologous to *TSX* and not *TSIX*.

cells, we were surprised to find the *TSIX* transcript in the two female EBD lines. In both of them, the transcript was ~3.8–5.6 kb shorter than that found in ES–10 cells. The start site was difficult to determine, because of the density of repeated sequences in this region (fig. 1), but transcription of *TSIX* most likely originates between nt 73254 and nt 75031 of U80460. The repeated sequences also made it difficult to design primers for 5' RACE. Since primers within the unique sequence failed to generate RACE products, it is difficult to determine the *TSIX* transcription start site in the human EBD cells more precisely. On the other hand, our analysis shows that the 3' end of the *TSIX* transcript was identical to that in ES-5 and ES-10 cells, since it extended only as far as *XIST* exon 5 (table 1).

TSIX is Transcribed in Mouse Embryonic Fibroblasts Carrying the Human Transgene

We examined ESch5 and ESch10 fibroblasts originating from 13.5-d and 18-d embryos chimeric for the ES-

5 or ES-10 transgenes. Both cell lines expressed not only *XIST*, but also *TSIX*. Assayed with *TSIX*-specific primer sets *l*, *n*, *o*, *t*, *w*, *aa*, *bb*, and *cc* (see fig. 1), these fibroblasts had RT-PCR products with all primer sets except *cc*, indicating the same expression pattern as the ES transfectants from which they were derived (cf. ES-5 and ES-10; table 1).

Characteristics of *TSIX* Expression in Embryonic Cells

According to observations of Lee et al. (1999) and Sado et al. (2001), murine *Tsix* is not present in eight-cell embryos, is expressed in blastocysts from both X chromosomes before X inactivation, and is expressed at the onset of inactivation only from the allele on the active X. *Tsix* expression disappears from ES cells during the 2–9 d following their differentiation into EBs. In chimeras, the level of expression is greater in the placenta than in the fetus proper and is absent from both by 12–15 d of gestational age. The expression of human *TSIX* seems to persist longer, at least in a mouse environment, since it is found in ES-10 cells long after their differentiation into EBs (at least to the 19th passage after differentiation) and in the fetal somatic cells derived from 18-d and 13.5-d ES-10 and ES-5 chimeric fetuses, at a time when murine *Tsix* transcripts are minimal or no longer expressed.

Another difference has to do with lack of evidence that human *TSIX* influences *XIST* expression. In the female embryo-derived cells, *TSIX* transcripts were coexpressed with those from *XIST*, so that human *TSIX* expression did not repress the expression of *XIST*. Since the male embryo-derived cells did not express *TSIX*, the transcript is not associated with active X chromosomes in males. Coexpression was also seen in the ES-10 and ES-5 EBs and their chimeric mouse derivatives. In this case, both human transcripts had to come from the transgene, and, hence, from the same chromosome.

Discussion

Comparing Human *TSIX* with Murine *Tsix*

The results of our analysis show that human *TSIX*, like its murine counterpart, is expressed only in embryo-derived cells, initiates downstream of the 3' end of *XIST*, and produces an untranslated RNA. This RNA is transcribed from the opposite strand as *XIST* and is, in part, antisense to *XIST*. However, the human *TSIX* gene differs from its murine counterpart in that it is truncated at the 5' end, does not cover the *XIST* promoter, and does not have the CpG island that is essential for function of murine *Tsix*.

The mouse *Tsix* gene was originally described as a 40-kb RNA without introns, originating 15 kb downstream of *Xist* and transcribed across the *Xist* locus. More-recent studies indicate that it is at least partially processed,

giving a 2.7-kb transcript originating from the major promoter located near *Tsix* exon 2. Sado et al. (2001) also found a 4.3-kb transcript originating from the minor promoter, located ~28 kb downstream of *Xist*, suggesting that *Tsix* starts 13 kb further downstream than originally reported. In addition, these investigators show that exon 2 is associated with a CpG island, which is known to be differentially methylated on active and inactive chromosomes (Prissette et al. 2001). Although the human *TSIX* transcripts from ES-10 and ES-5 cells also begin ~27 kb downstream of *XIST*, they have a much shorter overlap with *XIST*, ending in exon 5; they do not overlap the 5' end of the gene (table 1). The only transcript found in ES-10 in the region of the *XIST* promoter was shown by strand-specific RT-PCR to be attributable to *XIST* and not *TSIX* (table 1 and data not shown). Therefore, the *TSIX* transcript in ES-5 and ES-10 cells is ~35 kb in length. The *TSIX* transcript from human EBD cells has the same limited overlap but is 3.8–5.6 kb shorter; this difference may reflect the influence of the mouse environment on expression of the human transgenes.

Analysis of *TSIX* Identifies *XIC* Breakpoint

All of our evidence suggests that the 5' end of the human *TSIX* locus may have been decapitated by an evolutionary breakpoint. The existence of an *XIC* on Xq13 was originally postulated on the basis of the conserved order of genes on murine and human X chromosomes. However, subsequent studies have revealed many microrearrangements in this region, including an inversion, which affects the transcriptional orientation of the *XIST* locus (Brockdorff et al. 1992). Although the exact site of the breakpoint cannot be determined, it is clear that there was an evolutionary change in the human *TSIX* region that resulted in loss of the human counterpart of the mouse testes-specific gene (*Tsx*) distal to murine *Tsix*. This breakage event shattered the *TSX* gene, spreading exons throughout the long arm of the X chromosome in the ancestor who gave rise to the human population. The extraordinary distance between *TSX* exons 4 and 5 has also been observed by Nesterova et al. (2001, fig. 5e). This breakpoint at ~27 kb from human *XIST* probably defines one boundary of the essential *XIC* in both species (indicated by an arrow in fig. 4).

An important question is whether this breakpoint influences the function of *TSIX* or the process of X inactivation. Some insights come from deletions in the murine *Tsix* gene, induced by Sado et al. (2001). Disrupting *Tsix* exon 1 (located ~28 kb from murine *Xist*)—but not the CpG island (~14 kb from *Xist*)—neither affected the second promoter nor interfered with *Tsix* function. However, the deletion in the *Tsix* exon 2 region, including part of the CpG-rich region, blocked *Tsix* tran-

scripts running across the *Xist* gene, which these investigators suggest is responsible for the loss of *Tsix* function. In addition, the targeted (3.5-kb) deletion by Lee (2000) of this CpG island also disrupted imprinted X inactivation. The evolutionary breakpoint in human *TSIX* not only eliminated exon 1 and most likely part of exon 2, but also resulted in the loss of most of the CpG-rich region near the start of *TSIX*, either directly or through evolutionary drift. The lack of significant homology over the *TSIX* region (figs. 1 and 4) and the large number of repetitive sequences—mostly long tandem repeats (LTRs) (figs. 1 and 3) not seen in the murine *Tsix* region (data not shown)—suggest that significant drifting has occurred. In any case, although the exon 1 deletion may not interfere with *TSIX* function, it seems likely that elimination of the CpG island would interfere with parental imprinting.

Significance of Species Differences with Respect to X Inactivation

There are many ways to compensate for sex differences in X gene dosage; compensatory mechanisms are directed at transcription and range from up-regulating the transcriptional output of the single X in males to down-regulating that of the X chromosomes in females. However, only mammals compensate by X inactivation. Since both marsupials and placental mammals use X inactivation, the basic mechanisms should be the same for all mammals. Therefore, features of X inactivation that differ among mammals are merely variations on the basic theme (Migeon 1990). These species variations tell us which details of the X inactivation scheme are not essential for establishing a single active X in mammalian cells. Such variable features include time of onset, choice of active X (either random or maternal), and stability of inactivation once established. Variations affecting stability result from differences in the DNA methylation status of CpG islands on the inactive X chromosome (Migeon 1990). Most likely, differences in the time of onset are responsible, at least in part, for the paternal (imprinted) inactivation in marsupials and in placental tissues of rodents and for the well-documented lack of imprinted X inactivation in human placental tissues (Migeon and Do 1978, 1979; Migeon et al. 1985, 1986; Mohandas et al. 1989; van den Hurk et al. 1997). Other variable features include the *XIST* locus itself, which has undergone inversion event(s) during mammalian evolution and whose DNA sequence is so poorly conserved among mammals that the marsupial gene has not yet been identified. Such species differences in the details of X inactivation reflect variations that do not interfere with the basic process responsible for the single active X.

What do our studies reveal about the role of *Tsix*/

TSIX in imprinted X inactivation? On one hand, they support the notion that murine *Tsix* plays a role in imprinted X inactivation in mice, perhaps as a factor to block *Xist* expression on the maternal X chromosome. Studies of knockouts of the mouse *Tsix* locus clearly show that it has a role in paternal X inactivation in the placenta; *Tsix* deletions carried on the maternal X lead to silencing of the maternal as well as the paternal X chromosome in many cells, resulting in placental abnormalities and fetal death (Lee 2000; Sado et al. 2001). On the other hand, along with the lack of paternal X inactivation in human placental tissues, the evolutionary changes we observe suggest that human *TSIX* does not share this function. The structural features of the murine *Tsix* locus seem to be suited to its imprinting function. Like many imprinted genes, *Tsix* is initiated near a CpG island that is differentially methylated; it produces an antisense transcript, which, whether processed or not, overlaps the promoter of its target gene and is monoallelically expressed from the opposite chromosome as *Xist*. The nature of the interaction between the antisense transcript and its target is not yet known, and the possibility that the abbreviated human *TSIX* transcript interacts with *XIST* at the critical moment in development cannot be excluded. However, the fact that the human gene is a truncated version of the murine one has implications for its function. Because the human transcript does not cover the *XIST* promoter, the means by which it interacts with *XIST* cannot involve direct physical contact with the promoter. In addition, since *TSIX* lacks the differentially methylated CpG island, which, when deleted in the mouse gene, disrupts imprinted X inactivation, it probably cannot imprint *XIST* expression. Further, we obtained no evidence that human *TSIX* inhibits *XIST* expression; the fact that *TSIX* and *XIST* are transcribed simultaneously from the same transgene in ES-10 and ES-5 cells suggests that it does not.

Jacob (1977) has suggested that Mother Nature is more tinkerer than engineer and that she uses materials at hand to modify her work. *Tsix* in mice—and, perhaps, a comparable RNA in marsupials—may have been used to reinforce the parental imprinting needed because of the earlier onset of X inactivation in these species. In any case, our observations could explain why X inactivation is not imprinted in human placental tissues.

The role of *Tsix* in the basic scheme of random (non-imprinted) X inactivation is less clear. *Tsix* knockouts do skew the patterns of X inactivation in the embryo proper, since the chromosome bearing the deletion is exclusively the inactive one. However, the effect of a knockout of *Tsix* function on random (nonimprinted) X inactivation must be distinguished from its effect on imprinted inactivation in the placenta. In addition, a nonspecific effect of the deletion on *Xist* function, unrelated to antisense function, cannot be excluded. In fact,

Morey et al. (2001) studied mice carrying a 65-kb deletion that removed all of *Tsix* except that which overlaps *Xist* and which was associated with exclusive inactivation of the deleted X. They showed that restoration of *Tsix* function represses *Xist* RNA accumulation in *cis* but does not restore random inactivation. These investigators suggest that although *Tsix* represses the initiation of X inactivation in *cis*, it is insufficient for normal random choice. Conceivably, human *TSIX* is an evolutionary relic of the murine gene, which continues to be expressed in fetal tissues, in much the same way that *XIST* continues to be expressed long after the critical time in development—because its expression does no harm.

In addition, it is unlikely that any *cis*-acting X-linked gene could, by itself, bring about the initial choice of the single active X. The existence of a *trans*-acting factor to block *XIST* expression was postulated by Lyon (1996) and others before the discovery of *Tsix*. Evidence from triploid cells (Jacobs et al. 1979; Migeon et al. 1979) suggests that the factor(s) needed to induce the silencing of *XIST* on a single X chromosome in each cell most likely operates in *trans* and is autosomal in origin. In the presence of a *trans*-acting factor to repress *XIST* expression on the active X and DNA methylation to lock in the silence of the locus, *Tsix* would not be essential to the inactivation process but would merely reinforce the repression of maternal *Xist* in the species in which it is needed.

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Electronic-Database Information

Accession numbers and URLs for data in this article are as follows:

Celera, <http://public.celera.com/index.cfm> (for publicly available human genome sequence)
 GenBank, <http://www.ncbi.nlm.nih.gov/Genbank/> (for human *XIC* reference sequence [accession number U80460] and mouse *XIC* reference sequences [accession numbers AJ010350, M97167, L04961, U41394, X999460])

GeneBuilder, <http://125.itba.mi.cnr.it/~webgene/genebuilder.html> (for gene-structure prediction)
 HCtata, http://125.itba.mi.cnr.it/~webgene/wwwHC_tata.html (for identification of promoter elements)
 Oak Ridge National Laboratory Gene Recognition and Assembly Internet Link, <http://compbio.ornl.gov/Grail-1.3/> (Grail, version 1.3, for CpG island identification)
 Online Mendelian Inheritance in Man (OMIM), <http://www.ncbi.nlm.nih.gov/Omim/> (for *XIST* [MIM 314670] and *Tsix* [MIM 300181])
 ORFGene 2, <http://125.itba.mi.cnr.it/~webgene/wwworfgene2.html> (for identification of ORFs)
 PipMaker, <http://bio.cse.psu.edu/pipmaker/> (for homology analysis)
 RepeatMasker, <http://ftp.genome.washington.edu/RM/RepeatMasker.html>

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