Sex hormone modulation of proinflammatory cytokine and C-reactive protein expression in macrophages from older men and postmenopausal women

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Abstract

Inflammation plays a central role in the development and progression of coronary heart disease (CHD). The sex hormones estrogen and testosterone have been shown to modify the inflammatory response by influencing cytokine expression in human macrophages obtained from younger individuals. The effect of these hormones on the expression of proinflammatory markers in macrophages obtained from a CHD age-relevant population has not been studied. Human monocyte-derived macrophages (HMDMs) were obtained from healthy normolipidemic men and postmenopausal women (age 50–70 years), and cultured in autologous serum along with both physiological and supraphysiological concentrations of estrogen or testosterone. HMDMs were stimulated with oxidized low-density lipoproteins, and the expression of the cytokines tumor necrosis factor \( \alpha \) (TNF-\( \alpha \) or TFN), interleukin (IL)6, and IL-1\( \beta \) (IL1B) and of the acute-phase protein C-reactive protein (CRP) was measured. Both physiological and supraphysiological concentrations of testosterone reduced the expression and secretion of TNF-\( \alpha \) and reduced the expression of IL-1\( \beta \), but did not affect the expression of IL6 or CRP. Estrogen did not modify the expression of TNF-\( \alpha \), IL6, and IL-1\( \beta \). Estrogen caused a variable response in CRP expression that was positively associated with the plasma small dense LDL-cholesterol concentration of the donors. There were no gender differences in any of the observed effects. Our results indicate that testosterone may exert anti-inflammatory effects by reducing macrophage TNF-\( \alpha \) expression, while the effects of estrogen on macrophage CRP expression may depend upon the extracellular lipid environment.


Introduction

The pathology of coronary heart disease (CHD) is complex and multifaceted, with inflammation playing a central part (Libby 2002). Peripheral blood monocytes recruited into the intima–media layer of an artery play a pivotal role in the local content is locally produced (Kobayashi et al. 2001). Most circulating CRP is secreted by the liver, yet a small amount of CRP is produced by macrophages present in atherosclerotic plaques (Yasojima et al. 2001). Levels of CRP mRNA and protein have been found to be up to tenfold higher in arterial plaque tissue than in the normal artery, suggesting that at least a portion of atheroma CRP content is locally produced (Kobayashi et al. 2003). In vivo and animal studies indicate that CRP may actively participate in plaque development by promoting endothelial dysfunction (Bisogni et al. 2007), monocyte adhesion to the endothelium (Li et al. 2004), macrophage cholesterol accumulation (Singh et al. 2008), and fibrin breakdown (reduced plaque stability; Williams et al. 2004).

The steroid hormones 17\( \beta \)-estradiol (E\(_2\)) and testosterone are thought to play a role in modulating inflammation and thereby influencing atherogenesis. In postmenopausal women, E\(_2\) replacement therapy does not affect circulating TNF-\( \alpha \) and IL6 concentrations (Pradhan et al. 2002, Zegura et al. 2003). However, hormone treatment may cause changes in cytokine levels in the arterial lesions that may not be

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entirely predicted by plasma cytokine levels. Very little work has been conducted examining the effect of E2 on cytokine expression in human monocyte-derived macrophages (HMDMs), with one study showing that E2 withdrawal results in greater proinflammatory cytokine expression in female premenopausal HMDMs after 24 h of treatment than in HMDMs continually exposed to the hormone for 48 h (Kramer et al. 2004). Using testosterone, the expression of TNF-α, IL6, and IL-1β was shown to be reduced both in rodent macrophage cell models and in human monocytes obtained from younger individuals (Chao et al. 1995, Kanda et al. 1996, 1997, D’Agostino et al. 1999); yet, no studies have been done in HMDMs obtained from older individuals. The effect of E2 and testosterone on CRP expression in HMDMs has not been studied. The purpose of this study was to assess the effect of E2 and testosterone treatment on the expression of proinflammatory cytokines and CRP by macrophages obtained from a CHD age–relevant population.

Materials and Methods

Materials

E2 and testosterone were purchased from Sigma–Aldrich. Phenol-free RPMI1640 medium was purchased from Gibco. Ficoll-Paque was obtained from GE Healthcare (Piscataway, NJ, USA). RNeasy mini kit was purchased from Qiagen. Penicillin, streptomycin, and SuperScript III Reverse Transcriptase kit were obtained from Invitrogen. Power SYBR Green Master Mix was purchased from Applied Biosystems (Carlsbad, CA, USA). Bicinchoninic Acid Protein Assay kit was obtained from Pierce (Rockford, IL, USA).

Subjects

Male (n=10) and postmenopausal female (n=10) volunteers between 50 and 70 years of age were recruited. Subjects were included if they reported no history of CHD; cancer; diabetes; or renal, liver, or thyroid disease. Subjects who smoked or had hypertension were excluded from the study. Volunteers were not taking any medications to control plasma lipid or glucose levels. Inclusion criteria were low-density lipoprotein cholesterol (LDL-C) <160 mg/dl, high-density lipoprotein cholesterol (HDL-C) ≥40 mg/dl, triglycerides (TG) <150 mg/dl, and glucose ≤100 mg/dl. Women were considered postmenopausal if absence of menstrual periods exceeded 1 year. Most women (n=8) in this study had been postmenopausal for >5 years. Lastly, because it had been reported that the estrogen receptor α (ERα or ESR1) gene polymorphism IVS1-401 T/C, located within the first intron, can affect the plasma lipid response to E2 (Herrington et al. 2002), volunteers were genotyped for this mutation, and subjects with the IVSI C/C genotype were excluded. Characteristics of the subjects are given in Table 1.

<table>
<thead>
<tr>
<th>Age (years)</th>
<th>Women (n=10)</th>
<th>Men (n=10)</th>
<th>P value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>TC (mg/dl)</td>
<td>206 (31)</td>
<td>178 (34)</td>
<td>0.069</td>
</tr>
<tr>
<td>LDL-C (mg/dl)</td>
<td>111 (33)</td>
<td>87 (36)</td>
<td>0.136</td>
</tr>
<tr>
<td>sdLDL-C (mg/dl)</td>
<td>26 (8)</td>
<td>29 (13)</td>
<td>0.555</td>
</tr>
<tr>
<td>HDL-C (mg/dl)</td>
<td>66 (16)</td>
<td>46 (16)</td>
<td>0.011*</td>
</tr>
<tr>
<td>TG (mg/dl)</td>
<td>78 (28)</td>
<td>99 (46)</td>
<td>0.229</td>
</tr>
<tr>
<td>Glucose (mg/dl)</td>
<td>91 (7)</td>
<td>91 (8)</td>
<td>0.976</td>
</tr>
<tr>
<td>Plasma CRP (µg/ml)</td>
<td>3.9 (3.6)</td>
<td>1.4 (0.91)</td>
<td>0.053</td>
</tr>
<tr>
<td>ERα IVS1</td>
<td>T/C=5, T/T=5</td>
<td>T/C=8, T/T=2</td>
<td>0.349</td>
</tr>
</tbody>
</table>

*P value for gender difference.

Plasma lipid measurements and LDL isolation and oxidation

Plasma lipid levels were determined using enzymatic assays (Roche Diagnostics). Plasma levels of small dense LDL-C (sdLDL-C) were assessed using an enzymatic assay (Denka Seiken Corporation, Tokyo, Japan) as described previously (Ai et al. 2008). Plasma CRP levels were measured using a high-sensitivity immunoturbidimetric assay (Roche Diagnostics).

LDLs were isolated from the pooled plasma of the donors by rapid single-step ultracentrifugation using a Beckman NVT90 rotor as described previously (Vieira et al. 1996). After desalting, LDLs were oxidized by the addition of 100 µM CuSO4/100 µg protein. Oxidation extent was monitored by the formation of conjugated dienes at 234 nm. When absorbance began to increase exponentially (≈1.5 h), LDLs were placed on ice and immediately desalted using chromatography columns (Bio-Rad Laboratories) to stop further oxidation. This typically produced a thiobarbituric acid-reactive substance (TBARS) value of 6–8 nM malondialdehyde/µg protein. TBARS measurements were performed as described previously (Cathcart et al. 1991). The moderately oxidized LDLs (oxLDL) were stored at −80 °C in the dark for up to 2 months, as TBARS values and 234 nm readings were found to remain stable for this duration. The same batch of oxLDLs was used for all the experiments.

Isolation and culture of HMDMs

Blood was drawn in tubes containing 0-1% EDTA and centrifuged at 250 g for 30 min (25 °C) to remove plasma. Buffy coats were obtained by layering blood diluted 1:2 with RPMI culture medium over Ficoll-Paque followed by centrifugation (37 min, 500 g, 25 °C). White blood cells were collected and washed twice with RPMI medium. Cells were plated in RPMI medium containing 100 U/ml penicillin and 100 µg/ml streptomycin. After 3–5 h of incubation at 37 °C in 5% CO2, nonadherent cells were washed off, and the remaining monocytes were cultured in
RPMI medium containing 10% autologous serum. Cells were allowed to differentiate for 10 days in the presence of vehicle (ethanol), E2 (2 or 20 nM), or testosterone (10 or 100 nM), with the medium being changed every 3–4 days. E2 and testosterone solutions were prepared fresh under sterile conditions every 2 weeks by dissolving in 100% ethanol, and were stored in the dark at −80 °C. HMDMs were treated with 50 μg/ml oxLDL for 48 h (days 11–12) in the presence of 10% autologous serum and hormone, and were then exposed to hormones in addition to the medium without serum for an additional 24 h. The serum-free medium was collected, centrifuged to remove cell debris, and stored at −80 °C. Cells were collected in a lysis buffer (0.1 M KH₂PO₄, 0.05 M NaCl, 5 mM cholic acid, and 0.1% Triton X-100), and the cell protein was quantified using the bicinchoninic acid method with BSA as a standard.

Real-time PCR

Total cellular RNA was isolated using the RNeasy mini kit according to the manufacturer’s instructions. Hundred nanograms of RNA were reverse transcribed using the SuperScript III Reverse Transcription kit, and were amplified on a real-time PCR system 7300 using specific primers. The primer sequences were as follows: β-actin (F′-TGAAGTTGTGACGTGGACATCC and R′-CTCAGGAGGAGGCAATGATCTTG), TNF-α (F′-TGGAAGAGGTTGACCGACTC and R′-TCCCTCACAGGGGAC), IL-6 (F′-GTGGCTGCAGGACATGACAA and R′-TGAGGTTGCCCATGCTACATTT), IL-1β (F′-TTATTACAGTGCAATGAGGATGAC and R′-CGCCATCCAGAGGGCAG), CRP (F′-ATTCAGGCCCTTGT-ATCAGGGAC), androgen receptor (AR) (F′-GACTCCGTGCAGCCTATTGC and R′-GCGAGTCTCCTTGGCAG), and estrogen receptor (ER) or ESR1 (F′-CGGCATTCTACAGGCAA and R′-GGGAGCCACACTTCACC), and ERβ or ESR2 (F′-TACAAATCGATAAAACCGGGA and R′-GGGAGCCACACTTCACC). Primers were designed using Primer Express software (Foster City, CA, USA), included intron/exon boundaries, and were validated for efficiency and specificity using the standard curve dilution and melting point analyses. Real-time PCR was carried out for 40 cycles at 95 °C (15 s) + 60 °C (1 min) using the Power SYBR Green Master Mix. Changes in gene expression were assessed by ΔΔCt analysis with β-actin as the control/housekeeping gene. Changes were expressed as the percent of control (vehicle only).

TNF-α ELISA

TNF-α concentration in the cell culture medium was measured using an ultrasensitive human TNF-α ELISA kit according to the manufacturer’s instructions (Alpco Diagnostics, Salem, NH, USA). The final TNF-α concentrations were adjusted for cell protein.

Figure 1 Effect of E2 or testosterone treatment on proinflammatory cytokine expression in HMDMs. HMDMs were cultured in 10% autologous serum and differentiated over 10 days in the presence of vehicle (control), 2 nM E2, 20 nM E2, 10 nM testosterone, or 100 nM testosterone. HMDMs were then exposed for 48 h to oxLDLs with the hormone. RNA was isolated, and the expression of proinflammatory cytokines was measured. Data are expressed as the percent of control; *P<0.05.
Statistical analyses were performed using SAS software (version 9.1; SAS Institute, Cary, NC, USA), while correlations were determined using GraphPad Prism software (version 4; La Jolla, CA, USA). Gene expression results are expressed as the percent of control (vehicle treatment only). Means and S.D. are representative of the treatment response in macrophage cultures donated by ten females or ten males. Statistical differences were determined by two-way ANOVA for both treatment effect and sex effect using Tukey’s Student Range test.

Results

Both hormone receptors AR and ERα were expressed in HMDMs (average Ct of 33 and 32 respectively, on real-time PCR, compared with an average Ct of 24 for β-actin). There was no difference in the expression of AR between female and male HMDMs, while the expression of ERα was threefold higher in female HMDMs than in male HMDMs. The expression of ERβ was too low to be accurately quantified (Ct > 36).

E2 treatment did not significantly affect the expression of TNF-α, IL6, and IL-1β in HMDMs (Fig. 1). Relative to control, both physiological and pharmacological concentrations of testosterone significantly reduced TNF-α expression in HMDMs obtained from both males and females (Fig. 1). These reductions averaged 20–25% at 10 nM and 25–30% at 100 nM (Fig. 1). TNF-α concentration in the medium was measured in the vehicle-, 2 nM E2-, and 10 nM testosterone-treated cells (Fig. 2). Similar to the effect on gene expression, testosterone treatment, but not E2 treatment, significantly reduced HMDM TNF-α secretion, compared with the control. There was no statistical difference between the genders. Testosterone treatment significantly reduced IL-1β expression at 10 nM in males and at 100 nM in females, but it did not affect IL6 expression (Fig. 1).

The effect of E2 on CRP expression was variable in both women and men (Fig. 3). The change in CRP expression by HMDMs in response to E2 treatment, but not in response to testosterone treatment, was positively associated with the concentration of sdLDL-C in the plasma of the donors (Fig. 4A and B). Donors with plasma sdLDL-C > 30 mg/dl showed a significant increase in HMDM CRP expression with estrogen treatment, whereas there was no effect in HMDMs obtained from donors with sdLDL-C < 30 mg/dl (Fig. 4C). This effect was gender independent. The change in CRP expression with E2 was not associated with other parameters (age, body mass index, other plasma lipid, or CRP levels).

Discussion

The effect of sex hormones on macrophage inflammation is an important area of investigation because of the contribution of inflammation to CHD. The hormone receptors AR and ER, which are expressed in both male and female HMDMs, are the likely mediators of the effects of sex hormones on macrophage function (Cutolo et al. 1996). Low serum testosterone levels in men have been associated with increased atheroma formation (Hak et al. 2002). In addition, hypogonadal men have been found to have higher serum cytokine levels than healthy men, and androgen supplementation was found to reduce these levels (Yesilova et al. 2000). Testosterone treatment has been shown to suppress cytokine expression in rodent macrophages and in human macrophages from healthy humans.
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Figure 4 Association between fasting plasma sdLDL-C concentration of the donors and HMDM CRP gene expression in response to E₂. (A) Association between the change in CRP expression during 2 nM E₂ treatment, relative to the control (vehicle), and the sdLDL-C levels of the donors. Both the correlations were significant (P<0.01) and were gender independent. (C) Effect of E₂ on CRP expression as a function of the plasma sdLDL-C level of the donors. The number of subjects with sdLDL-C<30 mg/dl is 12, and the number of subjects with sdLDL-C>30 mg/dl is 8. Data are expressed as the percent of control; *P<0.05.

monocytes obtained from younger individuals (Chao et al. 1995, Kanda et al. 1996, 1997, D’Agostino Milano et al. 1999). Therefore, we hypothesized that similar changes would occur in HMDMs obtained from older individuals. In cells that were exposed to moderately oxidized LDLs, testosterone significantly reduced proinflammatory cytokine expression, specifically TNF-α and IL-1β expression, in agreement with the previous studies (Chao et al. 1995, Kanda et al. 1996, 1997, D’Agostino Milano et al. 1999). The effects of testosterone on TNF-α and IL-1β were observed at physiological and supraphysiological concentrations. Macrophages express the enzyme aromatase, and are therefore able to convert testosterone to E₂ (Mor et al. 2001). Since reductions in TNF-α and IL-1β were only observed with testosterone, it is reasonable to assume that the effect observed with testosterone treatment was not due to aromatization to E₃, but due to an androgen-specific action of testosterone.

NF-κB (NFkB), a transcription factor that plays a direct role in the expression of numerous proinflammatory cytokines including TNF-α and IL-1β (Li & Verma 2002), may be the mediator of the repression of cytokine expression by testosterone. Several studies have shown that AR activation suppresses NF-κB activity (Hatakeyama et al. 2002, Libby 2002, Itoh et al. 2007), presumably by increasing the expression of IκB-α (NFKBLA; Death et al. 2004). NF-κB inhibition by testosterone is also associated with enhanced macrophage apoptosis (Cutolo et al. 2005), suggesting that the reduction in TNF-α expression is accompanied by an immunosuppressive effect.

The effect of E₂ on cytokine expression is less clear. Short-term exposure to E₂ (<1 h) in vitro has been reported to decrease the production of proinflammatory cytokines in lipopolysaccharide (LPS)-activated rodent macrophages (Ghisletti et al. 2005). In contrast, several studies have reported that long-term exposure to E₂ in vivo enhances the proinflammatory cytokine production in LPS-activated rodent macrophages (Soucy et al. 2005, Calippe et al. 2008). In our study, in which HMDMs obtained from 50- to 70-year-old male and postmenopausal female donors were cultured in the continual presence of E₂, we observed no effect on proinflammatory cytokine expression compared with the vehicle-treated cells.

Randomized, placebo-controlled trials in postmenopausal women have shown that orally delivered, but not transdermally delivered, estrogen therapy increases plasma CRP concentrations (Cushman et al. 1999, Hodis et al. 2008). It has been speculated that hepatic metabolism of the orally delivered therapy is responsible for this rise in plasma CRP levels (Zegura et al. 2003). However, since CRP is also produced by macrophages in the aortic lesions, plasma CRP level changes in response to E₂ treatment may not predict CRP changes in the arterial wall. Because CRP has been shown to elicit proatherogenic effects such as the promotion of macrophage cholesterol accumulation (Singh et al. 2008), CRP secreted by macrophages may play a role in lesion development. The CRP expression in response to E₂ treatment was quite variable in both female and male
HMDM donors. It has been shown previously that the lipoprotein composition of culture serum may influence the cellular response (de la Llera Moyet al. 1994). Since HMDMs were cultured in 10% autologous serum, we tested the hypothesis that changes in CRP expression by E2 may be dependent on the plasma levels of lipoproteins of the donors. The CRP gene response to E2 was significantly correlated with plasma sLDL-C levels, with a greater increase in CRP expression in subjects with higher sLDL-C levels. Individuals with a high concentration of sLDL-C often display some degree of dyslipidemia and a greater degree of chronic inflammation and CHD (Vakkilainen et al. 2003, Krauss & Siri 2004). This indicates that in a proatherogenic lipid environment, E2 may promote arterial disease, an effect that is not present under a healthy lipid environment. This concept is supported by clinical evidence suggesting that E2 therapy is beneficial in younger premenopausal women, but harmful in older postmenopausal women (Rossouw et al. 2007). In support of our findings, a recent study done by Norata et al. (2009) has shown a greater expression of several proinflammatory molecules in HMDMs obtained from individuals with high sLDL-C levels than in those obtained from individuals with lower sLDL-C levels. Furthermore, in postmenopausal women, plasma sLDL-C levels have been found to be significantly associated with plasma CRP concentration, further indicating the link between sLDL-C and inflammation (Muzzio et al. 2007). We did not observe a significant correlation between these two factors in our study (data not shown). However, this may be due to the small sample size.

To our knowledge, our study is the first to report a modulation of CRP expression by E2 in HMDMs. The mechanism by which high sLDL-C levels may alter the effect of E2 on CRP expression is not currently known. IL6 and IL-1β are the known stimulators of CRP expression (Calabro et al. 2003). In the current study and in the study done by Norata et al. (2009), sLDL-C concentrations were not associated with HMDM IL6 expression. Similarly, the expression of IL6 and IL-1β did not change with E2 treatment. Therefore, the observed up-regulation of macrophage CRP expression by E2 is not driven by greater macrophage IL6 or IL-1β expression. An IL6-independent stimulation of CRP by hormone replacement has also been suggested by observational and intervention studies (Lakoski & Herrington 2005). The expression of Stat3, a transcription factor that regulates the expression of Cyp, has been shown to be increased by estrogen in the ob/ob mouse model of obesity, possibly through E2 binding to the promoter region of Stat3 (Gao et al. 2006). How sLDLs may modulate the effect of E2 on CRP expression is currently not known. sLDLs are known to be in circulation longer than LDLs, and therefore more likely to undergo oxidation (Millar & Packard 1998).

Overall, these data suggest that testosterone may protect against the progression of atherosclerosis by inhibiting the expression of select proinflammatory cytokines in human macrophages, while E2 may not be as potent in this regard. Furthermore, E2 may actually encourage macrophage CRP production under conditions of high sLDL-C, thereby potentially exacerbating atherosclerosis in individuals at risk of the disease. Understanding the mechanism of sLDL-C-induced macrophage inflammation and how E2 modulates this is an important step in defining the role of both sLDL-C and E2 in the progression of CHD.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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