

Identification of Developmentally Specific Enhancers for *Tsix* in the Regulation of X Chromosome Inactivation

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X chromosome inactivation silences one of two X chromosomes in the mammalian female cell and is controlled by a binary switch that involves interactions between *Xist* and *Tsix*, a sense-antisense pair of noncoding genes. On the future active X chromosome, *Tsix* expression suppresses *Xist* upregulation, while on the future inactive X chromosome, *Tsix* repression is required for *Xist*-mediated chromosome silencing. Thus, understanding the binary switch mechanism depends on ascertaining how *Tsix* expression is regulated. Here we have taken an unbiased approach toward identifying *Tsix* regulatory elements within the X chromosome inactivation center. First, we defined the major *Tsix* promoter and found that it cannot fully recapitulate the developmental dynamics of *Tsix* expression, indicating a requirement for additional regulatory elements. We then delineated two enhancers, one classical enhancer mapping upstream of *Tsix* and a bipartite enhancer that flanks the major *Tsix* promoter. These experiments revealed the intergenic transcription element *Xite* as an enhancer of *Tsix* and the repeat element *DXPas34* as a component of the bipartite enhancer. Each enhancer contains DNase I-hypersensitive sites and appears to confer developmental specificity to *Tsix* expression. Characterization of these enhancers will facilitate the identification of *trans*-acting regulatory factors for X chromosome counting and choice.

In mammals, the expression of X-linked genes is equalized in males and females by the transcriptional inactivation of one X chromosome in the female (2, 25). During embryogenesis, random X chromosome inactivation (XCI) occurs as the epiblast lineage differentiates and is regulated by counting and choice mechanisms, such that one randomly selected X chromosome remains active in each cell (33). Appropriate developmental regulation of XCI requires the X chromosome inactivation center (*Xic*), a *cis*-acting locus on the X chromosome that functions as a binary switch between the on and off transcriptional states (34). Molecular and genetic analyses of the *Xic* have focused on two genes, *Xist* and *Tsix*, that are integral to its function. *Xist* produces a large noncoding RNA that triggers X chromosome inactivation and coats the inactive X chromosome (6, 7, 27, 31), while *Tsix*, the antisense partner of *Xist*, antagonizes *Xist* expression and negatively regulates X chromosome inactivation (11, 21, 22, 35). Thus, *Tsix* works genetically upstream of *Xist*, and the opposing interplay between these two genes determines the pattern of XCI in the female cell.

Many aspects of *Tsix-Xist* dynamics remain poorly understood. Because the suppression and induction of *Xist* are regulated by *Tsix*, understanding how XCI is regulated in turn requires an understanding of how *Tsix* expression is controlled. During cell differentiation, *Tsix* expression is silenced asynchronously on the active and inactive X chromosomes (X_a and X_i , respectively) (21, 22). On the X_i chromosome, the loss of *Tsix* expression permits the upregulation of *Xist* and chromo-

some inactivation in *cis*. On the X_a chromosome, the persistence of *Tsix* expression blocks *Xist* upregulation and preserves the activity of the chromosome, with *Tsix* expression continuing until just after *Xist* expression is silenced. Loss-of-function mutations that reduce, abolish, or truncate *Tsix* transcription cause *Xist* upregulation and XCI on the linked X chromosome (9, 22, 24, 28, 35, 37). Conversely, gain-of-function alterations that augment the strength and duration of *Tsix* expression from one allele prevent *Xist* upregulation and XCI in *cis* (24, 38). Thus, the appropriate temporal and allelic regulation of *Tsix* expression is necessary to specify the transcriptional fate of each X chromosome by suppressing *Xist* activity on one X chromosome and permitting its upregulation on the other. In this manner, the asymmetric profile of *Tsix* expression underlies the epigenetic choice of which X chromosome is inactivated.

How is *Tsix* expression controlled developmentally and allelically? Several elements were recently implicated. One study suggested that a regulatable chromatin insulator, CTCF, acts as a transcriptional switch for *Xist* and *Tsix* (8). CTCF binding occurs around *DXPas34* (10), a repeat sequence downstream of the *Tsix* transcription start site at a location between the promoters of *Tsix* and *Xist*. This genetic arrangement is reminiscent of that observed at the autosomally imprinted locus *H19/Igf2*, where CTCF insulators are thought to regulate the accessibility of *H19* and *Igf2* on reciprocal chromosomes to shared enhancers (4, 14, 16, 17). In the *Xic*, however, such enhancers have not yet been reported. Relevant to this information, another recent study identified *Xite*, an intergenic region that is located upstream of *Tsix* and that harbors strong DNase I-hypersensitive sites (DHS) and several intergenic transcripts (30). Because deleting *Xite* results in the downregulation of *Tsix* at the onset of XCI and a skewed pattern of XCI,

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Xite is a candidate regulator of *Tsix* expression. Its mechanism of action is presently unknown. Finally, knockout studies of a 65-kb region (9, 29) that includes *Tsix* and *Xite* suggested the presence of X chromosome counting elements that are distinct from the 5' ends of *Tsix* and *Xite*. Such elements are also candidate regulators of *Tsix* expression. Thus, there is considerable physical overlap among the various candidate regulatory elements of *Tsix*. However, although these processes are likely to be coupled, the molecular links have remained unclear.

Here we have taken a systematic and unbiased approach toward identifying regulatory sequences in the region downstream of *Xist*. We found that the major *Tsix* promoter is insufficient to confer appropriate developmental control, implicating additional elements in transcriptional regulation. Several enhancer-like sequences were identified within the *Tsix* and *Xite* gene bodies. As discussed below, the data suggest that the combination of these elements confers transcriptional regulation to *Tsix*. Because they reside in domains implicated in counting and choice, we speculate that counting and choice elements will be identified as transcriptional regulators of *Tsix*.

MATERIALS AND METHODS

RNase protection. Total RNA was isolated with Trizol (Invitrogen). RNase protection was performed essentially as described previously (1). For *Tsix* riboprobes, pNS2 and pNS3 were cut with BamHI and transcribed with SP6 polymerase and [α - 32 P]CTP. An actin template (Clontech) was transcribed with T3 polymerase. For hybridizations, 2.5×10^5 cpm of probe and 20 μ g of RNA were used. A radiolabeled DNA sequencing reaction served as a size standard.

Cell culture and transfection. Male J1, female 16.7, and transgenic 116.6 embryonic stem (ES) cells and their culture conditions have been described elsewhere (21–23). For transfection of undifferentiated ES cells, cells were passaged ~1:2.5 on gelatin-treated 24-well plates and cultured overnight. On the next day, cells were transfected with Lipofectamine 2000 (Invitrogen) according to the manufacturer's protocol. Briefly, transfection was performed with serum-containing antibiotic-free medium, 3 μ l of Lipofectamine 2000, and 1 μ g of DNA (100 ng of pCS2+c β gal [see below] and 900 ng of luciferase vector) per well. Transfection for 12 to 18 h yielded efficiencies of ~50 to 90% for J1 cells and ~10 to 20% for 16.7 cells.

To induce the differentiation of 16.7 ES cells, cells were aggregated into embryoid bodies, grown in the absence of leukemia inhibitory factor (LIF) for 4 days in suspension cultures, and grown for an additional 5 to 11 days on gelatin-treated plates. On the day prior to transfection, cells were trypsinized, used to seed each well of a 24-well plate at 10^5 cells, and cultured overnight. Transfection was performed with serum-containing antibiotic- and LIF-free medium, 2 μ l of Lipofectamine 2000, and 1 μ g of DNA (100 ng of pCS2+c β gal and 900 ng of luciferase vector) per well.

NIH 3T3 cells and immortalized female fibroblasts were grown in Dulbecco minimal essential medium containing 10% fetal bovine serum. For transfection of NIH 3T3 cells, 10^5 cells were used to seed each well of a 24-well plate. On the next day, cells were transfected in a manner similar to that used for ES cells with 3 μ l of Lipofectamine 2000 and 2 μ g of DNA (200 ng of pCS2+c β gal and 1.8 μ g of luciferase vector) per well. For immortalized female fibroblasts, 0.5×10^5 cells were used to seed each well of a 24-well plate and then were transfected with 3 μ l of Lipofectamine 2000 and 1 μ g of DNA (100 ng of pCS2+c β gal and 900 ng of luciferase vector) per well.

In each experiment, samples were transfected in duplicate wells.

Luciferase and β -galactosidase assays. Media were aspirated, and cells were lysed by the addition of 100 μ l of Glo lysis buffer (Promega) and incubation at room temperature for 5 to 15 min. For the luciferase assay, 80 μ l of lysate was mixed with 80 μ l of Steady-Glo reagent (Promega) in a 96-well black OptiPlate-96F (Packard). The plate was incubated at room temperature for 5 min and covered with TopSeal-A sealing film (Packard). Luminescence was measured with a Packard TopCount-NXT instrument (3-min precount delay, 2-s count time).

For the β -galactosidase assay, 20 μ l of lysate was diluted with 15 μ l of Glo lysis buffer in a flat-bottom 96-well clear polystyrene plate. A total of 35 μ l of ice-cold $2 \times \beta$ -galactosidase assay buffer (200 mM sodium phosphate [pH 7.3], 2 mM MgCl₂, 100 mM β -mercaptoethanol, 1.33 mg of *o*-nitrophenyl- β -D-galactopyr-

anoside/ml) was added, and the plate was incubated at 37°C in a humidified chamber. The reaction was monitored visually and terminated with 130 μ l of 1 M Tris base. The absorbance was measured at 405 nm with a multiwell reader (Coulter) and corrected against the value for a water blank.

To control for transfection efficiency in each well, luciferase activity was normalized to β -galactosidase activity. Values for duplicate wells were averaged, and the mean was subsequently normalized to the value for a reference promoter to enable comparison across multiple experiments. For each plasmid, luciferase activity was measured in two to six independent experiments.

Plasmids. pCS2+c β gal was described previously (39). Plasmids generated for this study are described in Table 1. Except where specifically noted otherwise, luciferase activity was assayed with pNS11, a *kan/neo*-marked promoterless luciferase vector derived from pGL3-Basic (Promega) as follows. A PvuII-ClaI fragment containing the herpes simplex virus thymidine kinase (TK) promoter (–196 to +18 bp [–196/+18] fragment; C to T at +13) was ligated in a three-piece reaction with 260-bp ClaI-PvuII and 3.5-kb PvuII fragments from pEGFP-1 (Clontech). In the resultant vector, pNS9, the TK promoter replaces the simian virus 40 origin-promoter-enhancer of pEGFP-1. pNS9 was subsequently cut with AflII, blunted, ligated to a BamHI linker, cut with BglII, blunted, and ligated to a NotI linker; the 3-kb NotI-BamHI fragment was liberated and ligated to a 2.2-kb NotI-BamHI fragment from pGL3-Basic, yielding pNS11. In pNS185, the BamHI site of pNS11 has been replaced with a BssHII site.

DHS mapping. DHS mapping was performed essentially as described previously (13). Briefly, 10^8 trypsinized cells were washed with phosphate-buffered saline, pelleted, and resuspended in 10 ml of ice-cold nuclear isolation buffer (NIB) (0.32 M sucrose, 3 mM CaCl₂, 2 mM magnesium acetate, 10 mM Tris-HCl [pH 7.5], 0.1 mM EDTA, 1 mM dithiothreitol). A 1/20 volume of NIB–0.3% NP-40 was added, and the suspension was subjected to Dounce homogenization 15 times with pestle B. Nuclei were centrifuged at $400 \times g$ and resuspended in ice-cold NIB at 0.2 mg/ml, as determined by total nucleic acid content. For ES cells, DNase I (Worthington) was added to aliquots at final concentrations of 0 to 10 μ g/ml, and samples were incubated at 37°C for 2 min. For fibroblasts, the final concentrations of DNase I were 0 to 80 μ g/ml, and samples were incubated for 5 min. Digestions were terminated by the addition of an equal volume of $2 \times$ stop solution (1% sodium dodecyl sulfate, 0.6 M NaCl, 20 mM Tris-HCl [pH 7.5], 10 mM EDTA, 0.2 mg of proteinase K/ml), and samples were incubated overnight at 37°C. DNA was precipitated with 1.25 M ammonium acetate, resuspended, and digested for Southern analysis.

For Southern blotting, HindIII- and BamHI-digested DNAs were probed with a 0.5-kb BamHI-AvrII fragment (bases 141031 to 141540 of the sequence assigned GenBank accession no. AJ421479) and a 0.5-kb BamHI-MscI fragment (bases 146400 to 146909 of the sequence assigned GenBank accession no. AJ421479), respectively.

RESULTS

The major *Tsix* promoter cannot recapitulate developmental regulation. Two CpG-rich transcription initiation sites of *Tsix* have been described previously, with major and minor promoters located, respectively, 12 and 28 kb downstream of *Xist* (Fig. 1A) (21, 35, 36). Because several different 5' termini have been described at the major initiation site (Fig. 1B), we performed RNase protection to determine the precise location of the major *Tsix* promoter. With a 220-nucleotide (nt) *Tsix* probe spanning the 5' termini, two clusters of protected bands specific to ES cells were observed at nt 140 and 110, corresponding, respectively, to initiation sites 45 and 75 bp downstream of a BamHI site (Fig. 1B and C). A less abundant, fully protected band of 185 nt was also observed, likely corresponding to transcription originating from the minor *Tsix* promoter and other upstream positions (Fig. 1C) (11, 30, 35, 36). Similar results were obtained with a second, overlapping *Tsix* probe (data not shown). These data define within the major promoter two transcription initiation sites that coincide with previously described 5' termini located 46 bp (36) and 75 bp (35) downstream of the BamHI site (Fig. 1B). We designate these termini bp +1 and bp +29, respectively.

To functionally define the major promoter, we cloned a

TABLE 1. Plasmids generated in this study^a

Figure	Plasmid	Insert (5' terminus-3' terminus)	Backbone	Description
1C	pNS2	<i>Tsix</i> -44/+237 (BamHI-MscI)	pGEM7	Riboprobe template
1C	pNS3	<i>Tsix</i> -44/+144 (BamHI-EcoRV)	pGEM7	Riboprobe template
1D	pNS11			<i>kan/neo</i> promoterless luciferase vector
1D	pNS58	Herpes simplex virus TK promoter -198/+18; C to T at +13	pNS11	Herpes simplex virus TK promoter
1D	pNS65	<i>Tsix</i> -557/+176 (KpnI-SmaI)	pNS11	733-bp major promoter
1D	pNS94	<i>Tsix</i> -17271/-16131 (NheI-NheI)	pNS11	1.1-kb minor promoter
2A	pNS78	<i>Tsix</i> -457/+176 (AhdI-SmaI)	pNS11	5' Truncation of major promoter
2A	pNS79	<i>Tsix</i> -279/+176 (BlnI-SmaI)	pNS11	5' Truncation of major promoter
2A	pNS80	<i>Tsix</i> -160/+176 (BanII-SmaI)	pNS11	5' Truncation of major promoter
2A	pNS81	<i>Tsix</i> -82/+176 (FspI-SmaI)	pNS11	5' Truncation of major promoter
2A	pNS82	<i>Tsix</i> -44/+176 (BamHI-SmaI)	pNS11	5' Truncation of major promoter
2A	pNS83	<i>Tsix</i> -24/+176 (Bsu36I-SmaI)	pNS11	5' Truncation of major promoter
2A	pNS84	<i>Tsix</i> +31/+176 (BsgI-SmaI)	pNS11	5' Truncation of major promoter
2A	pNS85	<i>Tsix</i> +67/+176 (BsiHKA1-SmaI)	pNS11	5' Truncation of major promoter
2A	pNS86	<i>Tsix</i> +95/+176 (BspMI-SmaI)	pNS11	5' Truncation of major promoter
2A	pNS87	<i>Tsix</i> +114/+176 (NarI-SmaI)	pNS11	5' Truncation of major promoter
2B	pNS66	<i>Tsix</i> -557/+144 (KpnI-EcoRV)	pNS11	3' Truncation of major promoter
2B	pNS67	<i>Tsix</i> -557/+116 (KpnI-NarI)	pNS11	3' Truncation of major promoter
2B	pNS69	<i>Tsix</i> -557/+63 (KpnI-BsiHKA1)	pNS11	3' Truncation of major promoter
2B	pNS70	<i>Tsix</i> -557/+29 (KpnI-BsgI)	pNS11	3' Truncation of major promoter
2B	pNS71	<i>Tsix</i> -557/-21 (KpnI-Bsu36I)	pNS11	3' Truncation of major promoter
2B	pNS72	<i>Tsix</i> -557/-40 (KpnI-BamHI)	pNS11	3' Truncation of major promoter
2B	pNS73	<i>Tsix</i> -557/-82 (KpnI-FspI)	pNS11	3' Truncation of major promoter
2B	pNS74	<i>Tsix</i> -557/-164 (KpnI-BanII)	pNS11	3' Truncation of major promoter
2B	pNS75	<i>Tsix</i> -557/-276 (KpnI-BlnI)	pNS11	3' Truncation of major promoter
2C	pNS215	<i>Tsix</i> -160/+116 (BanII-NarI)	pNS11	5' and 3' Truncations of major promoter
3	pNS64	<i>Tsix</i> -1420/+116 (SacI-NarI)	pNS11	-1.4 kb to promoter
3	pNS63	<i>Tsix</i> -2407/+116 (PmlI-NarI)	pNS11	-2.4 kb to promoter
3	pNS62	<i>Tsix</i> -3498/+116 (NheI-NarI)	pNS11	-3.5 kb to promoter
3	pNS61	<i>Tsix</i> -4880/+116 (NciI-NarI)	pNS11	-4.9 kb to promoter
3	pNS60	<i>Tsix</i> -5920/+116 (BamHI-NarI)	pNS11	-5.9 kb to promoter
3	pNS96	<i>Tsix</i> -9009/+116 (StuI-NarI)	pNS11	-9.0 kb to promoter
3	pNS228	λ 17053/18556 (KpnI-KpnI)	pNS65	λ 1.5 kb + <i>Tsix</i> promoter
3	pNS229	λ 39355/41275 (EcoRV-EcoRV)	pNS65	λ 1.9 kb + <i>Tsix</i> promoter
3	pNS231	λ 18560/21271 (KpnI-EcoRV)	pNS65	λ 2.7 kb + <i>Tsix</i> promoter
3	pNS232	λ 45829/652 (EcoRV-EcoRV)	pNS65	λ 3.3 kb + <i>Tsix</i> promoter
3	pNS234	λ 2087/6683 (EcoRV-EcoRV)	pNS65	λ 4.6 kb + <i>Tsix</i> promoter
3	pNS235	λ 33589/28214 (EcoRV-EcoRV)	pNS65	λ 5.4 kb + <i>Tsix</i> promoter
3	pNS236	λ 34503/27973 (BamHI-BamHI)	pNS65	λ 6.5 kb + <i>Tsix</i> promoter
3	pNS237	λ 17058/5505 (KpnI-BamHI)	pNS65	λ 11.5 kb + <i>Tsix</i> promoter
3	pNS113	<i>Tsix</i> -12045*/-6535 (ND-AvrII)	pNS65	(-12 kb to -6.5 kb) + <i>Tsix</i> promoter
3	pNS116	<i>Tsix</i> -15105/-10229 (XhoI-XhoI)	pNS65	(-15.1 kb to -10.2 kb) + <i>Tsix</i> promoter
3	pNS114	<i>Tsix</i> -18196/-13656 (KpnI-KpnI)	pNS65	(-18.2 kb to -13.7 kb) + <i>Tsix</i> promoter
3	pNS117	<i>Tsix</i> -23351/-17267 (ClaI-NheI)	pNS65	(-23.4 kb to -17.3 kb) + <i>Tsix</i> promoter
3	pNS115	<i>Tsix</i> -27322/-21489 (XbaI-XbaI)	pNS65	(-27.3 kb to -21.5 kb) + <i>Tsix</i> promoter
3	pNS123	<i>Tsix</i> -6535/-12045* (AvrII-ND)	pNS65	(-6.5 kb to -12 kb) + <i>Tsix</i> promoter
3	pNS109	<i>Tsix</i> -13656/-18196 (KpnI-KpnI)	pNS65	(-13.7 kb to -18.2 kb) + <i>Tsix</i> promoter
4A	pNS134	<i>Tsix</i> -11329/-6535 (ScaI-AvrII)	pNS65	5' Deletion of proximal enhancer + <i>Tsix</i> promoter
4A	pNS133	<i>Tsix</i> -10234/-6535 (XhoI-AvrII)	pNS65	5' Deletion of proximal enhancer + <i>Tsix</i> promoter
4A	pNS132	<i>Tsix</i> -9009/-6535 (StuI-AvrII)	pNS65	5' Deletion of proximal enhancer + <i>Tsix</i> promoter
4A	pNS129	<i>Tsix</i> -12045*/-9010 (ND-StuI)	pNS65	3' Deletion of proximal enhancer + <i>Tsix</i> promoter
4A	pNS130	<i>Tsix</i> -12045*/-10229 (ND-XhoI)	pNS65	3' Deletion of proximal enhancer + <i>Tsix</i> promoter
4A	pNS131	<i>Tsix</i> -12045*/-11330 (ND-ScaI)	pNS65	3' Deletion of proximal enhancer + <i>Tsix</i> promoter
4A	pNS135	<i>Tsix</i> -10234/-9010 (XhoI-StuI)	pNS65	1.2-kb enhancer + <i>Tsix</i> promoter
4A	pNS142	<i>Tsix</i> -9010/-10234 (StuI-XhoI)	pNS65	1.2-kb enhancer (reverse) + <i>Tsix</i> promoter
4B	pNS248	<i>Tsix</i> -10234/+116 (XhoI-NarI)	pNS11	-10.2 kb to promoter
4C	pNS155	<i>Tsix</i> -10234/-9010 (XhoI-StuI)	pNS11	1.2-kb enhancer
4C	pNS156	<i>Tsix</i> -9010/-10234 (StuI-XhoI)	pNS11	1.2-kb enhancer (reverse)

Continued on following page

TABLE 1—Continued

Figure	Plasmid	Insert (5' terminus-3' terminus)	Backbone	Description
4F	pNS258	<i>Xist</i> P1 promoter (−540/+9)	pNS11	<i>Xist</i> P1 promoter
4F	pNS262	<i>Tsix</i> −10234/−9010 (XhoI-StuI)	pNS258	1.2-kb enhancer + <i>Xist</i> P1 promoter
4F	pNS263	<i>Tsix</i> −9010/−10234 (StuI-XhoI)	pNS258	1.2-kb enhancer (reverse) + <i>Xist</i> P1 promoter
4F	pNS259	<i>Xist</i> P2 promoter (−601/+21)	pNS11	<i>Xist</i> P2 promoter
4F	pNS264	<i>Tsix</i> −10234/−9010 (XhoI-StuI)	pNS259	1.2-kb enhancer + <i>Xist</i> P2 promoter
4F	pNS265	<i>Tsix</i> −9010/−10234 (StuI-XhoI)	pNS259	1.2-kb enhancer (reverse) + <i>Xist</i> P2 promoter
5A	pNS106	<i>Tsix</i> −4880/+116 (NciI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +145/+724 (EcoRV-AgeI) placed 3' to luciferase gene	pNS11	Combined <i>Tsix</i> upstream and downstream sequences
5A	pNS126	<i>Tsix</i> −4880/+116 (NciI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +145/+1867 (EcoRV-SalI) placed 3' to luciferase gene	pNS11	Combined <i>Tsix</i> upstream and downstream sequences
5A	pNS127	<i>Tsix</i> −4880/+116 (NciI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +145/+4049 (EcoRV-HindIII) placed 3' to luciferase gene	pNS11	Combined <i>Tsix</i> upstream and downstream sequences
5A	pNS107	<i>Tsix</i> −9009/+116 (StuI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +145/+724 (EcoRV-AgeI) placed 3' to luciferase gene	pNS11	Combined <i>Tsix</i> upstream and downstream sequences
5A	pNS108	<i>Tsix</i> −9009/+116 (StuI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +145/+1867 (EcoRV-SalI) placed 3' to luciferase gene	pNS11	Combined <i>Tsix</i> upstream and downstream sequences
5A	pNS128	<i>Tsix</i> −9009/+116 (StuI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +145/+4049 (EcoRV-HindIII) placed 3' to luciferase gene	pNS11	Combined <i>Tsix</i> upstream and downstream sequences
5B	pNS255	<i>Tsix</i> −9009/+116 (StuI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +721/+4049 (AgeI-HindIII) placed 3' to luciferase gene	pNS11	Combined <i>Tsix</i> upstream and downstream sequences
5B	pNS256	<i>Tsix</i> −9009/+116 (StuI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +1863/+4049 (SalI-HindIII) placed 3' to luciferase gene	pNS11	Combined <i>Tsix</i> upstream and downstream sequences
5C	pNS254	λ 27479/36895 and <i>Tsix</i> −557/+116 (KpnI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +145/+4049 (EcoRV-HindIII) placed 3' to luciferase gene	pNS11	λ substitution, <i>Tsix</i> promoter, and downstream sequences
5C	pNS251	<i>Tsix</i> −9009/+116 (StuI-NarI) placed 5' to luciferase gene, λ 18555/22350 placed 3' to luciferase gene	pNS11	<i>Tsix</i> upstream sequences and λ substitution
5D	pNS190	<i>Tsix</i> +145/+4049 (EcoRV-HindIII) and <i>Tsix</i> −557/+144 (KpnI-EcoRV) placed 5' to luciferase gene, <i>Tsix</i> −9009/−557 (StuI-KpnI) placed 3' to luciferase gene	pNS185	Swap of <i>Tsix</i> upstream and downstream sequences
5D	pNS195	<i>Tsix</i> +145/+4049 (EcoRV-HindIII) and <i>Tsix</i> −557/+144 (KpnI-EcoRV) placed 5' to luciferase gene, <i>Tsix</i> −4880/−557 (NciI-KpnI) placed 3' to luciferase gene	pNS185	Swap of <i>Tsix</i> upstream and downstream sequences
5F	pNS249	<i>Tsix</i> −10234/+116 (XhoI-NarI) placed 5' to luciferase gene, <i>Tsix</i> +145/+4049 (EcoRV-HindIII) placed 3' to luciferase gene	pNS11	Combined <i>Tsix</i> upstream and downstream sequences

^a Plasmids are listed in the order in which they first appear in the figures, from left to right and top to bottom. The orientation of λ inserts is not known. An asterisk indicates that the terminus is approximate. For *Tsix* sequences, +1 corresponds to base 77675 of the sequence assigned GenBank accession no. X99946. Construction details and plasmid sequences are available on request. ND, end not determined.

−557/+176 fragment upstream of a promoterless luciferase reporter. In transiently transfected ES cells, this fragment provided luciferase expression comparable to that of a control herpes simplex virus TK promoter (Fig. 1D). The minor upstream *Tsix* promoter was found to be five- to sixfold less active when assayed in parallel (Fig. 1D), consistent with the lower abundance of upstream *Tsix* transcripts (Fig. 1C) (11, 30, 35, 36). Because the upstream promoter produces a small fraction of *Tsix* transcripts and its removal causes no obvious consequences for XCI (30, 35), our subsequent experiments focused exclusively on the major promoter.

To define the sequences that confer full promoter activity, we next generated 5' and 3' deletion derivatives of the −557/+176 region and compared their activities to that of the parental fragment. As measured in ES cells, 5' truncations as far as bp −160 did not decrease transcriptional activity (Fig. 2A). Further truncations sharply reduced activity, with discrete reductions suggesting the presence of positive elements between bp −160 and −82 and between bp −44 and −24 that are distinct from the two initiation sites (Fig. 2A and E). For deletions at the 3' terminus, fragments truncated at bp +144 or +116 showed full or slightly reduced promoter activity, respec-

tively (Fig. 2B). More severe 3' truncations elicited discrete reductions in activity, consistent with the loss of a positive regulatory element between bp +63 and +116 (Fig. 2E) or with alterations in transcript stability or translation efficiency. As predicted by this deletion analysis, a −160/+116 fragment that combined both 5' and 3' deletions was sufficient for full promoter activity (Fig. 2C).

Tsix is expressed in undifferentiated embryonic cells and is silenced during differentiation. Because this property is a key criterion for judging whether isolated regulatory elements faithfully recapitulate *Tsix* expression, we compared the activities of the promoter in undifferentiated ES cells, which express *Tsix*, and in immortalized female mouse fibroblasts and male NIH 3T3 fibroblasts, which do not (Fig. 1C and data not shown) (21). Additionally, we assayed the promoter in female ES cells that had been allowed to differentiate for 11 to 17 days in the absence of LIF, a time at which *Tsix* expression has been silenced in most cells (38). In each of these assays, the *Tsix* promoter was active (Fig. 2D). Thus, the *Tsix* promoter is constitutively active and does not recapitulate the developmental dynamics of *Tsix* expression. We concluded that additional

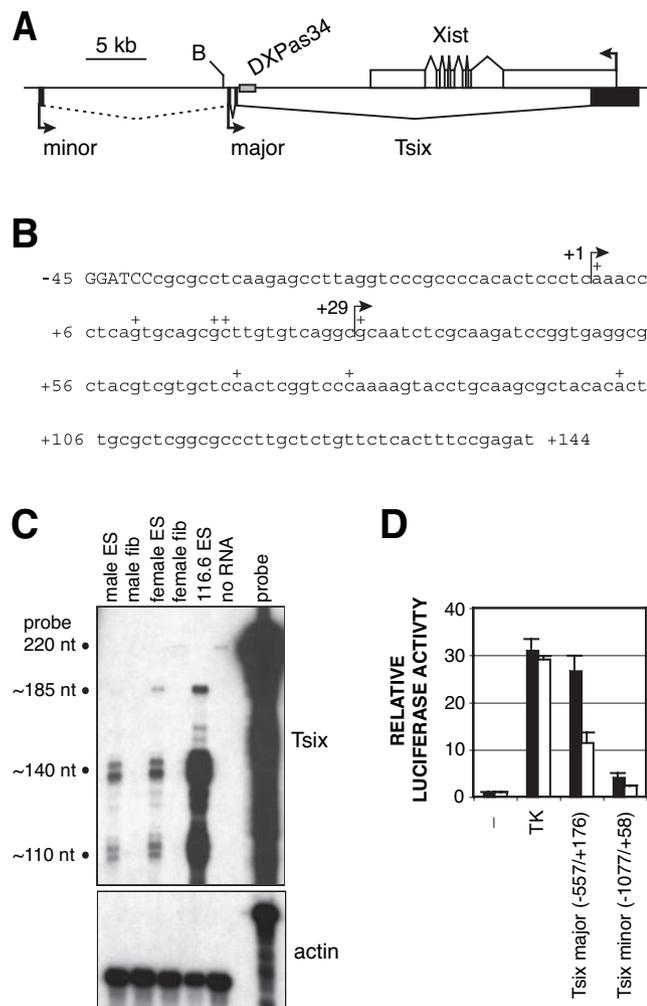


FIG. 1. Functional delineation of the major *Tsix* promoter. (A) Map of the *Xist/Tsix* locus. White boxes, *Xist* exons; black boxes, *Tsix* exons; broken line, less abundant *Tsix* transcripts; grey rectangle, *DXPas34* repeat array; B, BamHI. (B) Sequence proximal to the major initiation site. The 5' terminus is the BamHI site in panel A. Previously described 5' RACE termini are indicated by plus signs (21, 35, 36; D. Warshawsky, unpublished data). Arrows denote start sites defined by RNase protection analysis (see panel C). (C) RNase protection analysis at the major *Tsix* initiation site. Results obtained with *Tsix* (top panel) and actin (bottom panel) probes are shown. The *Tsix* riboprobe contains a 31-nt 5' leader sequence fused to a 189-nt *Tsix* sequence antisense to that shown in panel B. 116.6 ES cells bear ~25 copies of the *Xic* (23). (D) Activities of the major and minor *Tsix* promoters. Mean luciferase activities are shown for transiently transfected male ES cells (black bars) and female ES cells (white bars). Error bars denote one standard error. The activity of the backbone vector was set to 1. The major promoter is numbered as in panel B; the minor promoter is numbered with respect to the minor start site (35).

cis regulatory elements must be present in *Xic* and endeavored to identify them.

Identification of a *Tsix* enhancer that maps within *Xite*. To determine whether transcriptional control elements reside upstream of the major *Tsix* promoter, we performed two experiments. First, we examined whether contiguous 5'-flanking sequences extending up to kb -9 could modify promoter activity (Fig. 3, top panel). Because transfection efficiency is related

inversely to plasmid size (18), we also assayed control plasmids containing equivalent amounts of heterologous sequences from bacteriophage λ . When controlled for insert size, none of the *Tsix* fragments up to kb -9 significantly enhanced or repressed the *Tsix* promoter in transiently transfected ES cells. These results indicate that upstream *Tsix* sequences extending to -9 kb are by themselves unable to modify promoter activity.

Second, we examined whether sequences beyond kb -9 might contain *Tsix* regulatory elements. Five overlapping fragments extending from kb -6.5 to kb -27.3 were cloned upstream of the *Tsix* promoter-luciferase cassette in their natural orientation and assayed in ES cells. Two of these fragments, the first extending from kb -12 to kb -6.5 and the second extending from kb -18.2 to kb -13.7 , stimulated luciferase activity 4- to 12-fold relative to the results obtained with similarly sized control plasmids (Fig. 3, bottom panel). The stimulatory effects were orientation independent (Fig. 3), suggesting the presence of transcriptional enhancers in these regions.

To carry out finer mapping, we generated 5' and 3' deletion derivatives for both fragments and assayed their activities in ES cells. Deletion analysis of fragment from kb -18.2 to -13.7 ($-18.2/-13.7$ fragment) defined a 1.1-kb MluI-XhoI segment that was required for promoter stimulation (data not shown). This segment overlaps the minor upstream promoter defined previously (30, 35). However, because genetic analyses have shown no apparent function for this region in either random or imprinted X chromosome inactivation in vivo (30, 35), we chose not to pursue its analysis further.

Deletion analysis of the proximal element defined a 1.2-kb XhoI-StuI segment between kb -10.2 and -9.0 that was both necessary and sufficient for *Tsix* promoter stimulation, yielding a 2.5- to 3.5-fold stimulatory effect in male and female ES cells in an orientation-independent manner (Fig. 4A). In these experiments, the tested fragments were juxtaposed to the *Tsix* promoter, a synthetic configuration lacking intervening sequences. To test whether the 1.2-kb segment was able to stimulate the promoter in a more natural context, we assayed the entire upstream region from kb -10.2 to the promoter. In ES cells, this fragment stimulated the promoter approximately 3.5-fold relative to the results obtained with a similarly sized fragment extending to kb -9 (Fig. 4B). Thus, the 1.2-kb element was able to stimulate the *Tsix* promoter not only when placed immediately upstream but also from a significant distance in its natural context. Thus, we conclude that the $-10.2/-9.0$ region is able to function as a classical transcriptional enhancer of *Tsix*.

The position of the enhancer between kb -10.2 and -9.0 places it within the *Xite* locus (30). Indeed, *Xite* contains multiple DHS, a hallmark of enhancers, and has been shown to positively regulate *Tsix* expression. The 1.2-kb enhancer coincides with the most prominent DHS in the region and with initiation sites for the intergenic *Xite* transcript (30). In the course of the present analysis, we found that the 1.2-kb region contains bidirectional promoter activities. When joined to a promoterless luciferase reporter in its natural orientation, the 1.2-kb region provided luciferase expression at a level comparable to that provided by the major *Tsix* promoter (Fig. 4C). Promoter activity was slightly weaker in the reverse orientation (Fig. 4C). Given the bidirectional promoter activities, one potential explanation for the enhancing effect observed earlier is

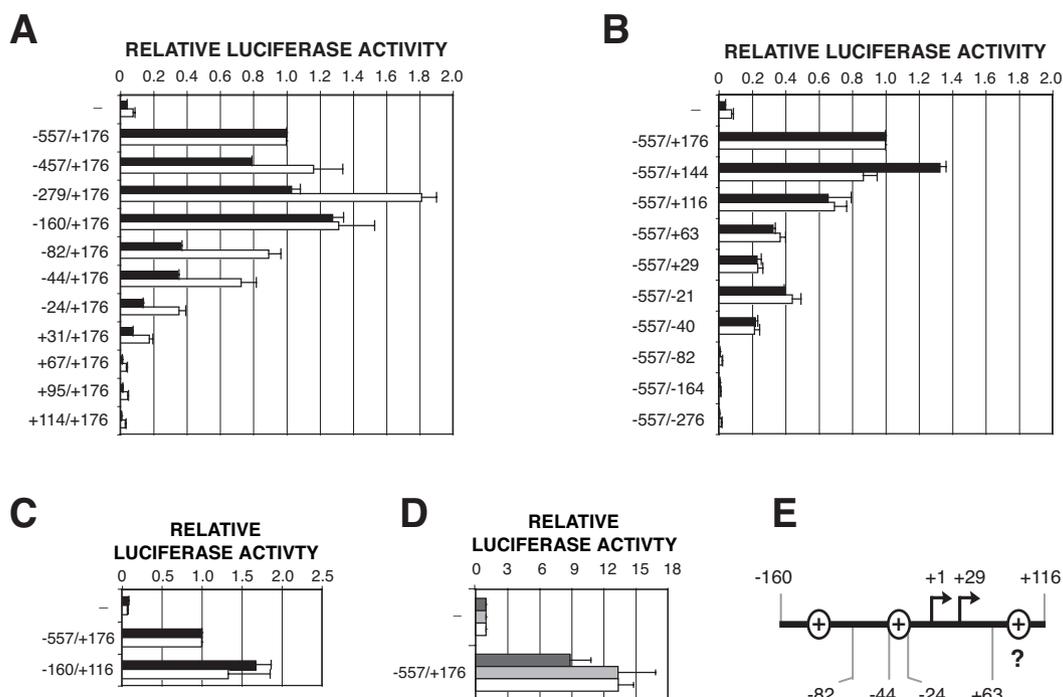


FIG. 2. The major *Tsix* promoter is constitutively active and is contained within a 276-bp region. (A to C) Activities of *Tsix* promoter deletion derivatives. Mean luciferase activities are shown for transiently transfected male ES cells (black bars) and female ES cells (white bars). Error bars denote one standard error. The activity of the +557/+176 fragment was set to 1. (D) The major *Tsix* promoter is constitutively active. Mean luciferase activities are shown for transiently transfected immortalized female mouse fibroblasts (dark grey bars), male NIH 3T3 fibroblasts (light grey bars), and differentiated female ES cells (days 11 to 17) (white bars). Error bars denote one standard error. The activity of the backbone vector was set to 1. (E) Anatomy of the major *Tsix* promoter. Inferred locations of positive elements are indicated by ovals within denoted sequence intervals.

that the 1.2-kb region simply adds to luciferase expression through its promoter activity and a mechanism of transcriptional readthrough. However, we do not believe this to be the case, as the *Xite* promoter would provide a maximum twofold increase in expression, significantly lower than the observed level of stimulation. Furthermore, potential readthrough transcripts from the 1.2-kb region are unlikely to increase luciferase expression when initiated at a significant distance from the reporter (Fig. 4B). Therefore, we conclude that the 1.2-kb region contains an enhancer that is functionally distinct from any inherent promoter activity.

The 1.2-kb *Xite* enhancer exhibits proper developmental regulation and is specific for *Tsix*. To test whether the *Xite* enhancer stimulates *Tsix* expression in a developmentally specific manner, we compared its activities in undifferentiated ES cells, where *Tsix* is active, and in differentiated ES cells and fibroblasts, where *Tsix* expression has been reduced or extinguished. In contrast to its ability to stimulate the *Tsix* promoter in undifferentiated ES cells (Fig. 4A and B), the *Xite* enhancer was inactive in male NIH 3T3 fibroblasts, immortalized female fibroblasts, and female ES cells that had been allowed to differentiate for over 11 days (Fig. 4D and data not shown). This profile of enhancer activity correlates with the presence of a DHS within the 1.2-kb *Xite* enhancer specifically in undifferentiated cells (30). Interestingly, the bidirectional promoter activities of this element were also specific to undifferentiated ES cells (Fig. 4C and E), consistent with the loss of intergenic

Xite transcripts upon cell differentiation (30). Thus, both enhancer and promoter activities were specific to undifferentiated ES cells, suggesting that the 1.2-kb region mediates many of the properties of *Xite* and contributes to the developmental regulation of *Tsix* expression in vivo.

One proposal for the control of *Xist* and *Tsix* expression invokes an enhancer that is shared by both genes and that is regulated in an allele-specific manner by an intervening chromatin insulator (8). To determine whether the *Xite* enhancer might be shared by *Tsix* and *Xist*, we tested whether it could stimulate the *Xist* promoter in undifferentiated ES cells and found that it induced neither the *Xist* P₁ nor the *Xist* P₂ promoter (15) when placed upstream in either orientation (Fig. 4F). Because *Xist* expression increases significantly in differentiated female cells after the onset of XCI, we also tested somatic cells and female ES cells that had been allowed to differentiate for 11 to 17 days. However, neither immortalized female fibroblasts nor differentiated ES cells exhibited any stimulation of the P₁ and P₂ promoters when the *Xite* enhancer was placed upstream in either orientation (data not shown). Thus, the *Xite* enhancer acts specifically on *Tsix* and has no apparent effect on the *Xist* promoter in the context of these experiments.

Identification of a bipartite enhancer in *Tsix* with developmental specificity. To identify additional regulatory elements, we screened for stimulatory sequences downstream of the *Tsix* promoter. Some attention has been focused on the repeat

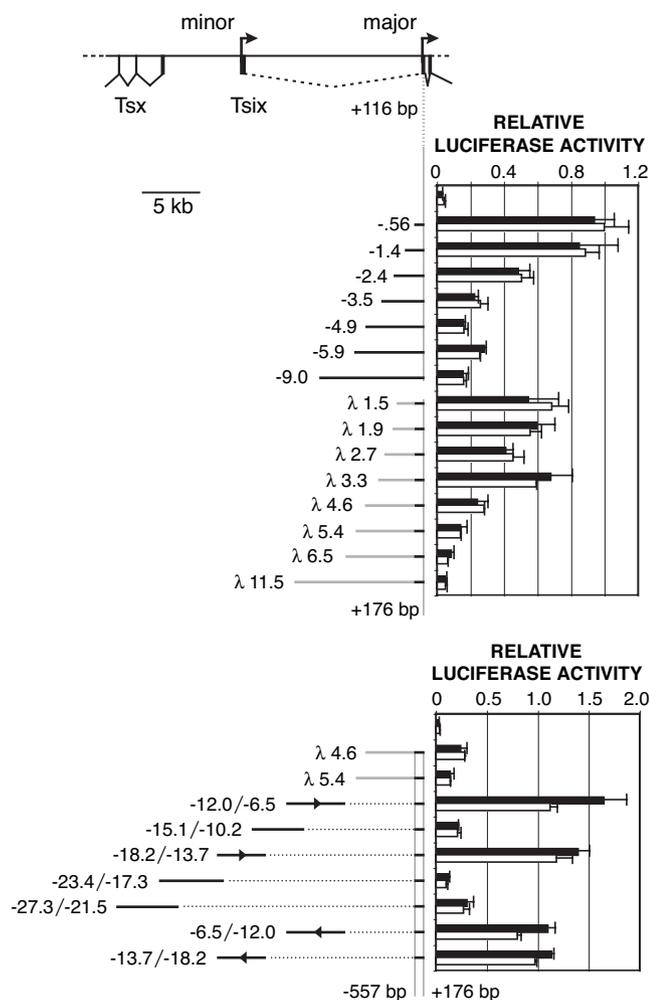


FIG. 3. Identification of enhancing activities in the *Xite* region. Mean luciferase activities are shown for transiently transfected male ES cells (black bars) and female ES cells (white bars). Error bars denote one standard error. The activity of the *Tsix* promoter ($-557/+176$; not shown) was set to 1. A map of the *Tsix* upstream region is shown at the top, with terminal *Tsx* exons at the left. (Upper panel) Assay of contiguous *Tsix* sequences extending from bp +116 to kb -9 ; λ sequences are coupled to the *Tsix* promoter ($-557/+176$). (Lower panel) Assay of distal sequences extending from kb -6.5 to -27.3 . The orientations of selected fragments are indicated.

array *DXPas34* (10) and a number of flanking CTCF-binding sites (8). To test the idea that *DXPas34* might be a *Tsix* control element (8, 11), we examined whether it could stimulate the *Tsix* promoter in various contexts. When placed upstream of the *Tsix* promoter, various fragments containing *DXPas34* alone were unable to confer stimulation in either orientation (data not shown). When placed 3' of a luciferase reporter driven by the *Tsix* promoter, *DXPas34* conferred a low level of stimulation, but only in some vector backbones (data not shown). These findings indicated that *DXPas34* by itself has little to no stimulatory capability.

We next considered whether *DXPas34* might regulate *Tsix* expression in a context-specific fashion that requires additional sequences. To test this idea, we assayed various combinations of upstream and downstream sequences (relative to the *Tsix*

start site) for their ability to stimulate the promoter. As described earlier, upstream sequences extending to kb -4.9 or -9 were by themselves unable to augment promoter activity significantly (Fig. 3, top panel). However, combinations of these upstream fragments with downstream segments terminating at kb $+1.9$ or $+4$ elicited potent luciferase stimulation, ranging from three- to ninefold, relative to that obtained with the upstream fragments in isolation (Fig. 5A). When each upstream fragment was combined with a downstream sequence extending to kb $+0.7$, stimulation was not observed, indicating that *DXPas34* was required for this effect. Inclusion of additional sequences from kb $+1.9$ to $+4.0$ further increased stimulation, especially in the context of upstream sequences extending to kb -9 . Although the 6.8-kb region extending from kb -4.9 to the *DXPas34* 3' terminus at kb $+1.9$ appeared to provide the bulk of this stimulation, the apparent contributions of distal upstream and downstream sequences suggested that as many as four distinct elements within a 13-kb region might mediate this effect. Thus, despite the inability of these elements to stimulate the promoter individually, combinations of these elements could stimulate *Tsix* expression in a highly cooperative manner.

In all contexts, *DXPas34* is essential for this cooperative stimulation. While the $+0.1/+0.7$ region (including *Tsix* exons 2 and 3) was dispensable, extending the deletion further to include *DXPas34* caused a loss of stimulation. Moreover, although sequences between kb $+1.9$ and $+4$ provided additional stimulation (Fig. 5A), they were unable to do so in the absence of *DXPas34* (Fig. 5B). We also found that when upstream or downstream sequences were replaced with stuffer fragments from bacteriophage λ , stimulation was severely reduced (Fig. 5C). Thus, the regulatory region has a bipartite structure, with an upstream region that contains sequences extending to kb -9.0 and a downstream region that contains *DXPas34* and additional sequences extending to kb $+4.0$.

To probe the context dependence of these interactions further, we assayed the consequences of swapping upstream and downstream sequences with respect to the *Tsix* promoter (Fig. 5D). Significantly, such swapping did not affect cooperative stimulation of the *Tsix* promoter for most substitutions (Fig. 5D and data not shown). These results demonstrate that combinations of upstream and downstream sequences can interact in a modular fashion and, like classical enhancers, tolerate orientation and position changes. *DXPas34* appears to have a central role in mediating the cooperative interactions of upstream and downstream regions within the enhancer.

Because the downstream sequences were placed 3' of the reporter in these assays, one potential concern is that the stimulation attributed to a bipartite enhancer might instead result from posttranscriptional alterations, such as alternative splicing or polyadenylation. Several experiments argued against this possibility. The removal of *Tsix* exons and splice donors within the downstream region did not affect cooperative stimulation (Fig. 5B). Furthermore, Northern blotting and 3' rapid amplification of cDNA ends (RACE) indicated that, in all instances, the sizes and 3' termini of luciferase messages were indistinguishable from those produced by a control simian virus 40 promoter-enhancer (data not shown). Thus, we conclude that the bipartite enhancer stimulates *Tsix* expression

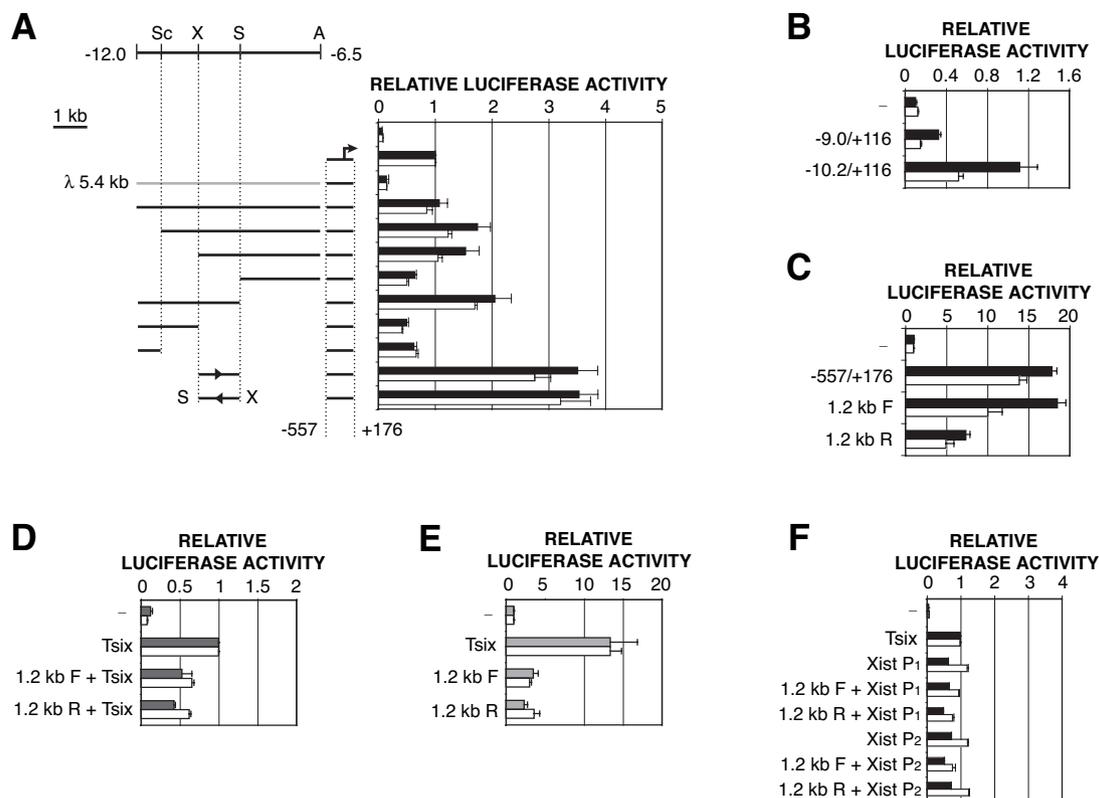


FIG. 4. *Xite* contains a 1.2-kb enhancer that is developmentally specific. (A) Deletion analysis of the upstream enhancer. Sc, ScaI; X, XhoI; S, StuI; A, AvrII. The orientations of selected fragments are indicated. Mean luciferase activities are shown for transiently transfected male ES cells (black bars) and female ES cells (white bars). Error bars denote one standard error. The activity of the *Tsix* promoter ($-557/+176$) was set to 1. (B) The 1.2-kb enhancer stimulates the *Tsix* promoter from its natural context. The activity of the *Tsix* promoter ($-557/+116$; not shown) was set to 1. (C) Bidirectional promoter activities of the 1.2-kb element. In this and subsequent panels, the forward (F) orientation of the 1.2-kb element is that in which the downstream (kb -9) terminus abuts the luciferase reporter; in the reverse (R) orientation, the upstream (kb -10.2) terminus abuts the reporter. The activity of the backbone vector was set to 1. (D) The 1.2-kb enhancer is unable to stimulate the *Tsix* promoter ($-557/+176$) in differentiated cell types. Mean luciferase activities are shown for immortalized female fibroblasts (grey bars) and differentiated female ES cells (days 11 to 17) (white bars). The activity of the *Tsix* promoter ($-557/+176$) was set to 1. (E) Bidirectional promoter activities are reduced in differentiated cell types. Mean luciferase activities are shown for transiently transfected NIH 3T3 fibroblasts (grey bars) and differentiated female ES cells (days 11 to 17) (white bars). The activity of the backbone vector was set to 1. (F) The enhancer is unable to stimulate the *Xist* promoter in undifferentiated ES cells. The 1.2-kb enhancer was assayed in the indicated orientation upstream of the *Xist* P₁ promoter ($-540/+9$) or the *Xist* P₂ promoter ($-601/+21$). Mean luciferase activities are shown for transiently transfected male ES cells (black bars) and female ES cells (white bars). The activity of the *Tsix* promoter ($-557/+176$) was set to 1.

at the transcriptional level rather than through alterations of splicing or polyadenylation.

To determine whether the stimulation provided by the bipartite enhancer is developmentally specific, we assayed its activity in NIH 3T3 fibroblasts and in differentiated female ES cells. In both backgrounds, the enhancer did not stimulate the *Tsix* promoter (Fig. 5E). Additionally, replacement of upstream or downstream sequences with heterologous bacteriophage sequences had little to no effect on luciferase activities in these differentiated cell types. Thus, as with the *Xite* enhancer, the stimulation conferred by the bipartite enhancer was specific to undifferentiated ES cells. We suggest that, together with the *Xite* enhancer, the bipartite enhancer contributes to the developmental specificity of *Tsix* transcription.

We next examined whether the bipartite enhancer might add to the effect of the *Xite* enhancer in regulating *Tsix* (Fig. 5F). Indeed, the presence of both the *Xite* enhancer and the bipartite enhancer augmented luciferase expression, yielding ~40% greater stimulation than the bipartite enhancer alone (com-

pare $-10.2/+4.0$ and $-9.0/+4.0$). This finding is consistent with the ideas that the *Xite* enhancer and the bipartite enhancer can function in parallel and that these elements work together to control both the level and the proper developmental timing of *Tsix* expression in vivo.

***Tsix* enhancers are associated with developmentally specific DHS.** In vivo, transcriptional enhancers are often associated with an open chromatin conformation. DHS mapping in the *Xite* region previously revealed several DHS specific to undifferentiated ES cells, with a strong DHS coinciding with the 1.2-kb enhancer and several weaker ones located in close proximity, approximately -7.7 and -8.8 kb from the major *Tsix* promoter (30). Thus, both the *Xite* enhancer and the 5' portion of the bipartite enhancer are associated with developmentally specific DHS. To determine whether the remainder of the bipartite enhancer also lies in specialized chromatin, we performed DHS mapping for the $-5.9/+4.0$ region.

We found several novel DHS in this region. First, within the upstream half of the bipartite enhancer, a single DHS was

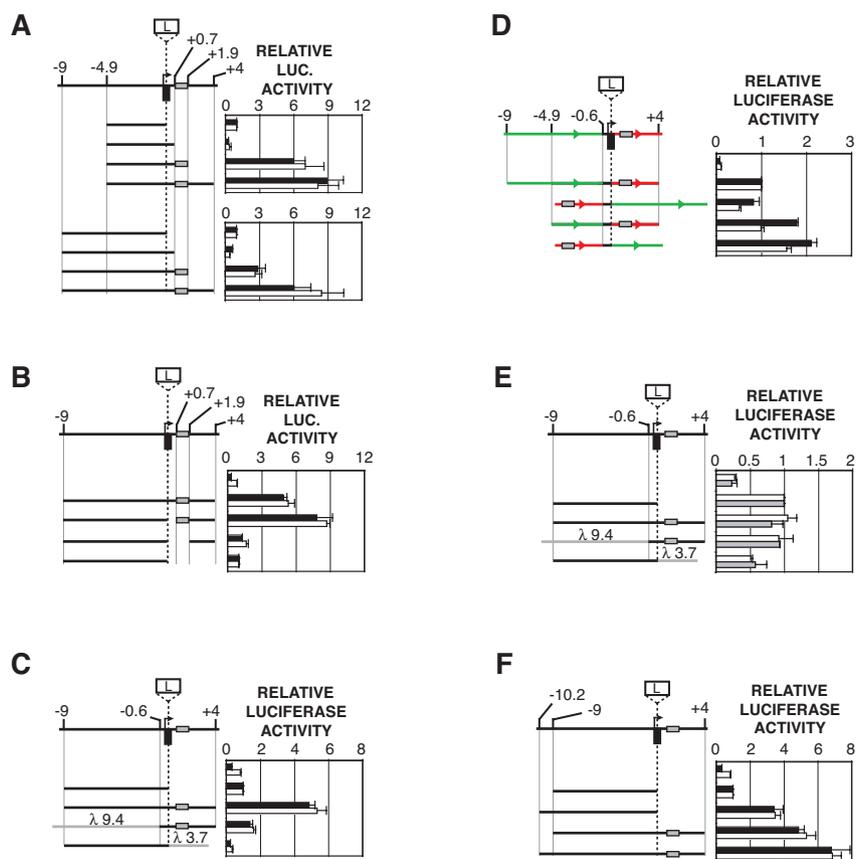


FIG. 5. A bipartite enhancer flanks the *Tsix* promoter and is developmentally specific. In sequence schematics, alternatively spliced exons 2' and 3 are omitted for clarity; L, luciferase cassette. *DXPas34* is shown as a rectangle between kb +0.7 and +1.9. For all panels except E, mean luciferase activities are shown for transiently transfected male ES cells (black bars) and female ES cells (white bars). For all panels except D, the activity of upstream sequences in isolation was set to 1. Error bars denote one standard error. (A) Synergistic stimulation of the *Tsix* promoter by combinations of upstream and downstream sequences. (B) *DXPas34* is required for stimulation. (C) Stimulation depends specifically on *Tsix* sequences. Bacteriophage λ sequences are labeled. (D) Swap of upstream and downstream sequences preserves stimulation. Orientations of upstream (green) and downstream (red) sequences are indicated. The activity of the native configuration was set to 1. The backbone was pNS185. (E) Combinatorial stimulation is absent in differentiated cells. Mean luciferase activities are shown for transiently transfected differentiated female ES cells (days 11 to 17) (lower bar in each set) and male NIH 3T3 fibroblasts (upper bar in each set). (F) Stimulation of the promoter by the 1.2-kb enhancer and more proximal upstream and downstream regions.

observed approximately -3.7 to -3.6 kb upstream of the *Tsix* promoter in both undifferentiated male and female ES cells (Fig. 6A and data not shown). This DHS was also present in differentiating ES cells and in fibroblasts, indicating that it is constitutive. Second, within the $-0.7/+4.0$ region, analysis of undifferentiated female and male ES cells revealed three DHS. Two were observed in a prominent doublet upstream of the EcoRV site at bp +144, corresponding to the two start sites of the major *Tsix* promoter (Fig. 6B and data not shown). This doublet was also present in differentiating ES cells, where *Tsix* expression is reduced, but was absent in primary fibroblasts, where *Tsix* expression is silenced. Thus, hypersensitivity at the *Tsix* promoter correlates with the pattern of expression during differentiation.

The third DHS was located downstream of the promoter at kb +1.4, approximately within or just downstream of *DXPas34*. The band corresponding to this DHS was consistently diffuse in female ES cells, perhaps because of the length polymorphism in *DXPas34* between the 129 allele and the *Mus. castaneus* allele (3). Consistent with this interpretation, the DHS band in

male ES cells (carrying only the 129 X chromosome) was consistently more distinct (data not shown). The DHS was observed in undifferentiated female and male ES cells, was markedly reduced in differentiated ES cells, and was undetectable in fibroblasts (Fig. 6B and data not shown). These results indicate that the downstream half of the bipartite enhancer is associated with specialized chromatin that changes dynamically with cell differentiation and the onset of XCI. Interestingly, the DHS at *DXPas34* was lost before that of the promoter (Fig. 6B), supporting a model in which the bipartite enhancer regulates the activity of the *Tsix* promoter in vivo.

Taken together with previous results (30), these findings map eight DHS in the *Tsix-Xite* region from kb +4.0 to -12.0 . While two map to the major *Tsix* promoter, the remainder correspond to regions that are associated with transcriptional enhancer elements. All but one of these DHS display developmentally specific changes that correlate with the pattern of *Tsix* expression during differentiation. We propose that the dynamic changes in *Tsix* transcription are regulated by the coordinate action of enhancer elements that span the *Xite* and *Tsix* loci.

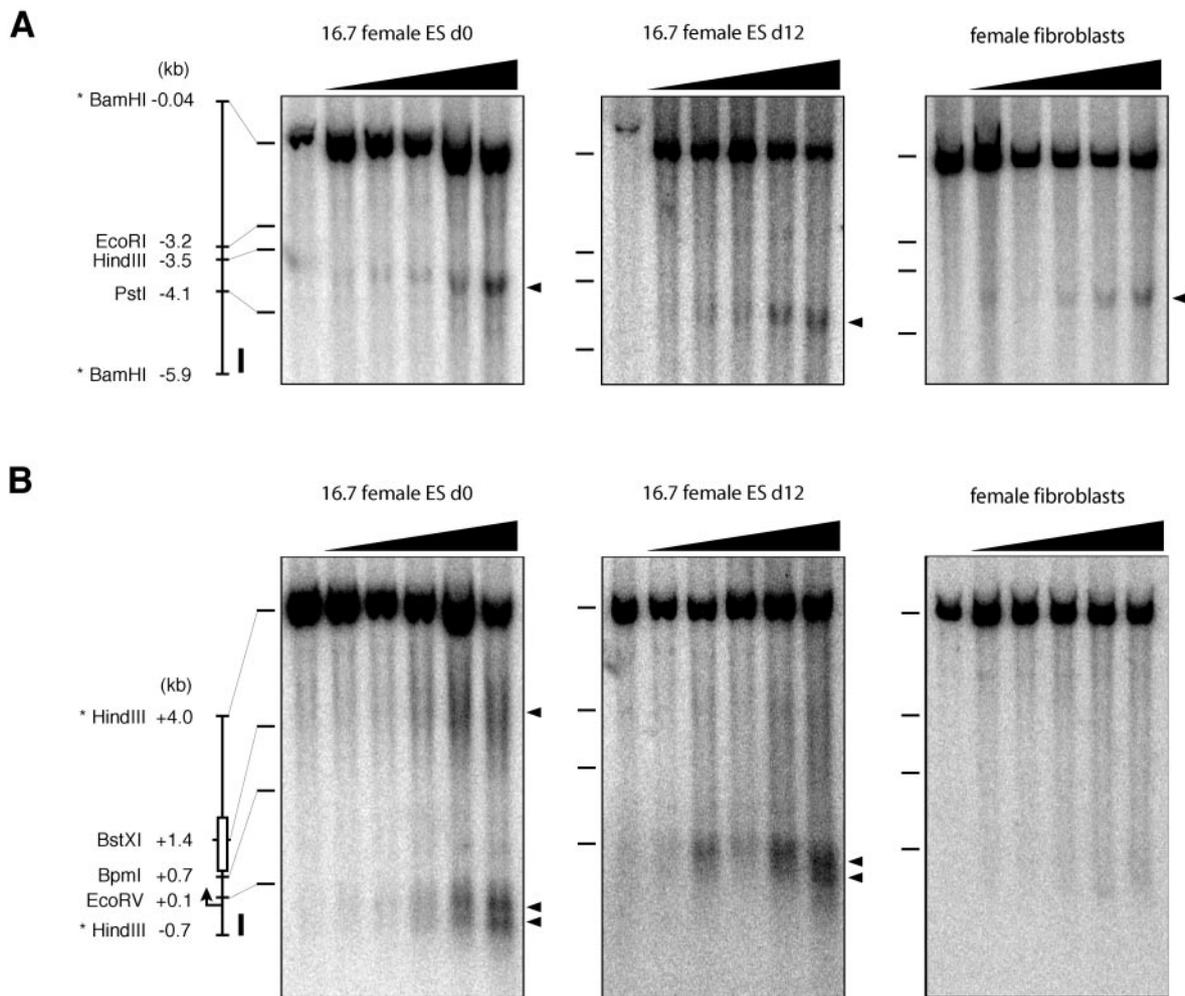


FIG. 6. DHS mapping in the bipartite enhancer. Sequence schematics of analyzed regions are shown at the left. The positions of *DXPas34* (white rectangle), the major *Tsix* promoter, and hybridization probes (black rectangles) are indicated. Mobilities of control DNA fragments terminating at the indicated restriction sites are shown to the left of each panel by horizontal tick marks. Positions of hypersensitive sites are shown to the right of each panel by arrowheads. From left to right, the DNase concentrations used for ES cells were 0, 0.5, 1, 2.5, 5, and 10 $\mu\text{g/ml}$ and those used for fibroblasts were 0, 5, 10, 20, 40, and 80 $\mu\text{g/ml}$. (A) DHS mapping from -5.9 to -0.04 kb. (B) DHS mapping from -0.7 to $+4.0$ kb.

DISCUSSION

Novel transcriptional switches for *Tsix*. The initiation of XCI in female cells is controlled by antagonistic interactions between *Xist*, the trigger for chromosomal silencing, and *Tsix*, its antisense regulator. During cell differentiation, this molecular interplay leads to opposite fates on the two X chromosomes: on the X_a chromosome, the persistent expression of *Tsix* prevents the spread of *Xist* RNA, and on the X_i chromosome, the extinction of *Tsix* expression allows *Xist* RNA to initiate the formation of heterochromatin. Thus, the control of XCI depends upon the mechanisms that regulate *Tsix* expression during differentiation. In the present work, we have identified the first enhancers at the X chromosome inactivation center (Fig. 7A). These enhancers constitute novel activities which act as developmentally appropriate transcriptional switches for *Tsix*.

One enhancer maps within *Xite* and appears to behave as a classical enhancer. The second lies at the 5' end of *Tsix* and

possesses a bipartite structure. While each is a self-sufficient enhancer unit, the combination of the two results in greater stimulation of the *Tsix* promoter. DHS mapping reveals that the 16-kb domain from kb -12.0 to $+4.0$ around the *Tsix* promoter is replete with DHS, suggesting the occurrence of a specialized chromatin structure. Therefore, we propose that *Tsix* is coordinately regulated by enhancer activities in this 16-kb domain. These enhancers work together to establish proper kinetics for *Tsix* expression and reciprocal behavior of its two alleles—that is, persistence of *Tsix* on the future X_a chromosome and extinction on the future X_i chromosome.

A potential mechanism of enhancer action is promotion of the persistence of *Tsix* expression. At what level might these enhancers work on *Tsix*? Several studies of the *Xite* and *Tsix* loci provide clues to their molecular function. A 12-kb deletion of *Xite* that encompasses the 1.2-kb enhancer does not significantly affect *Tsix* expression in undifferentiated ES cells but causes the premature extinction of *Tsix* expression in *cis* during

cell differentiation and the onset of XCI (30). This analysis suggests important mechanistic attributes of the *Xite* enhancer. First, *Tsix* expression does not require the *Xite* enhancer in undifferentiated ES cells, despite the fact that the enhancer appears to be active in these cells, as indicated by transfection and DHS analysis. Second, the *Xite* enhancer functions to promote the persistence of *Tsix* expression rather than to increase the absolute level of *Tsix* expression. Together, these findings suggest that the environment for *Tsix* expression becomes progressively more restrictive as differentiation proceeds, with the *Xite* enhancer functioning to counteract these restrictive influences and promote the monoallelic persistence of *Tsix* expression in *cis* (Fig. 7B).

Although specific knockouts of the bipartite enhancer have not been made, several deletions have included *DXPas34* and downstream portions of the enhancer (19, 22, 35). Although these deletions include the major *Tsix* promoter, they cause skewed XCI, with a bias toward inactivating the mutated X chromosome, a phenotype that is broadly consistent with the proposed role of *DXPas34* and downstream elements in positively regulating *Tsix* expression. In a 49-kb knockout in which the upstream half of the bipartite enhancer and all of *Xite* are deleted (29), XCI is also biased in favor of silencing the mutated X chromosome, supporting a role for both regions in regulating *Tsix* expression. Significantly, the 49-kb deletion displays a more severe bias in the skewing of XCI than does the deletion of *Xite* alone, suggesting that the *Xite* and bipartite enhancers each contribute to promoting the persistence of *Tsix* expression. Clearly, however, knockouts of specific elements in the bipartite enhancer, including *DXPas34*, are required to confirm this function and to determine whether the bipartite enhancer functions in a manner similar to that of the *Xite* enhancer.

Activator versus insulator models. The identification of CTCF-binding sites within and around *DXPas34* has suggested two potential (but not mutually exclusive) mechanisms of action (8). One model of *Tsix-Xist* regulation invokes enhancer competition, in which the two genes compete for access to a shared enhancer (8). In this model, differential access on the two X chromosomes is determined by a chromatin insulator (CTCF), whose allele-specific binding to *Tsix* on the future X_a chromosome blocks the shared enhancer from the *Xist* promoter. Here we tested whether the 1.2-kb *Xite* enhancer could be the hypothetical shared enhancer and found that it could not stimulate *Xist* promoters in transient assays, suggesting that the 1.2-kb *Xite* enhancer might be specific for *Tsix*, at least in the context of these experiments. With other DHS in the region, it is possible that the 1.2-kb region (which spans just one DHS) is not sufficient to recapitulate full enhancer function and that, through the inclusion of additional sequences, the *Xite* enhancer actually stimulates *Xist* as well as *Tsix*. It is also possible that the shared enhancer lies elsewhere in *Xic*.

The location and structure of the bipartite enhancer at the 5' end of *Tsix* make it unlikely to be the shared enhancer. While mammalian enhancers tend to be intronic or distinct from the structural genes that they stimulate, the *Tsix* enhancer embodies about 9 kb of sequence around the promoter itself. Most curiously, the downstream half of the bipartite enhancer actually overlaps the proposed CTCF insulator domain in *Tsix* (Fig. 7A). This finding raises several points. First, in the insulator-

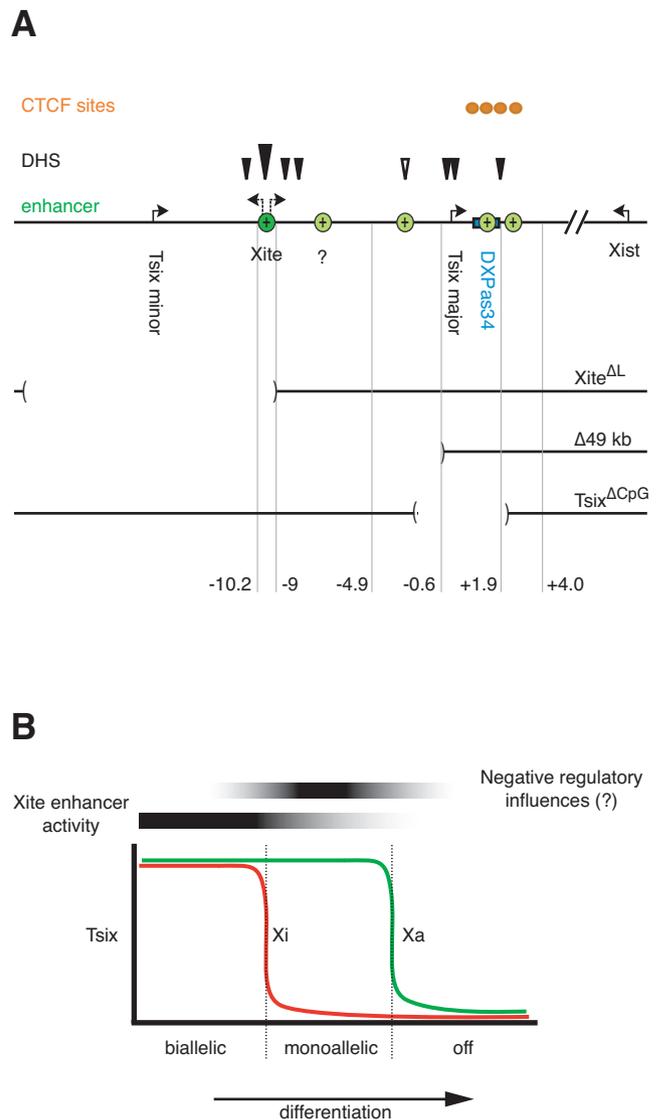


FIG. 7. *Tsix* control elements and *Tsix* binary switch model. (A) Summary of *Tsix* transcriptional regulatory sequences known to date. Embryonic specific stimulatory elements are shown in the context of genes, functional elements, and selected deletions in *Xic*. The novel enhancers are shown in green: dark green, 1.2-kb enhancer; light green, bipartite enhancer with multiple elements therein. The exact position of the light green element marked by a question mark is not known. Circles, CTCF elements; closed arrowheads, ES-specific hypersensitive sites; open arrowheads, constitutive DHS; largest arrowhead, strongest DHS. (B) Model for the binary switch through positive and negative control of *Tsix* expression. In this model, the *Xite* enhancer promotes the persistence of *Tsix* expression in *cis* but does not exert a major influence on the quantitative level of *Tsix* expression. The activity of the *Xite* enhancer is lost gradually as differentiation proceeds and, in conjunction with the induction of unidentified negative regulatory influences, yields the asynchronous silencing of *Tsix* expression (21).

mediated enhancer competition model, the insulator must lie between the promoters of the two competing genes and the enhancer must lie on the other side of either promoter (8). Thus, the placement of the bipartite enhancer around the *Tsix* promoter violates the requirement for distal placement, mak-

ing it an improbable shared enhancer. Second, the overlap of CTCF-binding sites in *Tsix* poses a certain quandary: what is the functional consequence of juxtaposing enhancer and insulator activities? In *Tsix*, CTCF-binding sites are dispersed across the region from kb +0.7 to +4.0, with *DXPas34* itself harboring some of these elements. As proposed previously for *Tsix*, CTCF might have both activating and insulating properties (8)—a scenario similar to what has been postulated for CTCF action at *H19* (12). The juxtaposition of an enhancer and an insulator has also been described for the β -globin locus (5). At *Tsix*, this configuration might be responsible for some observed orientation-dependent effects. For example, while *DXPas34* works well as a chromatin insulator in one orientation, it has a relatively weaker effect in the opposite orientation (8). Juxtaposition of insulator and enhancer sequences might impart directional character to enhancer action. This architecture might be integral to the mechanism of *Tsix* control—a possibility to be tested in the future.

A second model of *Tsix-Xist* regulation envisions CTCF as a direct transcriptional activator of *Tsix* rather than as a chromatin insulator between *Tsix* and *Xist* (8). In other systems, CTCF apparently has direct activating potential (40, 41). The data accrued here are consistent with such a model for XCI regulation. Indeed, we have shown that the bipartite element (in which the CTCF-binding sites reside) is a transcriptional enhancer of *Tsix*. Given its context dependence, this enhancer appears to be highly specific for *Tsix*. Given the similar *Tsix* specificity of the *Xite* enhancer, we propose that the two enhancers might actually work synergistically to bolster *Tsix* expression. Furthermore, the activator or enhancer must work allele specifically so that stimulation takes place only on the future X_a chromosome. The activator model is also consistent with other lines of evidence. For example, deletions which encompass one or both enhancers (9, 28–30) compromise *Tsix* expression and favor *Xist* expression in *cis*. Importantly, however, while the data here are consistent with an activator model, the activator and insulator models are not mutually exclusive. Both mechanisms might be required to set up the critical pattern of *cis* regulation between *Tsix* and *Xist*.

The binary switch, X chromosome counting, and epigenetic choice. With the delineation of activating sequences, our present work addresses how *Tsix* expression is preserved on the future X_a chromosome. However, the constitutive activity of the major *Tsix* promoter indicates that *cis*-acting silencers or other repressive mechanisms must also exist (Fig. 7B). This notion is further supported by the fact that *Tsix* silencing is obligatory for *Xist* upregulation on the future X_i chromosome (24, 38). Clearly, to gain a full understanding of *Tsix* regulation, future work must address how *Tsix* is turned off on the X_i chromosome. We suggest that enhancer action on the future X_a chromosome, together with negative regulatory influences, comprises the binary switch for X chromosome activity.

The molecular bases of X chromosome counting and choice remain two major goals of ongoing research. X chromosome counting elements have been shown to reside in a 37-kb region downstream of *Xist* that includes *Tsix* and *Xite* (29). In particular, a comparison of various deletion mutants has further suggested that counting elements are located within a discontinuous 20-kb sequence (29) that overlaps the bipartite enhancer. Thus, the bipartite enhancer may function in X chro-

mosome counting. Similarly, elements involved in X chromosome choice have been argued to reside in *Xite* (30), in *DXPas34* and the region flanking the major *Tsix* promoter (20, 22, 37), and in *Xist* itself (26, 32). Indeed, we believe that counting, choice, and the *Tsix* binary switch are tightly coupled events and predict that the counting and choice mechanisms will be revealed through further dissection of the enhancers identified here.

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