Identification of the gene coding for the Endo B murine cytokeratin and its methylated, stable inactive state in mouse nonepithelial cells

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The Endo B type-I keratin intermediate filament protein is first expressed at the 4- to 8-cell stage of mouse development. In the adult, its expression is restricted to a variety of simple epithelial cell types. To investigate the mechanisms responsible for the restricted expression of Endo B, the gene coding for Endo B has been identified from among the five different Endo B genes found in the mouse genome by Southern hybridization analysis and cloning all or part of four of the genes. Nuclear run-on experiments demonstrate that Endo B expression is regulated at the level of transcription. The 5' end of the active gene, designated Endo B1, was found to be highly methylated and in a relatively nuclease-resistant chromatin conformation in fibroblasts and myoblasts that do not express Endo B, but undermethylated and relatively sensitive to nuclease digestion in endodermal cells or F9 embryonal carcinoma cells. The inactive state of the Endo B B1 gene in fibroblast appears to be very stable, because somatic cell hybrids formed by the fusion of HeLa cells, which express the homologous human protein, keratin 18, and mouse fibroblasts, continue to express keratin 18 but do not activate Endo B expression. Similarly, the fusion of mouse endodermal cells and fibroblasts results in hybrids that do not extinguish Endo B expression. These results suggest that Endo B transcription is limited by two different mechanisms. In somatic cells such as fibroblasts or myoblasts, expression may be restricted by methylation and a stable, nonpermissive transcriptional state. However, in embryonal carcinoma cells, the Endo B B1 gene is undermethylated and in a relatively nuclease-sensitive conformation, but it is restricted by an additional, negative regulatory mechanism.

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Endo B [Oshima 1981] (also referred to as cytokeratin D, Franke et al. 1981a,b) and its homologous human form, keratin 18 (K18) are type-I keratin intermediate filament proteins (for reviews on intermediate filament proteins, see Lazarides 1980, 1982; Steinert and Parry 1985) found in a variety of simple epithelial tissues (Kemler et al. 1981; Moll et al. 1982; Schiller et al. 1982). Endo B and Endo A, its complementary type-II keratin subunit [Brulet et al. 1980; Oshima 1982], are first expressed at the 4-8 cell stage of mouse embryogenesis but are limited to trophoblastic and extraembryonic endodermal tissues of the blastocyst stage embryo [Brulet et al. 1980; Oshima et al. 1983]. Murine embryonal carcinoma (EC) cell lines provide a convenient system for investigating early development. The in vitro differentiation of EC cell lines provides a convenient system for investigating early developmental cellular transitions that would be difficult otherwise.

Two aspects of the regulation of Endo B expression are of particular interest. The first is the mechanism responsible for the expression of the same gene in a restricted but relatively large number of differentiated cell types as diverse as parietal endoderm and liver [Trevor and Oshima 1985]. In this respect, the regulation of Endo B might be considered intermediate between specialized genes expressed in only a very few cell types such as β-globins, immunoglobulins, α-fetoprotein, or epidermal keratins and ubiquitously expressed genes such as those for many metabolic functions. The second aspect is its activation and expression at very early times in development. Endo A and B are among the first gene products differentially expressed during mouse development [Chisholm and Houliston 1987].

In this study we identify the single Endo B gene among the five found in the mouse genome that is likely responsible for expression in diverse permissive cell types and show that it is regulated at the transcriptional level. Furthermore, we provide evidence that in differentiated cells that do not express Endo B, the gene is in a stable, inactive state characterized by a highly methylated and relatively nuclease-resistant chromatin structure.

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Results and discussion

Endo B genes in rodents and primates

The tissue distribution of Endo B and its homologous human form, K18, are very similar, if not identical (Moll et al. 1982; Schiller et al. 1982) in the adult. However, the expression of K18 and K8 in human EC cells (Damjanov and Andrews 1983; Damjanov et al. 1985; R.G. Oshima, unpubl.) suggests that human EC cells and perhaps human embryonic cells differ from mouse EC cells and their respective mouse embryonic counterparts in this respect. Previously, we have estimated that the mouse and human genomes contain ~5 and 20 copies, respectively, of genes very similar to Endo B (Trevor and Oshima 1985). The larger number of genes similar to Endo B found in human DNA may reflect a difference in early developmental expression, because expression of K18 in the equivalent of the mouse inner cell mass could provide a target for retrovirus-mediated pseudogene generation in the germ line (Linial 1987). To determine whether the large number of human genes similar to Endo B is characteristic of primates in general, a number of mammalian DNAs was screened by the method of Southern (1975) for sequences similar to Endo B under conditions that do not permit detection of related epidermal keratin or other intermediate filament genes. Figure 1 shows that mouse, rat, horse, and whale DNAs have a relatively small number of fragments that hybridize strongly to the Endo B cDNA, whereas all primates, with the possible exception of lemur, have a much larger number of similar sequences. These animals are representatives of evolutionarily diverse primate lineages, including prosimians (potto), old world monkeys (macaque), new world monkeys (saki and spider monkeys), great apes (orangutan, chimpanzee, and gorilla), and humans. This suggests that the duplication of Endo B-like genes occurred early in primate evolu-

Figure 1. Primates have many genes closely related to Endo B. DNAs from the indicated species were digested with BamHI, separated by agarose gel electrophoresis, transferred to nitrocellulose, and hybridized to nick-translated Endo B cDNA. Final washes were in 0.1 x SSC, 0.1% SDS at 50°C, which eliminates detectable hybridization to related epidermal keratin sequences. [1] Mouse; [2] rat; [3] horse; [4] sperm whale; [5] lemur; [6] potto; [7] saki monkey; [8] spider monkey; [9] macaque; [10] orangutan; [11] chimpanzee; [12] gorilla; [13] human. Size markers are indicated on the left of each panel (kb).

Figure 2. Comparison of cloned genes to mouse endogenous Endo B genes. [A] Multiple aliquots of 10 μg of F9 mouse genomic DNA were digested with EcoRI, separated in an agarose gel, transferred to nitrocellulose, and hybridized to a sense RNA probe representing the first 461 bp of the Endo B cDNA. Strips of the filter were hybridized together but washed independently in 0.1 x SSC, 0.1% SDS, at the temperature indicated at the top of the panel. The individual Endo B genes are indicated at the left of the panel. An autoradiographic image obtained with an intensifier screen is shown. Note that after washing at 75°C (lane 4), only the signal of the β1 gene is still detectable. [B] Phage λ DNAs containing cloned Endo B genes [EB31, lane 1; EBp3, lane 2, EB1, lane 5], plasmid DNA containing the 5’ end of the Endo B β1 gene [SK3112, lane 4], and mouse genomic DNA isolated from a near diploid parietal endodermal cell line [HR9, lane 3] were digested with EcoRI, separated in an agarose mini gel, transferred to nitrocellulose, hybridized to the same probe used in panel A, and washed in 0.1 x SSPE, 0.1% SDS at 50°C. The probe contains all of exon I and only nine nucleotides of exon II sequences of the β1 gene. An autoradiographic image obtained with an intensifier screen is shown. Carrier undigested Escherichia coli DNA (2 μg) was added to ~60 pg of phage DNA or 15 pg of plasmid DNA before loading. Two micrograms of mouse genomic DNA were used. The Endo B genes are indicated on the left, and size markers (in kb) are indicated on the right. The variation in the rate of migration of the hybridizing band of EBp3 DNA (lane 2) and the endogenous β5 Endo B gene (lane 3) is probably due to effects of the undigested carrier DNA.
The presence of processed pseudogenes for Endo A (Vasseur et al. 1985) and Endo B (see below) and the evidence for expression of only a single Endo B gene (Trevor and Oshima 1985) in both mice and humans (Kulesh and Oshima 1988) suggest that many of the hybridizing bands found in primate DNA may represent pseudogenes.

To identify which of the mouse Endo B genes might code for the expressed protein, Southern analysis was performed under increasing stringency. The results are shown in Figure 2B. After EcoRI-digested mouse DNA is hybridized with an RNA probe of the first 461 nucleotides of the Endo B cDNA and washed at 60°C, five fragments are detected [lane 1], which are designated β1–β5. If replicate filters are washed at 65°C or 70°C, only three of the fragments are detectable [lanes 2 and 3]. At 75°C, only a single hybridizing fragment is detected. This fragment, designated β1, likely contains sequences most similar to the cDNA probe and suggests that it represents a part of the gene transcriptionally active in endodermal or liver cells. The same experiment performed on BamHI-digested mouse DNA resulted in a single detectable fragment of ~6.5-kb after washing at 70°C [data not shown].

Figure 2B compares the hybridization pattern of the endogenous Endo B genes to four genes that were isolated by cloning. Repeated screening of total genomic DNA libraries resulted in the isolation of the β2, β3, and β5 genes. However, the β1 gene, which was the best candidate for the active gene, was apparently underrepresented in these libraries (Table 1). Part of the β1 gene was isolated by screening a library prepared from size-selected DNA derived from a cell line that expressed Endo B. This resulted in the isolation of the 5′ end of the β1 gene. Comparison of restriction enzyme maps of the β2, β3, and β5 genes to that of the cDNA and Southern analysis using probes of the cDNA specific for the 5′, middle, or 3′ portions of the cDNA revealed that none appeared to code for the Endo B cDNA (Fig. 3). The sizes of different hybridizing portions of the genes suggested that each might represent a pseudogene derived from reverse transcription of the Endo B RNA. Partial sequence analysis of two portions of the β2 EB3 isolate confirmed this suggestion for the β2 gene [data not shown]. In contrast, the results of Southern analysis under very stringent conditions [Fig. 2A], DNA sequencing of the β1 gene, and analyses of DNase-digested nuclei and the methylation state of the Endo B genes [see below] suggest that the 3-kb EcoRI fragment of the β1 gene represents part of the gene expressed in endodermal cells.

### Table 1. Summary of the isolated Endo B genes

| Gene | Isolate | Strain | Tissue | Vector | Library type | Plaques | Number |
|------|---------|--------|--------|--------|-------------|---------|--------|
| β1   | SK3112  | 129 Sv/J | HR9 cells | AZAP   | EcoRI partial | 4 \times 10^6 | 5      |
| β2   | EB3     | BALB/c  | embryo | Charon 28 | MboI partial | 3 \times 10^6 | 20     |
|      | EB25    | C57BL/6 | T cell  | EMBL3  | MboI partial | 5 \times 10^6 | 2      |
|      | EBp11   | BALB/c  | embryo | Charon 4A | EcoRI partial | 6 \times 10^6 | 1      |
| β3   | EB1     | 129     | liver   | EMBL3A | MboI partial | 6 \times 10^6 | 11     |
| β5   | EBp3    | BALB/c  | embryo | Charon 4A | EcoRI partial | 6 \times 10^6 | 2      |

**Sequence of the 5′ end of Endo B β1**

The sequence of the 5′ end of the Endo B β1 gene is shown in Figure 4. A 452-nucleotide portion of the gene was found to match the 5′ end of the Endo B cDNA exactly, including the 5′-noncoding leader. In addition, the 3′ end of this region [nucleotide 451, AGGTAAGG] matches the sequence expected for a 5′ donor splice junction (Ohshima and Gotoh 1987). Thus, the underlined portion of the sequence represents the first exon of the Endo B β1 gene, and the sequence following it represents part of the first intron. The first exon contains all but 9 nucleotides of the 461-nucleotide coding portion of the Endo B RNA probe used for screening and Southern hybridization.

Nuclease S1 protection analysis utilizing a fragment of the gene overlapping the expected 5′ end of the Endo B mRNA mapped the major transcriptional start site designated nucleotide +1 [Kulesh and Oshima 1988]. The TATA box motif [ATATAA], found immediately upstream of the transcriptional start site of many eukaryotic genes, is found at nucleotide −27. In comparison with the K18 gene [Kulesh and Oshima 1988 and unpubl.], eight regions of 8–11 identical nucleotides, but none longer than 11 nucleotides, are found. Some of these upstream regions may represent functionally important sequences because the TATA box is among them. No core sequences for the potential binding of the Sp1 transcription factor [GGGCGG, CCGCCC] [Dynan and Tjian 1985] are found, even though the K18 gene contains four such sites within 200 bp upstream of the human gene [Kulesh and Oshima 1988]. Two regions of short repeated sequences [nucleotides −816 and −448] and one imperfect long direct repeat [nucleotide −402] are found. One region within the first exon, starting at nucleotide 176, may be of particular importance. Of 18 nucleotides of this region, 17 are identical to the tRNA-binding site of murine retroviruses that has been implicated in the block of viral RNA expression in murine EC cells [Barklis et al. 1986; Loh et al. 1987; Weiher et al. 1987].

Two portions of the sequence from nucleotide −129 to +1 and from nucleotide 121 to 251 are particularly GC rich [68% and 71%, respectively]. In the first region, the dinucleotide CG is used almost as much as GC (10 to 12 times, respectively). The frequency of the CGs relative to GCs in the second region [58%, 10 CGs, 17 GCs] is similar to that of the sequence of the entire fragment [60%, 56 CGs, 93 GCs]. The 129 nucleotide region immediately upstream of the transcriptional start site of
Figure 3. Restriction maps of the isolated Endo B genes. Left and right cross-hatched and stippled regions of the Endo B cDNA and genomic fragments indicate the portions used as probes and their respective hybridizing fragments. Only hybridizing portions of the genomic fragments of β2 EB31, β3 EB1, and β5 EBp3 are shown. All three also contain additional unmapped flanking sequences. Enzymes shown are BamHI (B), EcoRI (E), PstI (P), EcoRV (RV), SstI (SI), and SstII (SII). The broken portion of the fragment of β2 EB31 indicates the approximate portion of the 7-kb fragment presumed to be responsible for the hybridization with the indicated probe and fragment. The broken EcoRI fragment of β3 EB1 indicates that this fragment hybridized to both the 5' and middle portion cDNA probes. The arrow beneath the map of β2 EB31 indicates the direction and approximate number of bases sequenced to confirm the pseudogene structure. The scale is indicated over the left portion of each map (bp).

the Endo B β1 gene appears to qualify as an ‘HTF island’ (Bird 1986). Such regions are usually unique sequences of genomic DNA associated with the 5’ end of constitutively active genes and additionally are undermethylated.

Endo B is transcriptionally regulated

Endo B is expressed in various simple epithelial cell types of the adult, parietal endodermal cells, and in EC cells that have differentiated to extraembryonic endodermal derivatives. Nuclear run-on assays were performed to determine whether the differential expression of Endo B is due to transcriptional or post-transcriptional mechanisms. Figure 5 shows the results of hybridizing radioactive RNA made by isolated nuclei from several cell types to various target DNAs. This assay measures the transcriptional activity of endogenous genes by permitting elongation of previously initiated transcriptional complexes in the absence of new initiation (McKnight and Palmiter 1979). Positive signals for the transcription of the Endo B gene were found only in nuclei from F9 cells induced to differentiate by exposure to retinoic acid and in HR9 parietal endodermal cells but not in 984 myoblasts, undifferentiated F9 cells, or STO fibroblasts (data not shown). Actin transcription served as a positive control for all cell lines. The differentiation of F9 cells also results in increased transcription of Endo A, the type-II keratin with which Endo B polymerizes. In contrast, the large increase in proteoglycan PG19 RNA in retinoic acid-treated F9 cells (Grover et al. 1988) appears to be regulated post-transcriptionally.

The Endo B β1 gene is differentially sensitive to DNase digestion

The chromatin states of the Endo B genes were investigated by digesting nuclei from various cell lines with varying concentrations of DNase I and analyzing the subsequently purified DNA by Southern analysis. Figure 6 shows the results of such an experiment for nuclei isolated from the MB4 endodermal and 984 myoblast cell lines. The Endo B β1 gene in endodermal cells is much more sensitive to digestion than the β2, β3, or β5 genes (Fig. 6, lanes 1–6). Weak hybridization precluded a firm conclusion concerning the sensitivity of the β4 presumptive gene. The unlabeled, large fragment detected above the β5 gene, which appears to be sensitive in both MB4 and 984 nuclei, does not hybridize to probes specific for the 5’ end of the Endo B cDNA (see Figs. 2 and 7) and may represent the 3’ end of either the β1 or β2 genes. In contrast to the case in endodermal nuclei, the β1 gene in 984 myoblast nuclei is relatively resistant to digestion. The β1 gene was detectable even after diges-
Figure 4. DNA sequence of the 5' end of the Endo B (31 gene. The sequence starts with BamHI site at -928 and ends at the EcoRI site at nucleotide 648. Nucleotide numbers on the left are relative to the start of the Endo B mRNA as +1. Solid underlined sequence represents exon I, which is 452 nucleotides. The ATG translational start codon and a potential TATA box transcriptional regulatory element are indicated by asterisks (*). Direct repeated sequences are indicated by dashed lines with arrow heads (-->). Plus signs (+) indicate the longest stretches of nucleotides identical in the homologous human K18 gene 5'-flanking region. The nucleotides of the first exon that are identical to the murine retroviral tRNA-binding site are indicated by arrowheads (>).

Differential methylation of the β1 gene

The methylation state of Endo B genes in the DNA of several cell types was investigated by Southern analysis utilizing the methylation-sensitive restriction enzyme HpaII and its methylation-insensitive isoschizomer MspI. After digestion with EcoRI, DNAs from HR9 endodermal cells, F9 EC, STO fibroblasts, and 984 myoblasts were treated with either MspI or HpaII and analyzed by blot hybridization. The results are shown in Figure 7. In both HR9 DNA and undifferentiated F9 DNA, the signal for the 5' end of the β1 gene is completely lost upon digestion with HpaII. Within the portion of the β1 gene that has been sequenced, there are 10 MspI/HpaII sites, six of which are located in the first exon. It is not possible by these results to ascertain directly the methylation status of the two individual MspI/HpaII sites located immediately upstream of the first exon (nucleotides −84 and −78) in the region of particularly high GC content. However, complete methylation of the four sites upstream of the transcriptional start site of the β1 gene should have resulted in a detect-
Figure 5. Nuclear run-on analysis of Endo B expression. Nuclei were isolated from F9 EC cells (F9), F9 cells that had been induced to differentiate by exposure to 1 μM retinoic acid for 4 days (F9 + RA), HR9 parietal endodermal cells (HR9), or 984 myoblasts (984) and permitted to elongate previously initiated RNA transcripts in the presence of [32P]UTP. The radioactive RNAs were purified and hybridized to 5 μg of the denatured DNAs indicated at the left of the figure, which had been immobilized on nitrocellulose filters. After 4 days of hybridization, RNase treatment, and washing, the filters were exposed to film in the presence of an intensifier screen. (Endo B+) M13 single-stranded DNA containing the noncoding strand of the Endo B cDNA from nucleotide 461 to 1456, which will hybridize to Endo B mRNA; (Endo B-) M13 single-stranded DNA containing the coding strand of the same portion of the Endo B cDNA as Endo B+ (M13mp10) control M13mp10 single-stranded DNA containing no additional sequences; (Actin) human β-actin cDNA plasmid pHFpA-1 (Ponte et al. 1984); (Endo A) a-1 Endo A pseudogene plasmid (Vasseur et al. 1985); (Neo) control Sp 65 plasmid containing a portion of the Tn5 neo gene; (PG) rat proteoglycan plasmid PG-6 (Bourdon et al. 1987) plasmid.

Expression of Endo B and K18 in cellular hybrids

To test for the presence of possible trans-acting negative regulatory activities of Endo B, mouse fibroblasts were fused with HeLa cells, which express the homologous K18 gene, and hybrids were isolated for protein and DNA analysis. Long-term hybrids instead of hetero-

Figure 6. DNase sensitivity of Endo B genes in nuclei of endodermal and myoblast cells. Nuclei from MB4 endodermal cells (lanes 1–6) or T984cl10 myoblasts (lanes 7–12) were suspended at a concentration that resulted in an A260 of 20 in 1 M NaOH and digested with the indicated concentrations of DNase I (bottom) for 10 min at 37°C. The reaction was stopped by the addition of EDTA, SDS, and proteinase K. After organic solvent extraction and digestion with EcoRI, the samples were separated in a 0.5% agarose gel, transferred to nitrocellulose, and hybridized to nick-translated Endo B cDNA. The Endo B genes are labeled on the left, and size markers are indicated on the right (in kb). (Band c) A small amount of control plasmid DNA added after digestion with DNase that permits evaluation of the loading and transfer efficiency. Lanes 13 and 14 contained samples 1 and 7, respectively, with 20 pg of additional Endo B cDNA to monitor the restriction enzyme digestion.
Figure 7. Differential methylation of the Endo B β1 gene. DNAs isolated from mouse HR9 parietal endodermal cells [lanes 1–3], F9 EC cells [lanes 4–6], STO fibroblasts [lanes 7 and 8], or 984 myoblasts [lane 9] were digested with EcoRI alone [lanes 1 and 4], or in combination with MspI [lanes 2, 5, and 7] or HpaII [lanes 3, 6, 8, and 9]. The digests were separated by agarose gel electrophoresis, transferred to nitrocellulose, and hybridized to an RNA probe corresponding to the first 461 nucleotides of the Endo B cDNA. Final washes were at 50°C. Individual Endo B genes are indicated on the left, and size markers are indicated on the right (in kb). Note that the β3, β4, and β5 genes are resistant to digestion with HpaII in all cell lines. However, the β1 gene is sensitive to digestion with HpaII only in HR9 and F9 cells.

Karyon analyses were necessary for this study because of the extremely slow turnover of Endo B intermediate filaments and the difficulty in distinguishing Endo B and K18 by immunofluorescent methods. Figure 8A shows the result of immunoprecipitation analysis of a subset of all the hybrids analyzed. Three of the four hybrids shown continue to express K18, and none activates Endo B. The lack of expression of Endo B was not due to the absence of the Endo B β1 gene in these hybrids, because Southern analysis confirmed the presence of the gene in these cells (6.5-kb band, Fig. 8B, lane 12; additional data not shown). The varying level of expression of K18 correlates well with the presence of the active K18 gene sequences found in the hybrids, as shown in Figure 8B. The characterization of the gene coding for K18 [Kulesh and Oshima 1988] permitted the identification of a diagnostic 1.7-kb BamHI fragment containing sequences coding for the central portion of K18. This fragment was not detected in hybrid HeST 7.1 [Fig. 8B, lane 6], was barely detectable in HeST 3.7 [lane 5], and was present in higher amounts in HeST 7.5 [lane 9] and HeST 9.6 [lane 10]. Thus, there is a direct relationship between the presence of the coding K18 gene and K18 protein synthesis. Further analysis of the degree of K18 protein synthesis in the other HeST hybrids investigated in Figure 8B confirmed this relationship [data not shown]. The results provide no evidence for trans-acting negative regulators of the Endo B gene in mouse fibroblasts that could silence the homologous K18 gene. In addition, the silent Endo B gene in fibroblasts is not expressed even in a cellular environment that supports the expression of the homologous human gene.

To test whether the results of the analysis of the HeST hybrids were due to species-specific effects, hybrids were constructed between two mouse cell lines, which differed in their expression of Endo B. The isolation of authentic hybrids between two cells of the same species depends critically upon a double selective system that excludes the growth of both parents and the isolation of hybrids at sufficiently high frequency to make improbable the isolation of rare parental variants resistant to the selective conditions. Proliferating hybrids of HR9 endodermal cells and STO fibroblasts were found at a frequency of ~1 x 10^-4, whereas the frequency of variant parental cells was ~1 x 10^-6. All four hybrids isolated from the fusion of HR9 and STO cells and examined in detail continued to express Endo B protein [Fig. 8C]. These results reinforce the conclusion that STO fibroblasts do not appear to contain trans-acting regulatory activities that can suppress Endo B expression.

Conclusions

Endo B and its complementary type-II keratin, Endo A, which are first expressed at the 4–8 cell stage of development (Oshima et al. 1983; Duprey et al. 1985; Chisholm and Houliston 1987) are the first intermediate filament proteins to be expressed during mouse embryogenesis [Brulet et al. 1980; Jackson et al. 1980; Paulin et al. 1980; Oshima et al. 1983]. This appears to be the only time when Endo B is expressed in the mouse germ line. By the early blastocyst stage, these proteins are expressed almost exclusively in the trophectoderm [Chisholm and Houliston 1987]. Mouse EC cells do not express Endo B. However, human EC cells, which appear to be representative of a developmental stage equivalent to or even earlier than mouse EC cells, already express the homologous human gene, K18. This implies that very early human and mouse embryos differ in their expression of these homologous genes even though adult tissue expression is equivalent or identical. The larger number of genes similar to Endo B found in primate DNAs [Fig. 1] may reflect this difference in early developmental expression. Expression of K18 in the equivalent of the mouse inner cell mass would provide a target for retrovirus-mediated pseudogene generation in the germ line [Linial 1987]. The extremely limited time...
Oshima et al.

Figure 8. Expression of Endo B and K18 in cellular hybrids. Somatic cell hybrids were constructed and analyzed for the synthesis of Endo B and the homologous human protein K18 by immunoprecipitation \( [A \text{ and } C] \) or for the presence of Endo B and K18 genes by the method of Southern \( [1975] [B] \). \( [A] \) \(^{35} \text{S}\)-methionine-labeled cell lysates of HeLa cells \( \text{[lane 1]} \) or STO mouse fibroblast \( \text{[lane 7]} \) parental cells and hybrid clones derived from the fusion of the two parental cells \( \text{[He}; \text{ST 3.7, 7.1, 7.5, and 9.6]} \) were immunoprecipitated with Endo B antiserum. The immunoprecipitates were separated by acrylamide gel electrophoresis in the presence of SDS and detected by fluorography. \( [B] \) Position of Endo B, [K18] position of keratin 18; [M] mouse endodermal cytoskeletal protein size markers. \( [B] \) Southern analysis of DNA from the HeLa and STO parental cell lines and from HeST hybrids. All DNAs were digested with \( \text{BamH} \), and the filter was hybridized with nick-translated pK187 K18 cDNA plasmid, which detects both K18 and Endo B genes. Final wash was performed at 55°C in 0.1 × SSC, 0.5% SDS. The filter was exposed for 6 days with two intensifier screens. Size markers are indicated on the right \( \text{[in kb]} \). \( [-] \) The 1.7-kb fragment corresponding to the middle portion of the K18 gene. Note the presence of the ∼6.5-kb mouse Endo B gene detected in STO DNA \( \text{[lane 12]} \) and all HeST hybrids, with the possible exception of HeST 10.10. \( [C] \) Immunoprecipitation analysis of Endo B synthesis in cellular hybrids derived from the fusion of HR9 endodermal cells \( \text{[lane 1]} \) and STO fibroblasts \( \text{[lane 6]} \). Endo B protein was detected by immunoprecipitation, as described for panel \( A \). Note that all HR9 × STO hybrids \( \text{[HRSTO 1–6, lanes 2–5]} \) continue to express Endo B.

Several different results indicate that the Endo B \( \beta 1 \) gene is the single mouse Endo B gene transcriptionally active in tissues as diverse as parietal endoderm and liver. The DNA sequence of the first exon of \( \beta 1 \), including the 5’-untranslated leader, matches the Endo B cDNA exactly \( \text{[Fig. 4]} \). The \( \beta 1 \) gene is distinguished from the other mouse Endo B genes in the stability of hybrids formed with a cDNA probe \( \text{[Fig. 2A]} \), its sensitivity to digestion by DNase I in chromatin from cells that express the gene \( \text{[Fig. 6]} \), and in its methylation state \( \text{[Fig. 7]} \). Finally, the compact structure of the regions of the isolated \( \beta 2, \beta 3, \) and \( \beta 5 \) genes that hybridize to the Endo B cDNA is consistent with the expected structure of processed pseudogenes. DNA sequence divergence, as judged by restriction enzyme site mapping or direct determination of the DNA sequence of portions of the \( \beta 2 \) gene, indicates that during which the germ line of rodents expresses Endo B could severely limit the incorporation of pseudogenes into the germ line. The acquisition of the differences in the very early developmental expression of Endo B and K18 may be coincident with the divergence of primates from other mammals.

The presence of five genes related to Endo B in the mouse genome requires the identification of the coding gene for detailed investigations of the regulation of Endo B. We have shown previously that tissues as diverse as parietal endoderm and liver express Endo B proteins that are indistinguishable in electrophoretic mobility, antigenicity, and peptide pattern \( \text{[Trevor and Oshima 1985]} \). In addition, the 5’ end of the Endo B mRNAs expressed in liver and parietal endoderm are identical, as judged by primer extension analysis \( \text{[Trevor and Oshima 1985]} \).
none of these three codes for the parietal endodermal Endo B mRNA. Similarly, the weak hybridization of the β4 putative gene and its hypermethylated state make it an unlikely candidate for an active Endo B gene fragment. We conclude that the cloned 3-kb EcoRI fragment of the β1 gene represents the 5' end of the single Endo B gene active in extraembryonic endoderm and liver.

The differential expression of Endo B in different cell types appears to be due to transcriptional regulation [Fig. 5]. However, at least two different methods of controlling transcription of the Endo B gene are evident. In fibroblasts or myoblasts that do not express Endo B, the 5' end of the β1 gene is both methylated [Fig. 7] and in a relatively DNase I-resistant chromatin state [Fig. 6]. This silent state appears to be quite stable because fusion of a fibroblast with HeLa cells, which express the homologous human protein, does not lead to the activation of Endo B [Fig. 8]. In addition, no evidence for trans-acting negative regulation of Endo B was found in either these hybrids or hybrids made from the fusion of mouse endodermal and fibroblast cells. We have shown recently that a transfected homologous human gene, K18, can be expressed in mouse fibroblasts from its own promoter while the endogenous Endo B gene remains silent (Kulesh and Oshima 1988). This result both reinforces our conclusion concerning the stability of the inactive state of the Endo B gene in mouse fibroblasts and suggests that the interaction of trans-acting factors with the promoters of these genes is not the principal reason for the differential expression of these keratins in fibroblasts and permissive cells. The demonstration that the Endo B gene in myoblasts can be activated by treatment with 5-azacytidine (Darmon 1985), a treatment that results in substantial demethylation of CG dinucleotides (Jones and Taylor 1980), is supportive of a direct role for methylation in the cis-acting negative regulation of Endo B in fibroblasts or myoblasts. However, it is possible that the methylation state of the Endo B β1 gene only reflects the inactive state and is not necessarily causal. Similar considerations must be given the altered chromatin state of the β1 gene in different cell types. It may now be possible to map sequences of the K18 and Endo B β1 genes necessary for the establishment of the stable inactive state found in fibroblasts by introducing recombinant constructions into pluripotent mouse EC cells or embryos and characterizing the activity, methylation status, and chromatin state of the constructions in differentiated cell types derived from the stem cells.

Although both undermethylation and a nuclease-sensitive chromatin state of the Endo B β1 gene are correlated with activity in endodermal cells, they are not sufficient to permit expression in all cell types because the β1 gene in EC cells is both sensitive to nuclease and undermethylated but is not expressed. Unlike fibroblasts, EC cells do not express transfected K18 genes efficiently (Kulesh and Oshima 1988). EC cells that have the potential of differentiating to cells that express Endo B appear to have an additional mechanism of suppressing transcription of the gene. This block may be related to recently reported evidence for a labile inhibitor of Endo A transcription in PCC4 EC cells (Cremisi and Duprey 1987). The sequence of the 5' end of the β1 gene revealed an interesting region of the first exon that might be involved in this regulation. At nucleotide 176, a region very similar to the tRNA-binding site of Moloney murine leukemia virus is found. Recently, this region of the virus has been implicated in the block to viral RNA expression in EC cells (Barklis et al. 1986; Loh et al. 1987; Weihr et al. 1987). It will be of great interest to determine whether this or other regions of the Endo B β1 gene is involved in the suppression of Endo B expression during very early development.

Methods

Cell culture

All cell lines and culture conditions have been described previously (Oshima 1981). Somatic cell hybrids between HeLa cells and STO mouse fibroblasts (HeST hybrids) and HR9 parietal endodermal cells and STO cells (HRSTO hybrids) were constructed as described previously (Oshima et al. 1981; Howe and Oshima 1982). HRSTO hybrids were selected in DME medium supplemented with HAT [100 μM hypoxanthine, 0.4 μM aminopterin, 16 μM thymidine] and 2 mM ouabain. HeST hybrids were selected in HAT medium containing 5 × 10⁻⁶ M ouabain. Primary isolates were subcloned to ensure the absence of contaminating parental cells due to metabolic cooperation. Endo B protein synthesis was measured by immunoprecipitation, as described previously (Oshima 1982).

Molecular probes

Endo B probes were prepared either by nick translation (Meinkoth and Wahl 1984) of the pUC9B7 Endo B cDNA [Singer et al. 1986] or in vitro transcription with bacteriophage RNA polymerases. A HindIII–EcoRI 487-bp fragment of the Endo B cDNA that represents the first 461 nucleotides of the Endo B mRNA was subcloned in the SP64 vector (Promega, Madison, Wisconsin). The EcoRI-digested plasmid was transcribed with SP6 polymerase in the presence of [32P]GTP [Melton et al. 1984] to generate a truncated Endo B mRNA probe.

The transcriptional start site of the Endo B β1 gene was mapped by the S1 protection method. A 658-bp fragment of the SK3112 plasmid derived from the SstI site at nucleotide −562 to the EcoRV site at nucleotide 196 was subcloned into M13mp18 that had been digested with SstI and HincII. The sequencing primer and the Klenow fragment of DNA polymerase was used to generate a single-stranded radioactive probe that protected the first 196 nucleotides of Endo B mRNA (Kulesh and Oshima 1988).

Isolation of Endo B genes

The BALB/c mouse embryo genomic library in Charon 4A (Maki et al. 1980) and the strain 129 mouse liver library in EMBL3A were gifts from R. Maki [La Jolla Cancer Research Foundation]. Philip Leder [Harvard Medical School] provided the Charon 28 mouse embryonic DNA library. The EMBL3 library of C57BL/6 T-cell DNA [provided by R. Maki] was prepared by standard methods [Maniatis et al. 1982] after partial digestion with MboI. All were screened by the method of Benton and Davis (1977), using either nick-translated Endo B cDNA or an RNA probe of the first 461 nucleotides of the Endo B cDNA. To obtain the Endo B β1 gene fragment, HR9 mouse parietal endo-
Oshima et al.

dermal DNA was digested with EcoRI, size fractionated by agarose gel electrophoresis, and recovered with powdered glass (Vogelstein and Gillespie 1979). The DNA of a fraction that hybridized strongly to a sense RNA probe of the 5' end of the Endo B cDNA was cloned into AZAP (Stratagene, La Jolla, California) and packaged and screened without amplification. K802 cells (Wood 1966) were used as host. Approximately 400,000 pfu were screened by transfer to nitrocellulose and hybridization with the 461-nucleotide RNA probe. After two additional rounds of purification and screening, eight of nine isolates were rescued from the phage as plasmids by coinfection of XL-1 cells with the individual AZAP isolates and R48 helper phage according to the suppliers instructions. Five of the isolates grew slowly, and gel analysis of the supernatants of these cultures revealed varying amounts of presumptive helper phage DNA, which appeared to correlate with their slower growth. Six of the rescued plasmids appeared to be very closely related by restriction enzyme mapping and Southern analysis. One of these was analyzed in detail after transfection into DH-1 cells to ensure the absence of helper phage and is designated SK3112. This plasmid represents ~3 kb of the Endo B β gene inserted in the EcoRI polylinker site of the Bluescript SK Ml 3m vector (Stratagene, La Jolla, California).

**DNA sequence**

For DNA sequence determination, a 1.6-kb BamHI fragment of SK3112, derived from sites located at -927 and 20 nucleotides downstream of the 3' EcoRI cloning site (nucleotide 648), was subcloned into M13mp18 in both orientations. Ordered deletions were constructed utilizing exonuclease III and nuclease S1 (Henikoff 1984) after digestion with both SphI and XbaI. The sequence was determined by the dyeoxy nucleotide chain termination method (Sanger et al. 1977) using modified T7 DNA polymerase (Sequenase, U.S. Biochemical) and 33S-labeled dATP (Biggin et al. 1983). The final sequence was assembled from data of 13 different deletions for one strand and 15 for the opposite strand, using the GEL program of the BIONET computer system (Intelligenetics, Mountain View, California).

**DNase digestion of Endo B genes**

Nuclei were prepared by rinsing cell cultures three times in cold PBS, scraping the cells into 25 ml of cold lysis buffer [0.25 M sucrose, 0.5% NP-40 nonionic detergent, 10 mM Tris–HCl (pH 7.2), 10 mM NaCl, 5 mM MgCl₂, 0.1 mM EGTA, phenylmethylsulfonyl fluoride (PMSF)] and low-speed centrifugation at 4°C. The nuclei were resuspended in lysis buffer with use of a Dounce homogenizer and centrifuged through a cushion of 30% sucrose in 10 mM Tris–HCl (pH 7.2), 10 mM NaCl, and 3 mM MgCl₂. The nuclei were then suspended in the same buffer without sucrose and containing 0.1 mM CaCl₂ at an A₆₀₀ of 20 (determined on an aliquot digested with micrococcal nuclease and dissolved in 1 mM NaOH), and digested for 10 min at 37°C with varying concentrations of protease-free DNase I (Otsuka and Price 1974). Digestion was stopped by the addition of EDTA to 10 mM and SDS and proteinase K to final concentrations of 1% and 200 μg/ml, respectively. After incubation overnight at 37°C, the DNA was purified by extraction with phenol-chloroform, chloroform, and ether and was ethanol precipitated. Aliquots of DNA were digested with restriction enzyme and analyzed by the method of Southern (1975).

**Nuclear run-on analysis**

Nuclei were prepared as described by others (Wang et al. 1985), with modifications. All cells were grown in roller bottles and washed with phosphate-buffered saline while still attached, scraped into Buffer A [0.3 M sucrose, 10 mM Tris–HCl (pH 7.2), 5 mM MgCl₂, 0.1 mM K₃EGTA, 0.4% NP-40, 0.5 mM dithiothreitol (DTT)], and recovered by low-speed centrifugation. After homogenization and washing once, as described (Wang et al. 1985), the nuclei were treated with protease and DNase-free RNase (20 μg/ml) for 5 min at 20°C, washed twice more by centrifugation through a sucrose cushion, washed once in 40% glycerol, 50 mM Tris–HCl (pH 8.3), 5 mM MgCl₂ and 0.1 mM EDTA, resuspended in the same buffer at 2 x 10⁶ nuclei/ml, and stored at -85°C. Transcription reactions were performed in 20% glycerol, 25 mM Tris–HCl (pH 8.3), 3 mM MgCl₂, 70 mM KC1, 0.5 mM MnCl₂, 2.5 mM DTT, 1 U/μl RNasin RNase inhibitor (Promega), 0.8 mM ATP, 0.4 mM CTP and GTP, 8 μM UTP, and 250 μCi of [α-32P]UTP (800 Ci/mM) in 200 μl containing 2 x 10⁷ nuclei for 10 min at 26°C. Radioactive RNA was purified (Groudine et al. 1981) and hybridized to denatured DNA immobilized on nitrocellulose in 50% formamide, 5 x SSPE, 0.1% SDS, 250 μg/ml salmon DNA, and 2 x Denhardt's solution (Maniatis et al. 1982) at 42–60°C for 4 days. Filters were washed several times in 2 x SSPE, 0.1% SDS at room temperature, rinsed in 2 x SSPE, and treated with 10 μg/ml RNase A in 0.15 M NaCl, 10 mM Tris–HCl (pH 7.2) and 1 mM EDTA for 30 min at room temperature and, finally, washed in 0.1 x SSPE, 0.1% SDS at 50–65°C. Filters were then exposed to film at -85°C with an intensifier screen.

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**Note**

Sequence data described in this paper have been submitted to the EMBL/GenBank Data Libraries under the accession number Y00217.

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