

# Inactive X chromosome-specific histone H3 modifications and CpG hypomethylation flank a chromatin boundary between an X-inactivated and an escape gene

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## ABSTRACT

In mammals, the dosage compensation of sex chromosomes between males and females is achieved by transcriptional inactivation of one of the two X chromosomes in females. However, a number of genes escape X-inactivation in humans. It remains poorly understood how the transcriptional activity of these 'escape genes' is maintained despite the chromosome-wide heterochromatin formation. To address this question, we analyzed a putative chromatin boundary between the inactivated *RBM10* and an escape gene, *UBA1/UBE1*. Chromatin immunoprecipitation revealed that trimethylated histone H3 lysine 9 and H4 lysine 20 were enriched in the last exon through the proximal downstream region of *RBM10*, but were remarkably diminished at ~2kb upstream of the *UBA1* transcription start site. Whereas RNA polymerase II was not loaded onto the intergenic region, CTCF (CCCTC binding factor) was enriched around the boundary, where some CpG sites were hypomethylated specifically on inactive X. These findings suggest that local DNA hypomethylation and CTCF binding are involved in the formation of a chromatin boundary, which protects the *UBA1* escape gene against the chromosome-wide transcriptional silencing.

## INTRODUCTION

The unbalanced gene dosage of sex chromosomes between males (XY) and females (XX) represents an impediment to normal development. In mammals, X-chromosome

inactivation (XCI) is achieved by transcriptional silencing of all but one of the X chromosomes in a diploid female cell, to equalize the gene dosage of X chromosomes between males and females (1). The *XIST/Xist* gene, which maps to the X-inactivation center, is expressed from the inactive X-chromosome (Xi) in female somatic cells (2). *XIST/Xist* RNA is essential for the initiation of XCI (3,4), playing a key role as a *cis*-acting factor in recruiting the silencing machineries. Once XCI is initiated, genes on the X chromosome undergo inactivation by sequential epigenetic modifications, and an inactive chromatin status characteristic of heterochromatin is stably maintained through cell generations. A number of genes, however, escape XCI in human cells and are transcriptionally active (5,6); more than 15% of human X-linked genes are reported to escape the inactivation (7). These escape genes can be categorized into two groups based on the presence or absence of Y-homologs.

The distal ends of the short and long arms of the human sex chromosomes are known as pseudoautosomal regions 1 and 2 (PAR1 and PAR2), which have homologous sequences with Y chromosome and are required for X–Y pairing in male meiosis. Most of the genes mapped on PARs escape XCI (7,8).

A number of genes that do not have Y-homologs, on the other hand, also escape inactivation in human cells. However, these genes are mostly inactivated in mice (9). Such different sets of escape genes between human and mice may reflect evolutionary differences and could explain the distinct phenotypes when one X is missing. The absence of a single X-chromosome in humans (45, X) results in Turner syndrome, which shows a more severe phenotype compared to X0 mice (10). This suggests that double dose of transcription found in escape genes without Y-homolog is crucial for normal female development in human but not in mice, but the

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significance of the dosage of escape genes remains poorly understood.

Post-translational histone modifications like acetylation and methylation play important roles in the epigenetic control of gene expression in eukaryotes, in addition to DNA methylation (11–13). Hypoacetylation of histone H4 and methylation at lysine 9 and 27 of H3 (H3K9 and H3K27), for instance, are associated with chromatin compaction and gene repression. In contrast, euchromatic domains display higher levels of histone acetylation (14–17). Recent studies established that the methylation of H3K9 and H3K27 is an early event in the XCI process (18–22) proceeding to DNA methylation on the promoters of inactivated genes (23). In contrast to the inactivated genes, the escape genes lack DNA methylation and H3K9 methylation and are associated with euchromatic features like histone acetylation, H3K4 methylation, and early replication (24,25).

Here we analyse the transcriptional status of escaped genes and their neighbouring genes to analyze chromatin boundaries in detail. For this, we used human–mouse hybrid cells carrying either active or inactive human X chromosome in mouse background. We then focused on the *RBM10-UBA1/UBE1* locus on Xp11.23, where the inactivated *RBM10* and escape *UBA1* are separated by only four kilobases of intergenic sequences. By profiling histone modifications using chromatin immunoprecipitation (ChIP), we detect a chromatin boundary in the *RBM10-UBA1* intergenic region. Trimethylated H3K9 and H4K20 (H3K9me3 and H4K20me3) were enriched in the last exon through the proximal downstream region of *RBM10* but were strongly diminished at ~2 kb upstream of *UBA1* on Xi. As previously found in other boundaries on Xi (26), ChIP also revealed association of CTCF to the intergenic region, suggesting the involvement of this zinc finger protein in maintaining the transcriptional activity of *UBA1* and its downstream escape genes.

## MATERIALS AND METHODS

### Cells and cytogenetics

A9 (7149)-5 (27) and CF150 (28) cells harboring human active and Xi chromosomes, respectively, were generous gifts of Dr M. Oshimura and Dr T.K. Mohandas. All cell lines were maintained in Dulbecco's modified Eagle's medium (DMEM) supplemented with 10% fetal bovine serum in a 5% CO<sub>2</sub> incubator.

For cytogenetic analysis, cells were incubated in 100 µg/ml 5-bromo-2'-deoxyuridine (BrdU) for 6 h, 1 µg/ml colcemid was added to the culture medium, and cells were further incubated for 1 h. Chromosome spreads and staining were prepared according to the method described previously (29). Briefly, cells were treated with 75 mM KCl for 10 min, fixed with 3:1 methanol: glacial acetic acid on ice, and air-dried on clean glass slides. Chromosome spreads were stained with freshly prepared acridine orange (Sigma) and examined under a fluorescence microscope (BX-81; Olympus) using an oil-immersion 100× UPlanApo objective lens (NA: 1.35)

equipped with a cooled CCD (ORCA-ER; Hamamatsu Photonics).

### RNA extraction and RT-PCR

Total RNA was prepared from each cell line using TRIzol (Invitrogen) and treated with RNase-free DNase I (Roche). To check the transcriptional status of X-linked genes, cDNA was synthesized from 1 µg RNA using SuperScript VILO cDNA synthesis kit (Invitrogen) as described by the manufacturer. To detect the sense/antisense transcripts in *RBM10-UBA1* intergenic region, cDNA was synthesized from 2 µg RNA using One-step RT-PCR kit (Qiagen) using a strand-specific primer as described by the manufacturer. To avoid primer-independent reverse transcription due to the secondary structure, the reaction mixture was incubated at 60°C.

Quantitative PCR was performed with Power SYBR Green PCR Master Mix (Applied Biosystems) using a 7500 FAST (Applied Biosystems). Each PCR was run in triplicate to control PCR variation. All primers used here (summarized in Table 1) have proven to be species-specific.

### Chromatin immunoprecipitation using native chromatin

We used mouse monoclonal antibodies specific to different H3 modifications (30) and commercial antibodies including di + tri-methyl H3K4 (H3K4me2 + 3; Abcam; ab6000), di-methyl H3K9 (H3K9me2; Abcam; ab1220), tri-methyl H3K27 (H3K27me3; Abcam; ab6002), tri-methyl H4K20 (H4K20me3; Abcam; ab9053). Control mouse IgG (Jackson Immunoresearch) was also used.

To profile the histone modifications, native chromatin was prepared and used for chromatin immunoprecipitation (ChIP) assay as described previously (15,31) with slight modifications. Briefly, ~15 µg of native chromatin (~2–4 nucleosomes) was incubated with 15–20 µl hybridoma culture sup or 10–20 µg IgG for 15 h at 4°C in ChIP buffer [25 mM Tris-HCl (pH 7.6), 500 mM NaCl, 5 mM EDTA and Protease inhibitor cocktail (Roche; Complete, EDTA-free)]. After further incubation for 2 h with 80 µl Dynabeads M-280 Sheep anti-mouse IgG (for mouse IgG; Dynal), or 50 µl Dynabeads Protein G (for rabbit IgG; Dynal), chromatin-antibody complexes ('bound' fraction) were separated from antibody-free chromatin ('unbound' fraction) using a magnetic stand (Dynal). After rinsing three times with ChIP buffer, bound-chromatin was eluted from the beads using ChIP buffer containing 1% SDS. At least three independent ChIP experiments were performed for each antibody.

### ChIP using cross-linked chromatin

To analyze the binding of CTCF and RNA polymerase with the chromatin, we used another ChIP method using cross-linked chromatin as described before (30,32). Sonicated chromatin (~300–1500 bp; ~15 µg) was incubated with mouse monoclonal anti-CTCF (BD transduction Laboratories, 612148; 15 µg), anti-RNA polymerase, B8-1 (33) (10 µg), 8WG16 (Covance; 10 µl) and H14 (Covance; 10 µl), or normal mouse IgG

**Table 1.** PCR primers

Application	Gene name	Forward (5' to 3')	Reverse (5' to 3')	
RT-PCR	<i>POLA</i>	aaagggcagatgaggaac	ttccaaacagaaataccacac	
	<i>PGK1</i>	aaagggaaagcgggtcgatc	tgcccagcagagatttgagt	
	<i>XIST</i>	gatgggtgccagctatc	ggatcctcatgcccatctc	
	<i>RBM10</i>	ggacctatgttcccgcc	ctcgtcatctcagaggga	
	<i>UBA1 TSS1</i>	ttccgagctccgcgcaaac	gcacttcggacaacacggactg	
	<i>UBA1 TSS1'</i>	ttgtgtcggcgtcggctgtaag	gcacttcggacaacacggactg	
	<i>USP11</i>	catgaaacatccattcttctcagtc	ctgagtcaggcatcgacgaa	
	<i>ZNF157</i>	aggaccctctacacagacagc	ctgtcgggtgaaatccacagccac	
	<i>KLHL15</i>	atcattcagaatatccggttttgct	ttcaatgcttggtcaactctgtaa	
	<i>EIF2S3</i>	ctgtcctgccacgataattt	aggtgttccgatgtctgagg	
	<i>ZFX</i>	ctgatcctctgactaccgactagtt	aatcataaggtagctcactcaggtgcc	
	<i>Mouse Thp</i>	accacccctgtacccttc	tttacagccaagattccagtgatg	
	ChIP and strand-specific RT-PCR	<i>RBM10 Ex20</i>	gaggactggtggcagccta	ccagtcggtgagcttctct
		<i>RBM10 Ex22</i>	caactgaacgcagagaaaag	atatgccgcctgacttctct
		<i>RBM10 Ex24</i>	ggcgaggggaaagacagagtg	gatttggccaagccctctctg
		<i>RU1</i>	ctagcaccaccaggcatgaa	tccagatctaggatggcaaa
		<i>RU2</i>	ccagcaatttaggggctga	tgcgccagcctaattttat
<i>RU3</i>		ggaacaacccaaatgtccaa	caccatgtgaaacatctacca	
<i>RU4</i>		tcaaatgccaaaactgttcca	cacatcatctggccctttt	
<i>RU5</i>		accacagatgaccccacta	tgaggcaggatgattgcttg	
<i>RU6</i>		ccacctgagctcggctcctc	gcaccagcgtccagaataatg	
<i>UBA1 Ex1</i>		ttccgagctccgcgcaaac	ttcctgcaaaagatgagctgaagc	
<i>Intron(In) 1-1</i>		ggaggctcaggactgacgagac	ggcgctctgacgctcagga	
<i>In1-2</i>		gggatgaggcctctacatgttt	cctcataccattattcaggttac	
<i>In1-3</i>		ctccatgcgaaggtaagtgtgctg	gggatgagccttagaacaattgctggc	
<i>In1-4</i>		gctgggattacaggcgtcatct	ccgctcagggaacaagg	
<i>In1-5</i>		gcagcaatagcggggcgt	gccacacacccccattcat	
<i>UBA1 Ex1'</i>		ttgtgtcggcgtcggctgtaag	ccgccaactcctcaaggagccgaa	
<i>In1'-1</i>		acatgctagccgcgctcag	ggaaagatgctaggcaggg	
<i>In1'-2</i>		tgcagtcaggggtttcagctt	cctggacacatagtgaaacaattctg	
<i>In1'-3</i>		caccctcagagcagctgtgatacc	gaggctgtgaggggtgagtg	
<i>In1'-4</i>		cccagaaaagcttgacatgctcc	gggtttgagggaggagctctggag	
<i>In1'-5</i>		cctcaggtcagtgagttctgaccacag	ctgactgacctggggtagagatctc	
<i>In1'-6</i>		tcagtgtgggaggactgtgtgaa	ctggtgtgggggtgactg	
<i>In1'-7</i>		cggttttcccctcccaaac	gagtgacaggaggaggtgagggc	
<i>UBA1 Ex2</i>		atgtccagctcggcgtgccaag	gcacttcggacaacacggactg	
<i>XIST + 1</i>		ctctcagttcttaagcgcgctcaa	ggatgttttttaagaaatagccat	
<i>XIST + 780</i>		tgggtgttgcaactctctgg	ctgctgacctgctatcatcc	
<i>PGK1 + 98</i>		gcacgtcggcagtcggct	cctcataacgaccgcttccc	
<i>PGK1 + 700</i>		acttgtggcctccaatctc	agaaaccgtgttgcaagt	
<i>RBM10 + 67</i>		gtagcggcagtgagtttcc	tctccaactctccagctc	
<i>RBM10 + 600</i>		ccacttattgggtgggaagg	tttccatggatgagcgtcag	
<i>H19 ICR</i>		ccgagaaaatagccattgcctacagt	catgttctttgagctcgggtgta	
<i>Eif2</i>		ggttgagggcagatgaga	cggcaacatgctgttaagac	
<i>KIAA0522</i>		agccagctgggataaggaat	ctggggtcttccactaaca	
Bisulfite sequencing		<i>RBM10 Ex24</i>	gggtaggggaaggatagagtgtg	aaactcacctcccccctcta
		<i>RU2 to RU3</i>	ggtttttgggtttgggtatagtgagt	aaaatttcaacttataccccaaac
		<i>RU5</i>	agtttgtaaagtgatgggattatagg	ccaaactaaactaaactcctaaactcaaac
		<i>RU6 to UBA1 Ex1</i>	gggtgtgtttttatgtggaggt	caccctctccaaataactatc
	<i>In1-1</i>	ggtattgtggaggtgttaggtttttgtg	caaaaacccccctcaacacgctcc	
	<i>In1-4</i>	ggtgtgaaattgaggtattgttaag	caccctaaactacctaaccacac	
	<i>UBA1 Ex1'</i>	gtttttgggggtgggttttaag	aaatccactccccctcctactc	

(Jackson Immunoresearch; 5 µg) for 15 h at 4°C in RIPA buffer [50 mM Tris-Cl (pH 8.0), 150 mM NaCl, 1 mM EDTA, 1% Triton X-100, 0.1% sodium deoxycholate and Protease inhibitor cocktail (Complete, EDTA-free; Roche)]. After further incubation for 2 h with 80 µl Dynabeads M-280 Sheep anti-mouse IgG (Dyna), or anti-mouse IgM (Dyna) for H14, immunoprecipitates were collected, washed with RIPA buffer containing 150 mM and 500 mM NaCl, and resuspended in 200 µl of 10 mM Tris-HCl (pH 8.0), 5 mM EDTA, 300 mM NaCl and 0.5% SDS. The chromatin-bound beads

(‘bound’ fraction) and the input fraction were incubated at 65°C for 12 h to reverse the cross-linking.

### Quantitative analysis of ChIP samples

DNA was extracted from the input, ‘bound’ and ‘unbound’ fractions with phenol/chloroform, precipitated with ethanol, and re-suspended in 60 µl TE [10 mM Tris-HCl (pH 7.6) and 1 mM EDTA]. The DNA sample (1 µl) was used for quantitative PCR with QuantiTect SYBR Green PCR (Qiagen) using an Opticon2 (BioRad). Each

PCR was run in triplicate to control PCR variation. Primers used here are listed in Table 1.

**DNA methylation assay**

Bisulfite treatment was carried out using EZ DNA methylation kit (ZYMO RESEARCH) according to the manufacturer’s instruction, and the completion of reaction was confirmed using control samples containing non- and fully-methylated templates. To amplify bisulfite-treated genomic DNA, primers for PCR amplification are designed to contain degenerate nucleotides (C to T) except CpG dinucleotide sequence (Table 1). PCR products were cloned into Topo-TA vector (Invitrogen) and the nucleotide sequence was determined using a PRISM3100-Avant (ABI).

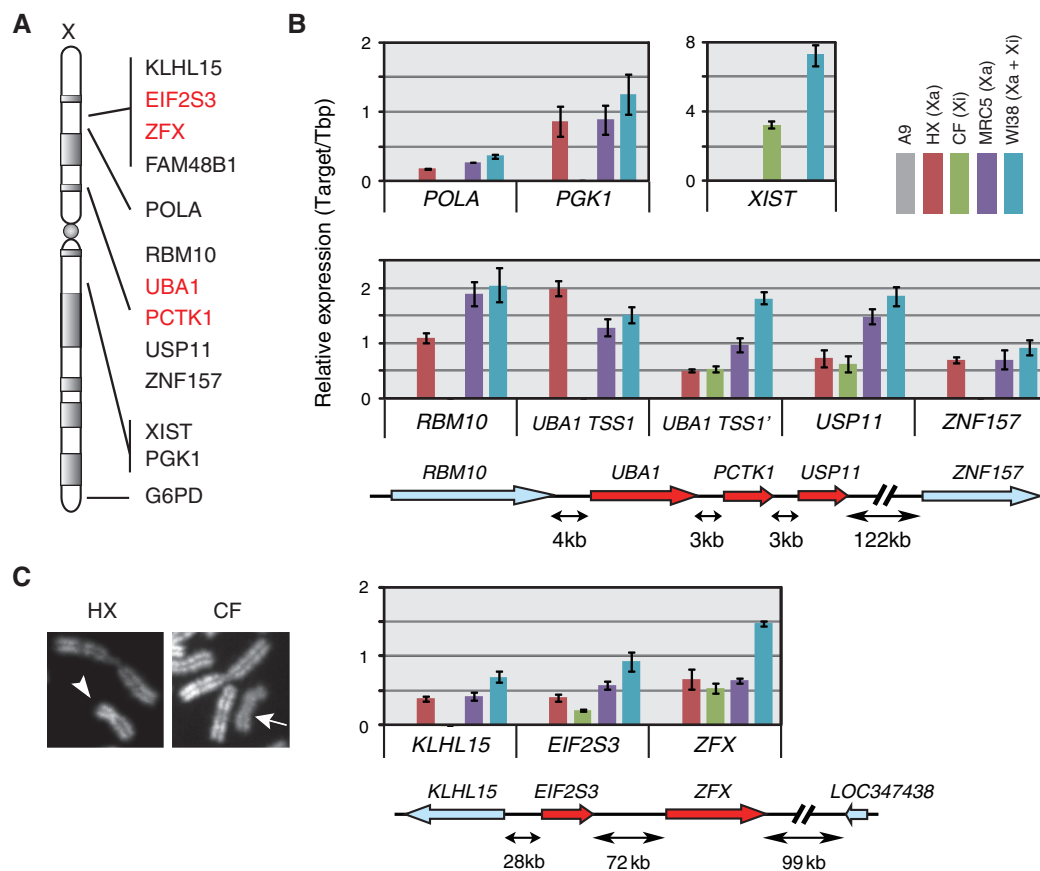
**RESULTS**

**Status of human X chromosomes in hybrid cell lines resembles that in diploid human cells**

To analyze active and inactive X chromosomes (Xa and Xi) separately, we used human–mouse hybrid cell lines,

A9 (7149)-5 [hereafter described as HX; (27)] and CF150 [CF; (28)] harboring active and inactive human X-chromosome, respectively. These were generated by microcell-mediated chromosome transfer into mouse A9 cells.

We first examined whether the active or inactive status of human X chromosomes was maintained in HX and CF cells. By quantitative RT-PCR, the expression of several human X-linked genes was examined in A9 (parent mouse cells; a negative control), HX and CF, and compared with that in diploid human male and female cells (i.e. MRC5 harboring Xa and WI38 harboring both Xa and Xi, respectively) (Figure 1A and B). The expression of typical human X-linked genes that are repressed on Xi (i.e. *POLA*, *PGK1* and *G6PD*) was detected in cells harboring human Xa, such as HX, MRC5 and WI38 (Figure 1B, top; data not shown). In contrast, *XIST* was expressed in cells harboring human Xi (CF and WI38; Figure 1B, top). To evaluate the replication timing of human X chromosomes in hybrid cells, the cytogenetic examination using R-banding (29) was employed. In chromosome spreads from HX cells, human X exhibited a typical banding pattern of active X (Xa) due to its asynchronous



**Figure 1.** The status of human X chromosomes in the human–mouse hybrid cell lines. (A) Schematic human X-chromosome map with the inactivated and escape genes examined in this study. (B) Quantitative RT–PCR analysis of human X-linked genes in different cell lines. Total RNA was prepared from A9 (parental mouse cell line for HX and CF: without human X), HX (mouse line harboring active human X), CF (mouse line harboring inactive human X), MRC5 (46, XY; male diploid human cells harboring active X) and WI38 (46, XX; female diploid human cells harboring active and inactive Xs). The levels of transcripts were analyzed using real-time PCR, and expressed as relative amounts to human TBP (for MRC5 and WI38) or mouse Tbp (for A9, HX and CF). (C) Representative X chromosome banding patterns. After the incubation with BrdU for 6 h to label late replicating DNA, cells were treated with colcemid for chromosome spreading and R-banding. A typical human active X chromosome is observed in HX cell spread (left panel; arrowhead). In CF cell spread, human inactive X chromosome shows homogenous dimmer staining (right panel; arrow).

replication, whereas all human X in CF spreads appeared as homogenous late replication staining, characteristic of the Xi (Figure 1C). These results indicated that human Xa and Xi were stably maintained in hybrid cell lines HX and CF, respectively (27,28).

To explore the chromatin boundary between inactivated and escape genes, we next validated the expression of three previously reported escape genes (2,7,34,35), *UBA1* (ubiquitin-like modifier activating enzyme 1; also known as *UBE1*) and *PCTK1* (PCTAIRE protein kinase 1), *EIF2S3* (eukaryotic translation initiation factor 2, subunit 3 gamma) and their flanking genes (Figure 1A and B). *UBA1* consists of two transcription start sites (TSSs) with the alternative non-coding first exons. Hereafter, the first TSS is numbered +1 and called *UBA1* TSS1, and the second, alternative TSS (at +3036) called *UBA1* TSS1'; accordingly, the first exons from TSS1 and TSS1' are called Ex1 and Ex1', respectively. Quantitative RT-PCR analysis showed that the transcripts from Ex1 were detected in HX, MRC5 and WI38, and those from Ex1' in all cells harboring human X including CF (Figure 1B, middle). The relative amount of *UBA1* Ex1' transcripts was roughly 2-fold in WI38 compared with MRC5, whereas that of Ex1 was comparable in the two. These results suggest that TSS1', but not TSS1, escaped inactivation in *UBA1* gene in WI38 and CF.

In the same locus, transcripts from 3' flanking genes to *UBA1* (i.e. *PCTK1* and *USP11*) were detected in all cell lines including CF harboring Xi (Figure 1B, middle), confirming previous data showing these genes escape from silencing as a cluster in other cell lines (7,34). In contrast, the expression of *RBM10*, located 4 kb upstream of *UBA1*, was detected only in the cells containing Xa (Figure 1B, middle), like other typical X-linked genes that undergo X-inactivation.

Another previously known escape gene *EIF2S3* was transcribed from both human Xa and Xi in hybrid cells (Figure 1B, bottom), as anticipated (7,35). Among the genes near *EIF2S3* (Figure 1A), *KLHL15* (Kelch-like 15) located 28 kb upstream *EIF2S3* was silenced in Xi (Figure 1B, bottom). In contrast, zinc finger protein, X-linked (*ZFX*), which is located 72 kb downstream *EIF2S3* and known to be an escape gene (7), was transcribed in all cells (Figure 1C). We did not obtain any evidence for the expression of *LOC347438/FAM48B1* (family with sequence similarity 48, member B1), a single exon gene located further 99 kb downstream, in any cell lines tested here (data not shown).

These results are entirely consistent with the previous data obtained by different hybrid cells (7,34), supporting the notion that clusters of genes spanning tens of kilobases escape X-inactivation constituting escape domains, and that chromatin boundaries between inactivated and escape regions should be present in the intergenic intervals.

#### Histone modification profile in *RBM10-UBA1* intergenic region

As the inactivated *RBM10* and escape gene *UBA1* are only ~4 kb apart, we focused on this region to investigate

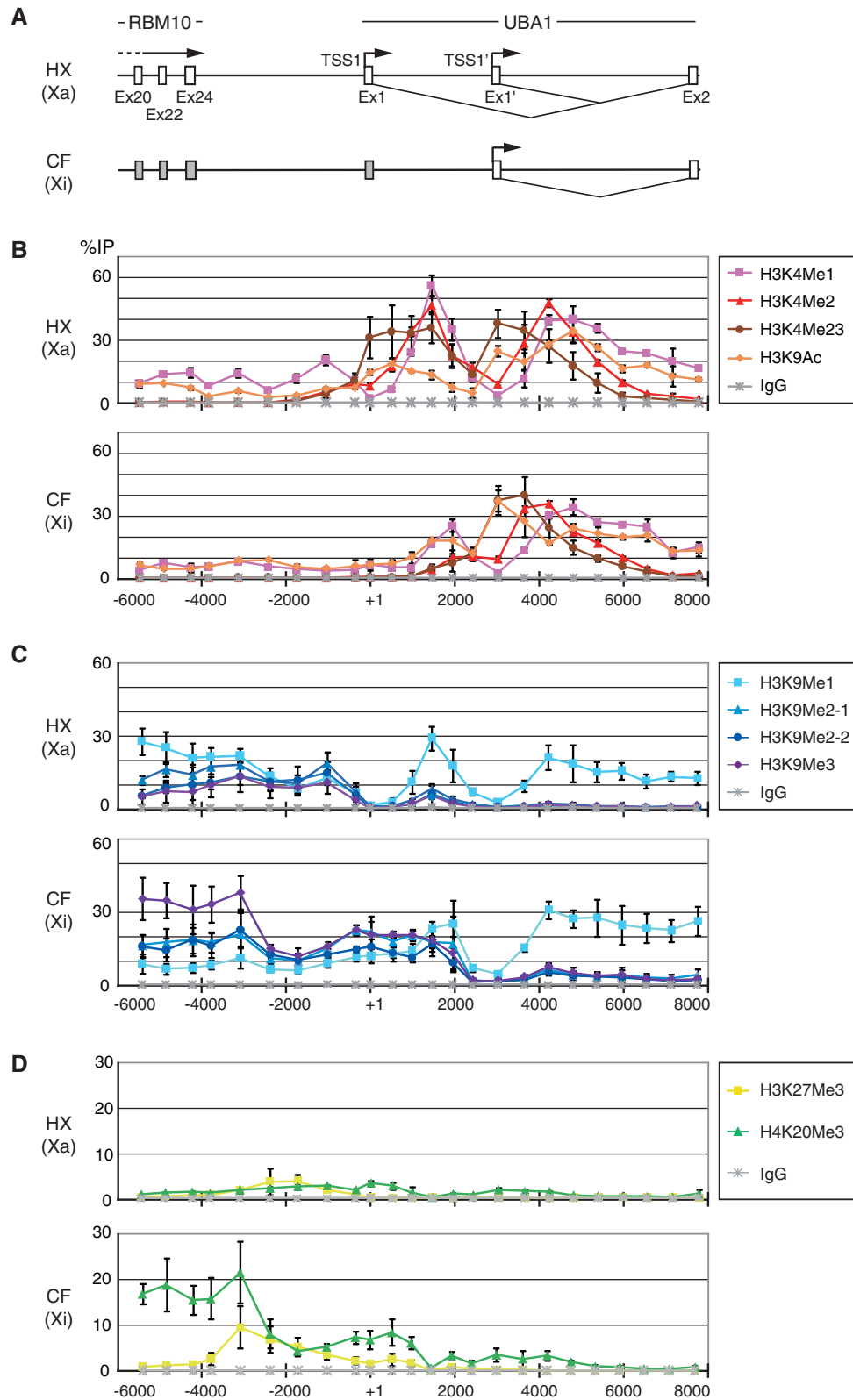
the boundary between silenced and active chromatin using hybrid cells. By using ChIP, we profiled histone modifications characteristic to euchromatin and heterochromatin along a 14 kb region covering the last exon of *RBM10* through the second exon of *UBA1* (Figure 2). Histone H3 K9 acetylation and K4 methylation are in general regarded as euchromatic modifications associated with the transcriptional activation (12,36–38). ChIP of di + tri-methyl H3K4 (H3K4me2 + 3) showed that these modifications were highly enriched around TSS1 and TSS1' on Xa (HX), and around TSS1' on Xi (CF) (Figure 2B), in good agreement with their transcription status (Figure 1B). As H3K4me2 distributed downstream H3K4me2 + 3, H3K4me3 appeared to be enriched more proximal to the TSSs. H3K4me1 was observed further downstream. H3K9ac was also enriched at active TSSs and downstream. Thus, all modifications characteristic to active chromatin were found at active TSSs and downstream but not in the intergenic region between the 3'-end of *RBM10* and *UBA1* TSS1. These results suggest that the active markers are simply correlated with transcription activities and may not be used as particular marks at the chromatin boundary in this locus.

H3K9me2 and H3K9me3 were reported as key modifications for heterochromatin formation and maintenance (12). Whereas H3K9me3 was observed little throughout the intergenic region of *RBM10-UBA1* locus on Xa, the same modification was highly enriched in inactivated *RBM10* last exons through the proximal downstream region on Xi (Figure 2C). The level of H3K9me3 was remarkably diminished at the intergenic region between *RBM10* and *UBA1* (~2 kb upstream *UBA1* TSS1) and disappeared at active TSS1' on Xi (Figure 2B). ChIP assay using two independent specific antibodies indicated that H3K9me2 showed similar distribution to H3K9me3. H3K9me1 was broadly detected from *RBM10* to *UBA1* except at around TSSs (Figure 2C). The level of monomethylation was relatively high in *RBM10* transcribed region on Xa. As H3K9me1 was also enriched in *UBA1* transcribed region on Xa and Xi, this modification seems to be correlated with active chromatin, in good agreement with recent genome-wide analysis (38).

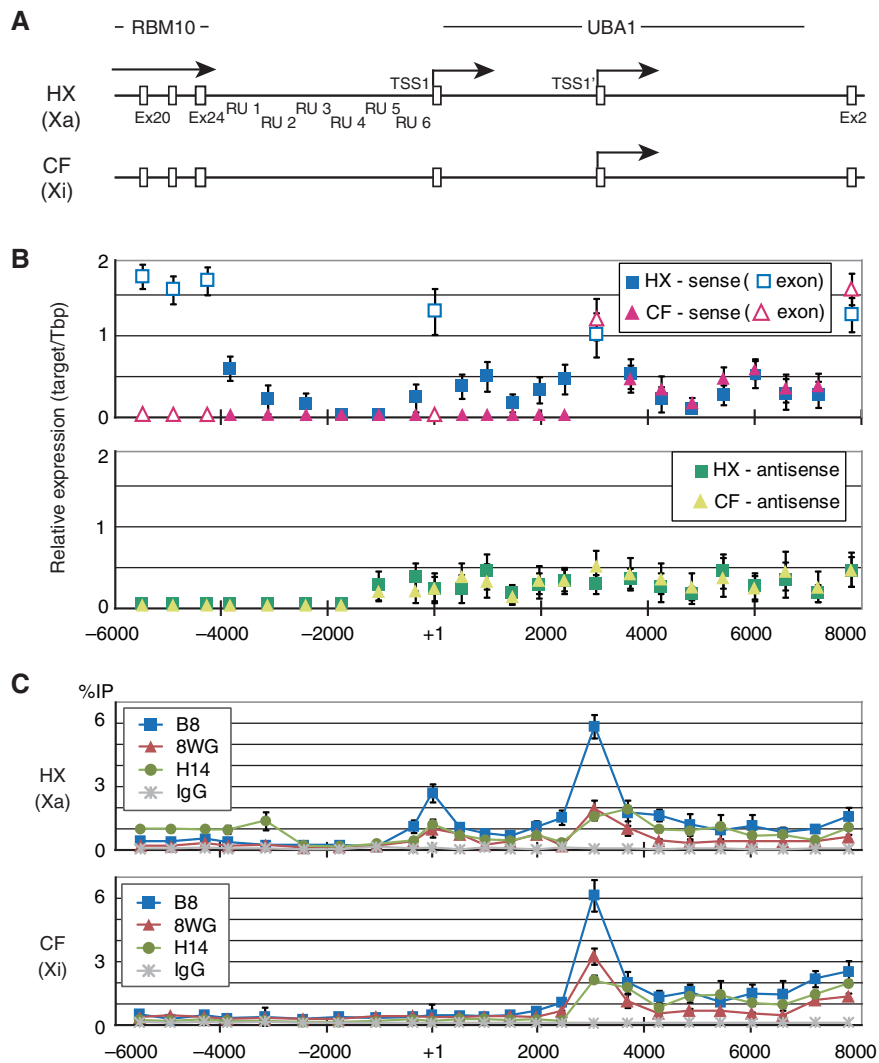
We further examined other modifications associated with repressed chromatin such as H3K27me3 and H4K20me3 (21,39). H4K20me3 was highly enriched in the inactivated *RBM10* to intergenic region on Xi similarly to H3K9me3 (Figure 2D). In contrast, enrichment of H3K27me3 was restricted to the transition point of H3K9me3 and H4K20me3 on Xi (Figure 2D). From these results, the heterochromatic feature of H3K9me3 and H4K20me3 enrichment in the intergenic proximal of *RBM10* appears to be disrupted at ~2 kb upstream of the *UBA1* TSS1, and a chromatin boundary is established at this region.

#### Little transcription around the *RBM10-UBA1* chromatin boundary

Non-coding transcripts are often found in locus control regions that regulate transcription of structural



**Figure 2.** Histone modification profiles in the ~12 kb region encompassing *RBM10* exon 20 to *UBA1* exon 2. ChIP was performed using antibodies against different modifications on histone H3K4, H3K9, H3K27 and H4K20. The enrichment of each region in immunoprecipitates was evaluated using real-time PCR and expressed as the percentage to the input (%IP). The mean values of three independent ChIP experiments are plotted with the standard deviations. (A) Schematic representation of the locus in HX (harboring Xa) and CF (harboring Xi). (B) ChIP with histone H3K4me1, me2, me2+3 and H3K9ac. (C) ChIP with histone H3K9me1, me2 and me3. (D) ChIP with histone H3K27me3 and H4K20me3.



**Figure 3.** The presence of transcripts and RNA polymerase II on *RBM10-UBA1* region. (A) Schematic representation of the locus in HX (harboring Xa) and CF (harboring Xi). (B) Relative amounts of sense and antisense transcripts measured by quantitative RT-PCR. (C) RNA polymerase II association. ChIP was performed using antibodies directed against Rpb8 (a shared subunit of RNA polymerase I, II and III; B8), unphosphorylated form of Pol II (8WG), and Ser5-phosphorylated form of Pol II (H14). The enrichment is expressed as the percentage to the input (%IP); the mean values from three independent experiments are plotted with the standard deviations.

genes (40–44) and siRNA could participate in establishing heterochromatin (45–48). Therefore, we investigated whether we could detect transcripts from the *RBM10-UBA1* intergenic region using strand-specific quantitative RT-PCR (Figure 3A and B).

*RBM10* mRNA was clearly detected along the last several exons by priming RT with the reverse primer in HX harboring Xa, but not in CF harboring Xi (Figure 3B). In *RBM10* proximal downstream regions (encompassing to the regions designated RU1 to RU3, from –3808 to –1595), sense transcripts primed with the reverse primers were also detected in HX (Figure 3B). These transcripts were likely to be read-through from *RBM10* transcription unit, as the exon sequences were amplified from cDNA primed with the reverse primers in the intergenic regions (RU1–3) (data not shown). Antisense transcripts in these *RBM10* proximal regions were, in contrast, undetectable in both cells under the

similar experimental conditions. Transcripts from either direction were not detected in RU4 region (from –1722 to –1595) in both tested cells (Figure 3B). Subtle antisense transcripts, primed with the forward primers, were observed in RU5 (from –1010 to –915) and RU6 (from –308 to –223) regions through *UBA1* gene on both Xa and Xi. These results suggest that *RBM10* read-through and *UBA1* antisense transcripts are terminated before approaching to RU4 region. We also detected weak signals of *UBA1* sense transcripts in RU6 region in cells harboring Xa (HX). In *UBA1* gene, the sense transcripts were clearly detected in the first exons (Ex1 and Ex1', starting from TSS1 and TSS1', respectively) and exon 2 (Ex2) in HX, whereas the transcripts were only detected in Ex1' and Ex2, but not in Ex1, in CF (Figure 3B), consistent with the ChIP data with methylated H3K4 and H3K9ac associating with active TSSs (Figure 2).

To refine these results, we also used ChIP assay to investigate the presence of RNA polymerases in *RBM10-UBA1* locus by using three monoclonal antibodies recognizing different forms (Figure 3C). B8 reacts with RPB8, a shared subunit of RNA polymerases I, II and III (33), and so chromatin fragments bound by any polymerase could be precipitated. 8WG16 and H14 recognize unphosphorylated and Ser5-phosphorylated forms of the C-terminal domain in the largest subunit of RNA polymerase II, respectively (49,50). While the pre-initiation complex on a promoter contains unphosphorylated form, Ser5 is phosphorylated during the initiation of transcription or when polymerases slow down at the posing and termination sites (51,52). ChIP with these antibodies revealed that RNA polymerase II was enriched around active *RBM10* TSSs (Figure 3C). Ser5-phosphorylated form (recognized by H14) was subtly detected from the last exons of *RBM10* to the proximal intergenic region (RU1 and in particular RU2, but not RU3), exclusively on Xa, which is probably associated with the termination of read-through transcripts. As polymerases were not detected in the intergenic region on Xi (Figure 3C), any specific transcripts, or transcription as such, does not appear to be crucial for the formation and maintenance of the chromatin boundary in this region.

### CpG hypomethylation and CTCF binding around the chromatin boundary on Xi

We next investigated DNA methylation at the *RBM10-UBA1* locus (Figure 4A). CpG dinucleotides on Xa were almost fully methylated from the last exon of *RBM10* through the intergenic regions (RU2 and RU5), and hypomethylated near TSS1 (RU6) and exon 1 of *UBA1*. CpG hypomethylation was also found in the 5' region of the first intron (In1-1) and around TSS1', but CpGs at the 3' region of the first intron (In1-4) were fully methylated. These results suggest that CpG hypomethylation is limited around active TSSs including promoter and enhancer elements on Xa. In contrast, while most CpG sites were fully methylated on Xi other than around TSS1', some CpGs in RU2 and In1-1 regions were remarkably protected from the methylation (Figure 4A). Such Xi-specific CpG hypomethylation might be involved in the chromatin boundary formation.

CTCF is known to be a methylation-sensitive DNA binding protein with chromatin insulator and enhancer blocking function (53-55). As CTCF also binds to 5' regions of escape genes like *Jarid1c* and *Eif2s3* (26), we investigated CTCF binding in *RBM10-UBA1* locus. By homology search with the previously reported CTCF binding motif CCGCNNGGNGGCAG (56), several potential CTCF binding sites were found in RU1, RU2, In1-1 and In1-4, close to the hypomethylated CpG sites (Figure 4A). We then examined the CTCF binding in these regions using cross-linked ChIP (Figure 4B). In both HX and CF cells, the positive-control loci, including mouse *H19* ICR [imprinting control region; (57)] and human *EIF2S3* promoter (26), were enriched

in anti-CTCF immunoprecipitates (data not shown). CTCF was distributed in the entire *RBM10-UBA1* intergenic region on Xi, with peaks at RU2-RU3 and In1-1-In1-2, in contrast to Xa, where little CTCF binding is observed (Figure 4B). Thus, the CpG hypomethylation and CTCF binding around RU2-RU3 region appear to be correlated with the chromatin boundary formation.

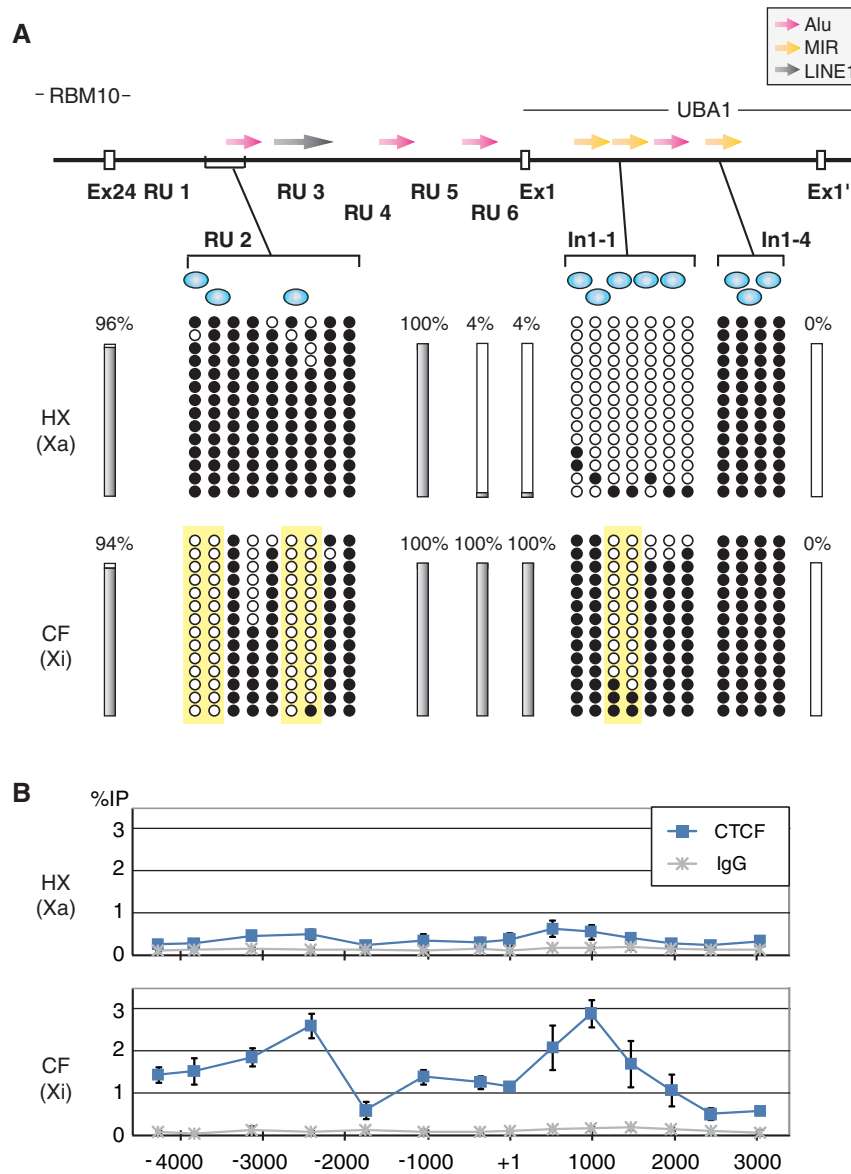
### DISCUSSION

A number of genes on human Xi chromosome are expressed, or escaped from the silencing, even after the whole chromosome is inactivated in somatic cells. To distinguish the Xa and Xi alleles, F1 hybrid mice from different parent strains are often used (26); however, the analysis is restricted to the regions containing small nucleotide polymorphisms on mouse X chromosome. Inter-species hybrid cell lines containing only either active or inactive human X chromosome allow higher resolution analysis without allelic examination (7,26). We here used human-mouse hybrid cells to investigate the chromatin boundary between inactivated *RBM10* and escape *UBA1* genes on human X chromosome. As summarized in Figure 5, the TSS1 was silenced on Xi, whereas the TSS1 and the downstream alternative transcription start site (TSS1') were both used on Xa. By detailed profiling of the histone modifications and transcription status using ChIP and real-time PCR at ~500 bp intervals across the intergenic region, a boundary between inactive and active chromatin was found around the middle of the intergenic region (~2 kb upstream the *UBA1* TSS1) on Xi. Importantly, similar histone modification profiles were observed in diploid human fibroblasts. In male cells harboring only Xa (MRC5), the distributions of H3K4me1-3 and H3K9me1-3 were similar to those in HX (Supplementary Figure S1). In female cells harboring Xa and Xi (WI38), their distributions appeared as the mixture of HX and CF (Supplementary Figure S1). These results support the idea that the epigenetic status of X chromosomes that was established in human cells is maintained in the hybrid cells, as seen by others (7,26). We also found CTCF binding and hypomethylated CpG in this chromatin boundary region specifically on Xi; in contrast, H3K9 acetylation and RNA polymerases were hardly detected (Figure 5).

### Histone modifications around the *RBM10-UBA1* boundary on Xi

In the *RBM10-UBA1* locus on Xi, H3K9me3 and H4K20me3 were enriched in *RBM10* gene body and the proximal downstream region but diminished at the middle in the intergenic region. H3K27me3, a modification enriched in facultative heterochromatin, was distinctively detected around this boundary region. These data are consistent with the mosaic structure of different levels of heterochromatin found on human Xi (58), and possibly correlated with the organization of chromatin boundary formation and maintenance of escape genes.





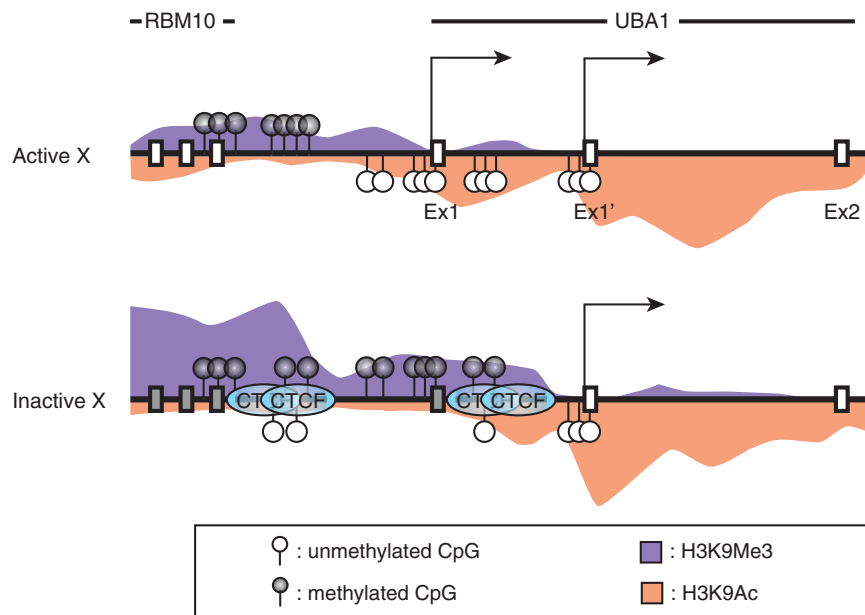
**Figure 4.** DNA methylation status and CTCF binding at the chromatin boundary and flanking region. (A) The methylation status of the regional and individual CpG dinucleotides in *RBM10-UBA1* region. Top: The gene structure is schematically represented with the positions of repeat sequences (Alu, red; MIR, yellow; and LINE1, grey) and putative CTCF binding sites (blue circle). Middle and bottom: DNA methylation status in HX and CF cells. Solid bars represent the methylation status of regions where no particular site exhibits a distinct pattern among several CpG sites. The percentages of methylation levels are indicated. Small circles represent the individual methylation status; methylated and unmethylated CpG sites are shown as filled and open circles, respectively. The inactive X-specific hypomethylated area is highlighted by yellow background. (B) ChIP with anti-CTCF. The enrichment is expressed as the percentage to the input (%IP); the mean values from three independent experiments are plotted with the standard deviations.

In contrast to H3K9me3 and H3K9me2, H3K9me1 was broadly distributed throughout the transcribed region other than around active *UBA1* TSSs, where H3K9 was heavily acetylated, suggesting that H3K9me1 is distributed broadly in relatively opened chromatin (38). All active TSSs also showed the prominent peak of H3K4me3, associated with those of H3K4me2 and then H3K4me1 shifted to downstream. These data are consistent with the previous genome wide and locus specific data (36,37,59). The monomethylated form may be a neutral state for H3K4 in open chromatin, because

unmethylated and higher methylated forms may recruit DNA methyltransferase 3L (60) and transcription machineries (61).

#### Possible mechanisms of chromatin boundary formation in *RBM10-UBA1* locus

CTCF-dependent insulators have been characterized in  $\beta$ -globin and imprinted *IGF2/H19* loci (53,54,57,62). Non-coding transcripts and hyper-acetylation of H3/H4 were often associated with these insulators (63,64),



**Figure 5.** Schematic representation of *RBM10-UBA1* region. The enrichment of histone modifications (H3K9Ac and H3K9me3), CpG methylation status and CTCF binding sites are shown.

suggesting the involvement of transcription in the insulator function. On mouse and human Xi, CTCF binding has also been reported at the 5' end of escape genes, *Jarid1c*, *Eif2s3x* and *EIF2S3*, adjacent to an inactivated gene (26). Although the boundary between *RBM10* and *UBA1* seems similar to these insulators in terms of CTCF binding, there are two notable distinct features. First, it does not appear to be a typical LCR/ICR-type insulator, as little RNA polymerase and acetylated H3 were associated around the boundary. Second, the insulator function may not be very strong, because the transcription from TSS1 was repressed on Xi. The presence of two TSSs (i.e. tandem promoter/enhancer elements) may also contribute to maintain the *UBA1* transcription activity by suppressing heterochromatin spreading from the upstream inactive region. In fact, chicken  $\beta$ -globin insulator, containing DNaseI-hypersensitive site 4, was not sufficient to intercept the X-inactivation, when GFP reporter gene flanked with the insulator was introduced into X-linked *Hprt* locus (65). Thus, CTCF-mediated insulators per se may not be sufficient to maintain the transcription activity on Xi due to the global silencing of the whole chromosome. Alternatively, protein complexes assembled on strong promoter/enhancer may also recruit CTCF to block the heterochromatin spreading.

The correlation between CTCF binding and DNA methylation has well been characterized in the insulators of the imprinted genes (54,57) as well as in the enhancer of *Tsix* (56). In the *RBM10-UBA1* intergenic region, the site-specific CpG hypomethylation was also detected exclusively on Xi in the methylated domains on Xa. From the ChIP data, CTCF binding does not appear to be restricted to this hypomethylated region, but the binding was rather spread throughout the intergenic region. It is thus suggested that the CTCF-mediated insulator activity of *UBA1* boundaries may be regulated by the specific CpG

methylation, but the insulator machinery may progressively spread from the DNA methylation-free cardinal point to their vicinities. In good agreement with this notion, among seven CTCF binding sites in human H19 differential methylation region (DMR), only the sixth site is reported to act as a key regulator for switching between *IGF2* and *H19* expression, whereas the other sites can be hyper-methylated (66).

Consistent with previous reports (7,34), we here showed that *PCTK1* and *USP11*, located 3 and 6 kb downstream *UBA1*, respectively, were also escaped in CF cells. As many escape genes are clustered (7), it is likely that the escaping from the inactivation is regulated at chromatin domain levels. Recent studies also suggest that the chromatin insulator activity is correlated with the formation of long-distance chromatin loops by CTCF and cohesin (67,68). During and after X inactivation, *Jarid1c* gene remains located outside the Xist-RNA accumulated, transcriptionally silent compartment (69), suggesting that the inactivated and escape domains can be separated into different chromatin loops anchored through CTCF and cohesin. If this is the case, the *UBA1-PCTK1-USP11* escape domain may be extruded from the inactivated X chromosome body. The 3' boundary of this escape domain should be laid in the intergenic region between *USP11* and *ZNF157*, which is located ~122 kb downstream, as *ZNF157* and further downstream genes are silenced [Figure 1B and data not shown; (7)]. The large intergenic sequence between *USP11* and *ZNF157*, however, presented a great difficulty to exploring the 3' boundary of *UBA1* escape domain.

#### Repetitive sequences in human *RBM10-UBA1* intergenic region

In human, more than 15% of the X-linked genes escape the inactivation, and escape genes are spread throughout

X chromosome with enrichments in PARs (7). Recent bioinformatic analyses revealed that the primary DNA sequence is correlated with the chromatin status on human Xi; *Alu* repetitive elements and CAG/CGT and GATA repeats are significantly enriched in the escape domain, whereas the long interspersed nuclear element (LINE) and mammalian-wide interspersed repeat (MIR) preferentially distributed in inactivated domains (70,71). The *UBA1* boundary region indeed contains an *Alu* variant, in which the hypomethylated CpG sites were mapped. As mouse *Ubal* is not an escape gene (72), the insertion of *Alu* element between *RBM10* and *UBA1* may contribute to the establishment of the escape domain in human. Although the downstream boundary region is not identified, fragmented *Alu* repeat-containing sequences account for ~45% of the entire 122 kb intergenic region between *USP11* and *ZNF157*, supporting the notion that *Alu* sequence may disturb the heterochromatin spreading on Xi. Further identification of the boundaries between the inactivated and escape domains may draw more clarified view how those domains are established and maintained.

## SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online.

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