Oestradiol decreases rat apolipoprotein AI transcription via promoter site B

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ABSTRACT

Oestrogens protect against ischaemic heart disease in the post-menopausal female by increasing serum concentrations of apolipoprotein (apo) AI and the abundance of high-density lipoprotein particles. In men and experimental male animals, the administration of oestrogen has variable effects on apo AI expression. As the major mode of oestrogen action on target genes involves regulating promoter activity and hence transcription, oestrogen is expected to alter transcription of the apo AI gene. To test this hypothesis, the effect of 17β-oestradiol (E₂), on rat apo AI promoter activity in male hepatoma HuH-7 cells, was tested by co-transfecting a reporter template, pAI.474.CAT containing −474 to −7 of the rat apo AI promoter and an oestrogen receptor (ER) expression vector, pCMV-ER. Transfected cells exposed to E₂ showed a dose-dependent decrease in chloramphenicol acetyltransferase (CAT)-activity, with a maximum 91% reduction at 1 µM E₂. Deletional analysis of the promoter localized the inhibitory effect of ER and E₂ to site B (−170 to −144) with an adjacent 5′ contiguous motif, site S (−186 to −171) acting as an amplifier. HuH-7 cell nuclear extracts showed binding activities with both sites S and B, but recombinant human ER did not. Furthermore, nuclear extracts from E₂-treated HuH-7 cells showed weaker binding activity to site B, but not to site S. In summary, the inhibitory effect of ER and E₂ on rat apo AI gene activity is mediated by a promotor element, site B. This inhibitory effect arises from a mechanism that does not involve direct ER binding to the B-element. The conclusion that E₂ inhibits apo AI transcription was confirmed in vivo. Treatment of male adult Sprague–Dawley rats with up to 200 µg E₂ for 7 days decreased apo AI protein and hepatic mRNA by 72±21% and 68±1·4% respectively. Results of ‘run-on’ transcription of the apo AI gene in isolated hepatic nuclei showed a 55% decrease in hormone-treated male rats. These findings suggest that E₂ exerts primarily an inhibitory effect within male hepatic nuclei.

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INTRODUCTION

Cardiovascular disease (CVD) is the leading cause of premature death of men and women in developed countries. Although death rates from CVD are similar, at 46% and 52% in men and women, the major difference is that the peak incidence in women occurs 10 years later than in men (Barter & Rye 1996). The protection afforded to premenopausal women is in large part due to the beneficial effects of endogenous oestrogens. Oestrogen deficiency, caused by surgical or natural menopause, increases the risk of CVD in women by threefold (Barrett-Connor & Bush 1991). Furthermore, epidemiological studies suggest that oestrogen replacement therapy in postmenopausal women reduces the risk of CVD by up to 50% (Bush 1990, Stampfer & Colditz 1991). In women with prior myocardial infarction, the benefit of hormone replacement may be as much as a 90% reduction...
in CVD risk (Sullivan et al. 1990). The precise mechanisms underlying the cardioprotective action of oestrogens are unknown. Although the mechanism is most probably multifactorial (Shewmon 1994), a major reduction in CVD risk is attributed to the effects of oestrogen on the metabolism of apolipoprotein (apo) AI and high-density lipoprotein (HDL) and low-density lipoprotein (LDL) particles (Bush et al. 1987).

Previous studies demonstrated that pre- or postmenopausal women treated with exogenous oestrogens have an increased abundance of apo AI and HDL (Walsh et al. 1991). However, oestrogen does not appear to augment the concentrations of apo AI and HDL in males and thus does not appear to protect individuals from CVD. Studies detailing the effect of oestrogens in males have shown contradictory findings. For example, male patients with prostate cancer or those undergoing sex-change procedures who are treated with exogenous oestrogen have a greater risk of developing new myocardial infarctions (Fortin et al. 1984, Henriksson et al. 1987). Similarly, hyperoestrogenaemia in males attributed to increased hepatic conversion of testosterone as a result of cigarette smoking, alcohol intake, diabetes, or drug abuse carried a greater risk of CVD, in some (Morgan 1982, Klaiber et al. 1984, Phillips 1984, Glazer 1991), but not all studies (Ruebner et al. 1961). In contrast, other data suggest that oestrogens reduce the risk of CVD in males (Marmorston et al. 1960, Farhat et al. 1996). In addition, newborn animals, pre-pubertal boys and men with cirrhosis of the liver—that is, those males with high concentrations of oestrogens but low testosterone–have reduced prevalence of CVD (Fangman & Hellwig 1947, Neufeld et al. 1962, Sukurai et al. 1960, O’Kane et al. 1997). In experimental animals the data are equally confusing; various studies showing decreased (Weinstein et al. 1986, Srivastava et al. 1993, Zou & Ing 1998), increased (Seishima et al. 1991) and no (Srivastava et al. 1997) oestrogenic effect on apo AI expression in the male.

The preceding data portray an unclear picture of the cardioprotective role of oestrogen in normal intact males. As oestrogens and other steroid hormones increase apo AI expression in the female through activation of the apo AI promoter, we examined the effect of 17β-oestradiol (E₂) on apo AI promoter activity in males. The following studies show that E₂ inhibits apo AI gene transcription via a cis-acting promoter motif designated site B. In addition, E₂ increases female, but decreases male, rat serum apo AI protein concentrations.

MATERIALS AND METHODS

Materials

The rat apo AI cDNA, rat apo AI antibody, wild-type, mutant and site-specific pAI.CAT constructs have been described previously (Romney et al. 1992, Taylor et al. 1996a). Both the human ribosomal RNA (Sylvester et al. 1986) and pRSV-β-galactosidase plasmids were gifts (Chan et al. 1993). The p5’S.CAT, p5’S2.CAT and p5’S3.CAT constructs were created by ligating one, two or three copies, respectively of site S (gatccGCAGCCCCGACGCTTCCTGTG)gibco BRL, Burlington, Ontario, Canada) into the SmaI site of pGEM-CAT.5’ (Romney et al. 1992). The SB.CAT construct was created by annealing the antisense strand of site S with site B (gatccAGGCTAGGCAACAGGTGGGCAA–ACAGGAAc), followed by a ‘fill in’ reaction with the Klenow fragment of DNA polymerase I. The product was then cloned into the SmaI site of the pGEM-CAT.5’ in the correct orientation as a monomer. Complementary strands of both sites S and B with protruding EcoRI ends and an oestrogen response element (ERE; aattCGGTCACTGTG–ACGGAaatt) were purchased from Integrated DNA Technologies, Inc., Coralville, IA, USA, for use in electrophoretic mobility shift assays (EMSA). Human recombinant oestrogen receptor protein (ERα) was purchased from Affinity Bioreagents, Inc., Golden, CO, USA. The pCMV-ER was a kind gift from Dr R Mischeck, University of New York.

Cell culture and transfection

Male human fetal hepatoma HuH-7 cells were cultured in serum-free RPMI-1640 medium, as previously described (Nakabayashi et al. 1989). This medium contains 1·4 × 10⁻⁵ M phenol red but, to observe the effect of exogenous E₂, cells are cultured in medium that lacks phenol red because this compound is weakly oestrogenic (Bindal et al. 1988). In the absence of phenol red, HuH-7 cells rapidly die (authors’ unpublished observations). Therefore, for E₂ studies, HuH-7 cells were subcultured to 25 cm² flasks for 48 h in RPMI-1640 medium and then transferred to medium containing a lower concentration of phenol red (7 × 10⁻⁷ M) for a further 24 h before transfection in the same medium. After transfection (Romney et al. 1992), cells were exposed to E₂ for 24 h and then harvested (Taylor et al. 1996a). Cell lystate was assayed for β-galactosidase and chloramphenicol acetyltransferase (CAT) activities as described elsewhere (Taylor et al. 1997).
intact nuclei for remainder of the fresh liver was used to prepare method (Promega, Madison, WI, USA). The guanidinium isothiocyanate liver were snap-frozen for RNA extraction using analyses. Approximately 500 mg freshly excised proteins for EMSA studies.

Northern analysis
Total hepatic RNA (10–40 µg) was analysed by northern blot analysis (Sambrooke et al. 1989). The 28S and 18S rRNA were visualized by staining with ethidium bromide to ensure that the RNA was intact, and equal amounts were loaded. Contents of the gel were blotted to Zetabind membrane (AMF Cuno Inc., Meriden, CT, USA) and probed with radiolabelled rat apo AI cDNA (Taylor et al. 1996a). Apo AI mRNA abundance was determined by video-assisted densitometry and normalized to the density of the 18S RNA obtained from a photographic negative of the ethidium-bromide-stained gel.

Preparation of nuclei and nuclear ‘run-on’
Preparation of rat liver nuclei and their use in the in vitro transcription assay were as described previously (Taylor et al. 1996a). Each labelling reaction contained 1.3–4.8 × 10⁷ nuclei. Blots were hybridized at 65 °C with 3.3–4.3 × 10⁶ c.p.m. ‘input’ RNA. Membranes were washed at high stringency and signal detected by autoradiography using Kodak XAR-5 film for 2–4 days at –80 °C. The hybridization signal was assessed by video-assisted densitometry. These values were expressed relative to rRNA standards and then normalized for both ‘input’ RNA per hybridization and the number of nuclei per assay.

Statistical analysis
Data are presented as mean ± s.d. and significance determined by either unpaired Student’s t-test or one-way analysis of variance (ANOVA) using the least significance difference test provided in the Clinstat software package (Bland 1990).

RESULTS
Site B mediates the repressive effects of E₂
To determine whether E₂ inhibits apo AI promoter activity, we measured activity of a construct, pAI.474.CAT containing the –474 to –7 bp fragment of rat apo AI DNA in a male human hepatoma cell line, HuH-7, co-transfected with an ER expression vector, pCMV-ER. Subsequent exposure of these cells to E₂ caused a dose-dependent decrease in promoter activity with a maximal 91% reduction at a hormone concentration of 5 × 10⁻⁶ M (Fig. 1).

To locate the motif that mediated the effects of E₂, we assayed the activity of deletional constructs
containing smaller amounts of the promoter (schematic map; Fig. 2A) in the presence of pCMV-ER and $10^{-6}$ M E$_2$. Removal of the DNA between $-474$ to $-187$ bp did not significantly alter the inhibitory actions of E$_2$ (pA1.186.CAT, Fig. 2B). However, most of the inhibitory effect of E$_2$ was lost in constructs lacking $-474$ to $-170$ bp or $-144$ bp ($60 \pm 7\%$ and $25 \pm 2\%$ respectively) compared with control. Together these findings show that the cis-acting element spanning $-170$ to $-145$ bp mediates a dominant role in the inhibitory actions of E$_2$, and that an adjacent element, $-186$ to $-171$ bp, has a minor role. These two elements correspond to two motifs, B and S, that we described previously (Taylor et al. 1996a). A minimal promoter, $-46$ to $-7$ bp, was non-responsive to E$_2$ (pA1.46.CAT; Fig. 2). These observations suggest that the inhibitory effect of E$_2$ is mediated by site B.

Next we examined the ability of site B to transmit the inhibitory actions of E$_2$. For these studies we created a template, p5'B.CAT, containing duplicate B sites placed in front of the heterologous
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SV40 promoter that is fused to the CAT-gene (Fig. 2A). CAT activity in E2-treated HuH-7 cells co-transfected with p5′B.CAT and pCMV-ER was decreased by 49 ± 13% (p5′B.CAT, Fig. 2). The specificity of this effect to site B was demonstrated in the study of similar constructs, pS1.CAT, pS2.CAT, pS3.CAT and p5′A.CAT, that contained one, two or three copies of site S and duplicate site A respectively. E2 had no effect on activities of these constructs (Fig. 2).

Potential roles of site S
To examine the potential roles of site S (−186 to −171 bp), we created pSB.CAT, which contains sites S and B fused in tandem to the SV40 promoter 5′ to the CAT-gene (Fig. 2A). E2 inhibited the activity of this construct by 72±4% (Fig. 2B). This level of repression was more pronounced than the sum of that of either site S (pS1.CAT; 2.3% reduction) or site B alone (pAI.170.CAT; 58.4% reduction). Together, these data suggest that E2 acts through site B and that site S amplifies or augments the inhibitory actions of E2 through site B.

E2 receptor does not bind to sites S and B
In an attempt to define the mechanism by which E2 inhibits the transcription of the rat apo A1 promoter, EMSA were performed with purified human recombinant ERα (hERα) to [32P]ERE. Purified hERα bound to [32P]ERE (lane 1, Fig. 3). The addition of 100-fold M excess unlabelled ERE blocked the formation of the ER–ERE complex, but excess site S or B (lanes 2–4 respectively, Fig. 3A) had no effect. These findings indicate that hERα binds to the ERE, but not to sites S or B.

Nuclear protein binding to sites S and B
Next we examined the ability of HuH-7 nuclear proteins to bind sites S and B (Fig. 3B). Incubation of nuclear extracts with radiolabelled site S revealed the presence of at least three binding activities (lane 2, Fig. 3B). These activities were unaffected by the treatment of cells with E2 (lane 3, Fig. 3B). In contrast, the pattern of nuclear protein binding to site B DNA was more complex (lanes 4–8, Fig. 3B). Nuclear proteins from HuH-7 cells had at least five site B binding activities. The intensities of the three complexes with the slowest mobilities (asterisks and arrow) were less abundant in E2-treated cells transfected with ER (compare lanes 5 and 6 with 7 and 8, Fig. 3B). In addition, there was a slight increase in intensity of the second most mobile band in extracts from E2-treated cells (Fig. 3B), but this complex represented non-specific binding. Only a single complex was specific in competition analysis (arrow, Fig. 3C). The abundance of this complex was decreased in response to E2. The most mobile band (denoted by a dot in Fig. 3B) remained unchanged. These data show that E2 decreases specific site B binding activity.

Mutation of HNF3 elements in site B abolish actions of ER
Site B in the apo A1 promoter is comprised of direct repeats, with two HNF-3 motifs of 9 bp separated by a 4 bp spacer (Fig. 4A). To determine whether this element mediates the repressive actions of ER and E2, we used a reporter template, BCD.CAT (Fig. 4A). The activity of this template was similar to that of pAI.170.CAT in the absence or presence of pCMV-ER and E2 (compare Fig. 4B and Fig. 2). CAT activity in cells transfected with BCD.CAT alone was not significantly different from control after treatment with E2. However, in the presence of ER there was a 89% reduction in CAT activity from 7.85% to 0.85% CAT conversion/µg protein/h (Fig. 4B). An additional 45% reduction was evident in the presence of exogenous E2 (from 0.85% to 0.473% CAT conversion/µg protein/h). The inset to Fig. 4B shows a western blot analysis of lysate from control cells or those transfected with pCMV-ER. ER was not detected in control cells, but is clearly evident in those containing the expression vector. This result supports the finding that E2 suppression of the reporter requires ER.

Next, we used three templates containing mutations to the HNF-3 sites. First, we found that pAI.Bm1.CAT, which contained a mutation of the HNF-3 motif at the 5′ end of site B, had a dramatic reduction in basal activity from 7.85% to 0.17% CAT conversion/µg protein/h (Fig. 4B). Despite this low expression, the activity of this construct was further reduced by 25% in the presence of ER (Fig. 4C). However, no further reduction was evident after the addition of E2. Another template, pAI.Bm2.CAT, containing a mutation of the HNF-3 motif at the 3′ end of site B, showed a similar reduction in basal activity and, like pAI.Bm1.CAT, it was no longer responsive to E2 (Fig. 4C). These data confirm that the inhibitory action of ER requires an intact site B. In addition, the construct pAI.158.CAT – a template that lacks the 5′ HNF-3 site – had very low basal CAT activity (0.47% CAT conversion/µg protein/h) compared with BCD.CAT (Fig. 4B). However, the activity of this construct, which retains an unmodified 3′ HNF-3 site, was reduced by 28% in the presence of
ER (Fig. 4C). Together, these mutants and deletion analyses show that intact HNF-3 sites are required for E2 to exert its inhibitory actions in the presence of ER and the 3’ HNF-3 motif alone appears capable of mediating the repressive actions of ER.

These data suggest that the HNF-3 sites have a critical role in mediating the actions of E2.

**E2 decreases apo AI abundance, expression and transcription**

As expected, administration of E2 to female rats produced a significant 2.9-fold up-regulation of serum apo AI concentrations (Fig. 5A, B). However, administration of E2 to male rats had the opposite effect on apo AI expression: male animals injected with E2 for 7 days had 72% less apo AI protein in serum (Fig. 5C, D) and a 68% lower hepatic mRNA content (Fig. 6A, B). Similarly, E2 also reduced the 'run-on' transcription rate of the gene by 55%, from $12.9 \pm 3.1$ to $5.8 \pm 0.77$ ng RNA/10^6 nuclei/h (Fig. 6). However, the rate of albumin gene transcription was not significantly affected by hormone administration (Fig. 6D).

These data suggest that E2 reduced male apo AI serum protein and hepatic mRNA levels via a reduction in transcription of the gene.

**DISCUSSION**

The major economic and social impact of CVD on members of our society makes this condition an important topic for biomedical research. Apo AI

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**FIGURE 3.** EMSA of ER binding to ERE but not to apo AI sites S or B. (A) Purified human recombinant ERα protein was incubated with [32P]-labelled ERE for 30 min in EMSA binding buffer (lane 1) supplemented with $5 \times 10^{-5}$ M E2. A 400-fold molar excess of unlabelled ERE (lane 2), site S (lane 3), or site B (lane 4) was added to the basic reaction mixture. Bound and free radioactivity were visualized as described in Materials and Methods. The autoradiogram is a representative of four separate experiments. (B) E2 decreases binding to site B but not site S. Nuclear extracts from ER-containing HuH-7 cells treated with or without E2 were incubated with either [32P]-labelled sites S (lanes 2 and 3 respectively) or B (lanes 5 and 6, and 7 and 8 respectively) for 30 min. DNA–protein complexes that were reduced after treatment with E2 are indicated by asterisks and an arrow. Non-specific complexes are indicated by asterisks or the dot. The autoradiogram is representative of three separate experiments. (C) Competition analysis. Site B binding activity in extracts from control cells is bound to radiolabeled DNA (lane 1) in the presence of 100-fold M excess unlabeled authentic HNF-3β (lane 2) or site B (lane 3) binding motif. Addition of excess non-specific DNA had no effect on intensity of the complexes (data not shown). The only specific complex is indicated by an arrow.
and HDL act as protectors against CVD by amelioration of hypercholesterolaemia via several mechanisms that decrease cholesterol serum concentrations (Kashyap 1998). Therefore, it seems reasonable to postulate that to increase rates of synthesis of serum apo AI would have beneficial effects on health. In order to work towards this goal, we have embarked on a detailed analysis of the apo

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**Figure 4**. Mutation of HNF-3 sites in site B abolishes oestrogen responsiveness. (A) Schematic diagram and site B sequences showing the rat apo AI promoter−CAT constructs used in the transfection studies. HNF-3 sites are represented by the arrows and mutations are underlined. Numbering is relative to the transcription start site. (B) HuH-7 cells were transfected with wild-type (pAI.BCD.CAT), mutated (pAI.Bm1.CAT, pAI.Bm2.CAT), or deleted (pAI.158.CAT) pAI−CAT constructs. Parallel cultures were grown and treated with 10⁻⁶ M E₂. Cells co-transfected with pCMV-ER were treated with vehicle and 10⁻⁶ M E₂ for 24 h. Cellular proteins were assayed for CAT activity and the data presented relative to pAI.BCD.CAT. Values are shown as means ± s.d. from two experiments performed in triplicate. The insert shows levels of ERα as detected by western blot of lysates from cells that were non-transfected or transfected with pCMV-ER. *P<0.05, **P<0.01; ANOVA with least significant difference test.
AI promoter. The clinical observation that E₂ in women reduces their risk of developing CVD prompted us to investigate whether this beneficial effect can be extended to males, by examining effects of the hormone on the apo AI promoter. Results of these studies show that E₂ decreases apo AI promoter activity in males.

In the initial transfection experiment with pAI.474.CAT and E₂, an inhibitory effect of E₂ was demonstrated and shown to be dependent on the dose of hormone in HuH-7 cells, which are derived from a male liver (Fig. 1). Similar studies performed on the human apo AI promoter in HepG2 cells (also derived from a human male liver),
co-transfected with a similar ER expression plasmid, showed similar dose-dependent decreases in apo AI promoter activity (Harnish et al. 1998). Maximal inhibition of reporter transcription from the human apo AI promoter in HepG2 cells was an order of magnitude lower than in the results presented here using a rat apo AI promoter construct (5 × 10⁻⁸ M compared with 5 × 10⁻⁷ M in HuH-7 cells). The difference in extent of inhibition by E₂ in these two sets of data may be due to differences in the nature of the human and rat promoters. Although these promoters share greater than 90% homology, there is a significant difference in their responses to many transcription factors, including ROR and PPAR (Vu-Dac 1997, Staels & Auwerx 1998). However, although the human and rat apo AI promoter differ in response to some factors, in both HepG2 cells and HuH-7 cells, the apo AI promoter activity is inhibited by E₂.

The expression of oestrogen-responsive genes such as ovalbumin and IGF-1 are also regulated by E₂ at the transcriptional level. In these cases, liganded ER interacts with ERE motifs in the promoter to decrease gene expression (Sanders & McKnight 1988, McCarthy et al. 1997). Other genes such as the EGF receptor, the progesterone receptor and ER itself are up-regulated at the transcriptional level via promoter-specific ERE motifs. However, an examination of the apo AI promoter DNA for consensus ERE motifs (GGT-CAnnTGACC), shows none to be present. As some E₂-dependent genes lack consensus EREs, E₂ action may also be mediated via non-genomic mechanisms or via other cis-acting elements. For example, E₂ regulates gene transcription via AP-1 and SP-1 sites (Cerillo et al. 1998, Xie et al. 1999) in which the liganded ER does not contact the DNA.

The lack of a consensus ERE in the rat apo AI gene led us to perform deletional analysis of the rat apo AI promoter. The element that mediates the inhibitory actions of E₂ is site B. This conclusion comes from the fact that removing DNA spanning –186 to –145 bp caused the inhibitory effect of E₂ on promoter–CAT activity to lessen from 85% to 20%. There are two cis-acting sites, which we have designated site S and site B, in this fragment of the promoter. Removal of site S (–186 to –171) caused a 25% change in E₂ effectiveness from 85% to 60%, indicating that this portion of the fragment has a minor role in the actions of E₂. Further removal of site B (–170 to –145) caused a 40% change in E₂ effectiveness, indicating that site B is the major mediator of E₂ inhibition. This idea was confirmed by the observation that E₂-inhibited the CAT activity of the construct, p5'B.CAT, but not that of p5'A.CAT, p5'S.CAT, p5'S2.CAT or p5'S3.CAT. As expected, the control template, p5'A.CAT containing site A, a thyroid hormone receptor (TR) and/or a retinoic acid responsive element (Taylor et al. 1996b) was not affected by E₂. Although ER can bind to 1,25-dihydroxyvitamin D₃ response elements (TREs) and TRs bind to EREs, they do not transactivate the genes involved (Dellovade et al. 1996). Indeed, bacterially-expressed TR1 binds to ERE and hERα binds to TRE (data not shown, and Klinge et al. 1997), but clearly ER does not affect the activity of the A-site (p5'A.CAT, Fig. 2).

Our data also show that site S is not an ERE, because E₂ failed to affect the CAT activity of constructs containing a monomer, dimer or trimer of site S (Fig. 2). In addition, site S must be present for site B to be fully active, as the activity of a construct containing only the B-element, p5'B.CAT, was inhibited to a lesser extent than that of pSB.CAT, which contained both sites S and B (Fig. 2). These data add further support to the idea that site S is an amplification sequence, as previously reported in the response of apo AI to glucocorticoid (Taylor et al. 1996a). At present, the trans-acting factors that may possibly bind to site S include Egr-1 and PEA-3 (Nerlov et al. 1992, Kilbourne et al. 1995) – developmental genes that are regulated by E₂ in vitro (Lockwood et al. 1998, Pratt et al. 1998). There are other developmental genes, such as the homeobox gene family member proteins Hox a10 and Hox a11, which are also regulated in an E₂-dependent manner (Ma et al. 1998). Although the responsive elements for the Hox family of transcription factors are not known, it is possible that Hox genes might interact with the apo AI promoter, and we speculate that a cascade of transcriptional factors and elements are involved in some of the actions of E₂ on rat apo AI promoter repression. Furthermore, trans-acting factors that bind to adjacent cis-acting elements – HNF-4, AR, and others – may play a role in the inhibitory effect of E₂ through protein–protein interactions (Harnish et al. 1998, Hargrove et al. 1999).

The results of the EMSA studies suggest that the inhibitory effect of E₂ on apo AI transcription does not require binding of hERα to either site S or site B. As expected, hERα binds to radiolabelled ERE (Fig. 3) and the formation of this complex is inhibited by unlabelled ERE. However, neither excess site S nor site B sequences are effective in displacing hERα from the radiolabelled probe. In addition, the converse experiment of hERα added to radiolabelled site S or site B produced no visible protein–DNA complexes (data not shown). These observations indicate that hERα does not directly
bind to site S or site B and ER probably decreases apo AI transcription via an indirect mechanism.

The preceding observations prompted us to search for a potential mechanism using EMSA. If ER does not transmit the inhibitory effects of E2 directly by binding to apo AI site B, then what does? Nuclear extracts from E2-treated HuH-7 cells had no effect on protein binding to radiolabelled site S, but did decrease nucleoprotein binding to site B (Fig. 3). This finding suggests that loss of hepatonuclear protein binding to apo AI site B is responsible for the observed decrease in apo AI transcription.

Site B contains two HNF-3 motifs (Fig. 4) that bind HNF-3 (Harnish et al. 1994). Constructs containing a mutation or deletion of either HNF-3 motif were no longer responsive to E2 (Fig. 4). These data are qualitatively similar to those from previous studies in which removal of the HNF-3 motifs from site B caused the loss of basal promoter activity (Harnish et al. 1994) and stimulated response to dexamethasone (Taylor et al. 1996a). The present observations show that both HNF-3 motifs are required by the promoter to maintain full responsiveness to ER and E2. It is conceivable that E2 influences the activity, abundance, or both, of HNF-3 isoforms (Lai et al. 1993, Harnish et al. 1998) or HNF-3-like proteins (Jacquemin et al. 1999) to transmit its effect on site B. This hypothesis also suggests that the HNF-3 isoforms or HNF-3-like proteins would have opposing activities on apo AI transcription via site B. Indeed, HNF-6 (an HNF-3-like transcription factor) has recently been shown to inhibit HNF-3 trans-activation in a gender-specific manner (Lahuna et al. 1997).

The literature provides a confusing picture of the effect of E2 in experimental animals when compared with that in humans, with the species and strain of experimental animal being of utmost importance. Our studies on serum protein concentrations showed that administration of E2 to female and male Sprague–Dawley rats increased and decreased serum apo AI concentrations respectively (Fig. 5). These observations negate our original hypothesis that the beneficial effect of E2 on apo AI serum concentrations in females would also be observed in males. In physiological doses, E2 had no significant effect on serum apo AI concentrations in male rats (data not shown). These data are consistent with the findings of a number of studies, showing either that male rats do not respond to administration of E2, or that it decreases their apo AI secretion rate (Weinstein et al. 1986, Srivastava et al. 1993, 1997, Zou & Ing 1998).

In some, but not all, studies transcription of apo AI was shown to be unaffected, and the main effect of E2 occurred through gene translation (Srivastava et al. 1997). However, these data are not consistent with other data showing that 17β-ethinyl oestradiol in male rats increased apo AI concentrations (Seishima et al. 1991). In the present study, we used intact animals to measure serum apo AI and used high doses of E2. Perhaps this combination could explain the discrepancy between the findings presented here and those of others that involved castrated animals, smaller doses of E2, measured plasma apo AI concentrations or used alternative E2 formulations or routes of administration. As reporter assays may be more sensitive than in vitro studies because of ligand availability, we decided to increase the dose of E2 to a pharmacological value that simulates the concentrations of E2 in a pregnant woman (Check 1999). Our data suggest that supraphysiological doses of E2 in the male rat decrease the transcription of apo AI, in keeping with the findings of some animal studies (Weinstein et al. 1986, Srivastava et al. 1993, Zou & Ing 1998). If decreased translation of apo AI mRNA is the main mechanism of decreased apo AI serum protein concentrations in rats (Srivastava et al. 1997), transcriptional rates and steady-state-levels of apo AI mRNA would presumably remain constant.

It is clear from our western and northern blot analyses of apo AI protein and hepatic RNA that E2 decreases both the abundance of apo AI protein in serum and mRNA in the liver of treated male rats. Furthermore, nuclear ‘run-on’ experiments show that the effects of E2 on apo AI gene expression arise from its ability to inhibit transcription of the gene in hepatic nuclei of hormone-treated male rats. The 55% reduction in transcription rate is similar to the 70% reduction in mRNA and 60% reduction in serum protein concentrations. Although we have observed a good correlation between decreased apo AI transcription rates, steady-state mRNA and secreted protein concentrations, we cannot rule out that E2 could still exert a major effect on apo AI translation. Indeed, this may explain the differential effect of E2 in the male and female rats used in the present study. First, the relative concentrations of apo AI in male and female serum were different: males had significantly greater amounts of apo AI/µg total serum protein. Secondly, in male rat hepatocytes, high doses of E2 decrease apo AI transcription, but the hormone may also alter the expression of other proteins involved in the secretion or processing of apo AI. In the female rat hepatocyte, high doses of E2 also decrease apo AI transcription, but the hormone may increase the activity of proteins involved in apo AI secretion and processing. This might lead to the observed gender differences in response to high-dose E2. In addition,
$E_2$ increases the expression of hepatic LDL receptor and lipoprotein lipase, which could increase the availability of re-circulated apo AI within the hepatocyte, and in turn may have an inhibitory effect on apo AI transcription via altered translation, as a result of apo AI/lipoprotein recycling events (Song et al. 1998).

The findings summarized above show that, in contrast to females, male rats treated with low-dose $E_2$ do not have the expected increases in the serum concentration of apo AI; furthermore, males treated with high doses of $E_2$ show decreased serum apo AI concentrations. These results support the idea that $E_2$ has a detrimental effect on lipid metabolism in the male (Kalin & Zumoff 1990) and are consistent with a recent report in which male rats treated with 17β-ethinyl oestradiol had undetectable apo AI (Landschultz et al. 1996). This information helps to explain the observations made in clinical studies (Fortin et al. 1984, Henriksson et al. 1987), in which the use of exogenous $E_2$ to prevent further myocardial infarctions had the opposite effect, and actually enhanced CVD risk in males. A decrease in apo AI expression may not be the only factor involved in the increased CVD risk in $E_2$-treated males, because $E_2$ has a number of additional effects on the progression of CVD. For example, $E_2$ has potent thrombogenic activity, and increases prostacyclin and prostaglandin secretion in the vasculature, but decreases intima thickening (Chang et al. 1980, Gisclard et al. 1988, Iafrati et al. 1997).

One could postulate that, as $E_2$ decreases apo AI in male Sprague–Dawley rats, the hormone would have a similar effect in the human male and, therefore, the use of $E_2$ in men would be contraindicated. It could actually enhance CVD risk in males. A decrease in apo AI expression might have a clinical relevance.

Our findings support previous reports that $E_2$ has a detrimental effect on male health and actually enhances the risk of CVD, possibly by decreasing the concentrations of apo AI. Although the mechanism by which $E_2$ increases apo AI concentrations in females but decreases it in males is not yet known, the data presented here help us to understand a fundamental aspect of apo AI gene expression. These data will be useful in the development of strategies to reduce the hypercholesterolaemic component of CVD. For example, because $E_2$ decreases male apo AI, a specific antioestrogen treatment that inhibits the effect of $E_2$ on hepatic apo AI expression might have a beneficial effect on CVD risk in the male.

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