Differential Effects of Oral and Transdermal Estrogen Replacement Therapy on Endothelial Function in Postmenopausal Women

Satu Vehkavaara, MD; Tiina Hakala-Ala-Pietilä, MD; Antti Virkamäki, MD; Robert Bergholm, MD; Christian Ehnholm, MD; Outi Hovatta, MD; Marja-Riitta Taskinen, MD; Hannele Yki-Järvinen, MD

Background—We determined whether the vascular effects of estradiol depend on the route of administration by comparing the effects of oral estradiol and transdermal placebo, transdermal estradiol and oral placebo, and transdermal placebo and oral placebo on in vivo endothelial function in 27 postmenopausal women.

Methods and Results—Endothelial function was assessed from blood flow responses to intrabrachial artery infusions of endothelium-dependent (7.5 and 15 μg/min acetylcholine) and endothelium-independent (3 and 10 μg/min of sodium nitroprusside) vasodilators at 0, 2, and 12 weeks. In the oral estradiol group, the increase in flow above basal during infusion of the low dose of acetylcholine at 0, 2, and 12 weeks averaged 6.0±0.8, 6.9±0.8, and 11.3±1.2 (P<0.01 versus 0 and 2 weeks) mL·dL⁻¹·min⁻¹ at 0, 2, and 12 weeks. The percentage increases versus 0 weeks averaged 21±14% at 2 and 120±34% at 12 weeks. During the high-dose acetylcholine infusion, the increase in flow above basal averaged 8.6±1.3, 10.2±1.5, and 15.1±1.8 (P<0.05 versus 0 weeks) mL·dL⁻¹·min⁻¹, respectively. The percentage increases versus 0 weeks averaged 22±10% at 2 weeks and 119±46% at 12 weeks. In the oral estradiol group, endothelium-independent vasodilatation also improved significantly, but less markedly than endothelium-dependent responses. In the transdermal and placebo groups, all vascular responses remained unchanged. Oral but not transdermal estradiol also induced significant decreases in LDL cholesterol and Lp(a) concentrations and an increase in HDL cholesterol within 2 weeks.

Conclusions—We conclude that oral but not transdermal estradiol induces potentially antiatherogenic changes in vivo endothelium-dependent vasodilatation and lipid concentrations. (Circulation. 2000;102:2687-2693.)

Key Words: atherosclerosis ■ blood flow ■ endothelium

Observational studies suggest that postmenopausal hormone replacement therapy, including various estrogen preparations with or without progesterin, approximately halves the risk of cardiovascular disease in women. A recent prospective study in postmenopausal women with coronary artery disease failed, however, to demonstrate benefits of estrogen plus progesterin on cardiovascular events. Whether the possible cardioprotective effects of estrogen preparations depend on the route of estrogen administration is currently unknown.

Several mechanisms have been suggested to contribute to the cardiovascular effects of oral estradiol. These include oral estradiol–induced increases in HDL cholesterol and decreases in LDL cholesterol and lipoprotein (a) [Lp(a)] concentrations. Compared with oral estradiol, transdermal estradiol appears to have no or only marginal effects on plasma lipid and lipoprotein concentrations. Estrogen replacement therapy could also directly affect blood vessels. In postmenopausal women, short-term infusions of estradiol potentiate endothelium-dependent vasodilatation in coronary arteries and forearm resistance vessels, but it is unclear whether this acute estradiol effect is responsible for enhanced endothelium-dependent function during long-term therapy. In none of the previous studies have effects of oral and transdermal estradiol on endothelial function been compared, and no study addressed the question of whether the acute effects of estradiol are responsible for enhanced endothelial function during long-term therapy. Because both oral and transdermal estradiol induce stable serum estradiol concentrations within a few days, we reasoned that if direct vascular effects of estradiol were to be responsible for improvements in endothelial function, maximal enhancement should be observed already after 2 weeks, with no further improvement during continued therapy in either group.
In the present study, we compared the effects of oral and transdermal estradiol on endothelial function in vivo. We chose the clinically most commonly used doses of estradiol, a patch delivering 50 μg/d and a daily 2-mg estradiol tablet, which both effectively relieve postmenopausal symptoms\(^\text{11}\) and were estimated to increase peripheral free estradiol concentrations similarly. A total of 27 healthy postmenopausal women were randomized to receive either oral or transdermal estradiol or placebo, and they underwent invasive measurements of endothelial function after 0, 2, and 12 weeks of treatment.

### Methods

#### Subjects and Study Design

**Screening Visit 1 (Internist)**

Postmenopausal women were recruited by a newspaper advertisement. To be acceptable for the study, the subjects had to fulfill the following inclusion criteria: (1) natural amenorrhea for \(\geq 12\) months or age \(\geq 52\) years, (2) follicle-stimulating hormone (FSH) \(>30\) U/L, (3) no contraindications for estrogen treatment, (4) no evidence of any disease on the basis of history and physical examination and standard laboratory tests (blood counts, serum creatinine, electrolyte concentrations, liver function tests, and ECG), and (5) use of no medications, including vitamins and antidepressants.

**Screening Visit 2 (Gynecologist)**

A total of 36 women visited the gynecologist. A gynecological history was obtained, and a gynecological examination, including transvaginal ultrasound, was performed. To be included in the study, the patients had to fulfill the following criteria: (1) no suspicion of malignancy, (2) no myomas \(>4\) cm, (3) no endometrial pathology, (4) endometrial thickness \(<6\) mm, (5) and normal ovaries and breasts.

Of the 36 women attending the first gynecological examination, 4 had bleeding after the first screening visit and were excluded from the study. Another 3 patients were excluded from the study because of myomas, and 2 patients did not wish to continue the study. The experimental protocol was designed and performed according to the principles of the Helsinki Declaration and was approved by the ethical committee of the Helsinki University Central Hospital.

**Randomization**

Eligible patients were randomly assigned to 1 of 3 groups (Table 1) by use of minimization of differences between the treatment groups as the method of randomization.14 The first group used an oral estradiol tablet of 2 mg (Estrofem, Novo Nordisk) and a placebo patch, the second group transdermal estradiol 50 μg/d (Menorest, Rhône-Poulenc Rorer) and a placebo tablet, and the third group a placebo patch and a placebo tablet for 12 weeks. Measurements of endothelial function in vivo and circulating lipid and hormone concentrations were performed at 0, 2, and 12 weeks as detailed below.

### In Vivo Endothelial Function Tests at 0, 2, and 12 Weeks

Endothelial function was assessed in forearm resistance vessels after an overnight fast by measurement of forearm blood flow responses to intra-arterial infusions of endothelium-dependent (acetylcholine, ACh) and endothelium-independent (sodium nitroprusside, SNP) vasodilators, as previously described in detail.15 Current smokers refrained from smoking for 12 hours before the study. An indwelling cannula was inserted in an antecubital vein for blood sampling. A 27-gauge unmounted steel cannula (Coopers Needle Works) connected to an epidural catheter was inserted into the left brachial artery. All drugs were infused at a constant rate of 1 mL/min in the following sequence: normal saline, SNP (Nitropress, Abbott Laboratories) 3 (low dose) and 10 (high dose) μg/min, and ACh (Miochol, OMJ Pharmaceuticals) 7.5 (low dose) and 15 (high dose) μg/min. Each dose was infused for 6 minutes, and infusion of each drug was separated by infusion of normal saline for 18 minutes, during which time blood flow returned to basal values. Forearm blood flow was recorded simultaneously in the infused (experimental) and control arms as previously described.16 The percentage changes in blood flow responses induced by treatment were calculated from the mean±SEM of individual changes, which were defined as 100×(flow after−flow before)/flow before treatment.

### Other Measurements

Serum total cholesterol and triglycerides and LDL and HDL cholesterol concentrations were measured as previously described.16 Serum concentrations of Lp(a) were determined by use of the Pharmacia Apolipoprotein (a) (RIA assay system, FSH and sex hormone-binding globulin (SHBG) (AutoDELFIA SHBG, Wallac) by fluoroenzymometric assays. Serum estradiol (Estradiol-2, Sorin Biomedica), estrone17 (Estrone-RIA, Bühlman), and testosterone

<table>
<thead>
<tr>
<th>TABLE 1. Baseline Characteristics of the Groups</th>
</tr>
</thead>
<tbody>
<tr>
<td>Oral Estradiol Group (n=9)</td>
</tr>
<tr>
<td>Age, y</td>
</tr>
<tr>
<td>Weight, kg</td>
</tr>
<tr>
<td>BMI, kg/m²</td>
</tr>
<tr>
<td>Lean body mass, kg</td>
</tr>
<tr>
<td>Body fat, %</td>
</tr>
<tr>
<td>Waist/hip ratio</td>
</tr>
<tr>
<td>SBP, mm Hg</td>
</tr>
<tr>
<td>DBP, mm Hg</td>
</tr>
<tr>
<td>Smoking, +/ex/−</td>
</tr>
<tr>
<td>Age at menopause, y, mean (range)</td>
</tr>
<tr>
<td>Hysterectomized, %</td>
</tr>
<tr>
<td>Serum FSH, IU/L</td>
</tr>
</tbody>
</table>

BMI indicates body mass index; SBP, systolic blood pressure; DBP, diastolic blood pressure; +, current smoker; ex, ex-smoker; and −, never smoked.
Serum total estradiol concentrations were below the limit of detection in the placebo group (<20 pmol/L) at 0, 2, and 12 weeks. In the oral estradiol group, serum total estradiol concentrations increased from <20 pmol/L at baseline to 378±249 at 2 weeks and 423±45 pmol/L at 12 weeks (P<0.01 for treatment effect). In the transdermal estradiol group, serum total estradiol concentrations increased from <20 pmol/L at baseline to 156±26 at 2 weeks and 216±31 pmol/L at 12 weeks (P<0.001 for treatment effect). Serum total estradiol concentrations were significantly higher in both the transdermal (P<0.05 and P<0.01 at 2 and 12 weeks versus placebo) and oral (P<0.001 at 2 and 12 weeks versus placebo) estradiol groups than in the placebo group and higher in the oral than the transdermal estradiol group at 2 (P<0.001) and 12 (P<0.01) weeks. Serum SHBG concentrations remained unchanged in the placebo (49±10 versus 48±9 nmol/L, 0 versus 12 weeks) and the transdermal estradiol (58±4 versus 65±7 nmol/L, 0 versus 12 weeks) groups but increased in the oral estradiol group by 133%, from 72±11 to 168±11 nmol/L at 12 weeks (P<0.001, Figure 1). Because of the increase in serum SHBG concentrations, free estradiol concentrations were similar in the oral (3.17±0.36 pmol/L) and transdermal (3.09±0.49 pmol/L) groups at 12 weeks (Figure 1). Serum estrone concentrations remained unchanged in the placebo and transdermal estradiol groups (Figure 2) but increased >10-fold in the oral estradiol group, from 238±34 pmol/L at baseline to 2727±316 pmol/L at 2 weeks and 2947±421 pmol/L at 12 weeks (P<0.001 for treatment effect). Serum total testosterone concentrations remained unchanged in the oral (0.99±0.02 versus 0.74±0.05 nmol/L, 0 versus 12 weeks), transdermal (1.13±0.17 versus 1.23±0.13, respectively), and placebo (1.07±0.12 versus 1.01±0.09, respectively) groups. Also due to an increase in serum SHBG concentrations, serum free testosterone concentrations decreased in the oral estradiol group from 11±1 pmol/L at baseline to 4±1 pmol/L at 12 weeks (P<0.001) and remained unchanged in the transdermal (14±2 versus 14±2 pmol/L, respectively) and placebo (15±2 versus 14±2 pmol/L, respectively) groups.

Serum Lipids and Lipoproteins and Apoproteins
In the oral estradiol group, LDL cholesterol decreased by 9%, from 3.6±0.2 mmol/L at baseline to 3.2±0.1 mmol/L (P<0.05) at 2 weeks, and remained at this level until 12 weeks. Also in the oral estradiol group, HDL cholesterol increased by 19% at 2 weeks and remained increased until 12 weeks. Lp(a) decreased significantly in the oral estradiol group at 2 and 12 weeks (P<0.01 versus baseline). In the transdermal and placebo groups, all lipid concentrations remained unchanged for the 12-week period (Table 2).

Endothelial Function
In the oral estradiol group, basal flow increased slightly, although not significantly, during the 12-week treatment period from 1.5±0.3 to 1.7±0.2 mL·min⁻¹ at 2 weeks (P=NS) and to 2.1±0.1 mL·min⁻¹ at 12 weeks (P<0.10). Total (basal and ACh-stimulated) flow during infusion of the low dose of ACh increased from 7.6±0.9 at 0 weeks to 8.9±0.9 mL·min⁻¹ at 2 weeks and by 92±26% to 13.0±1.1 mL·min⁻¹ at 12 weeks (P<0.01 versus 0 and 2 weeks).
by guest on April 18, 2017 http://circ.ahajournals.org/ Downloaded from
dose of ACh averaged 6.0±0.8, and 11.3±6.6 versus 0 and 2 weeks) (Figure 3). In the oral estradiol group, 634% at 12 weeks during the low-dose at 2 weeks and 120±10% to 17.9±110 by 110±6 versus 0 weeks) and high-dose (8.8±0.9 versus 10.6±0.8 versus 13.5±11.1 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P<0.05 for 12 versus 0 weeks) SNP infusions in the oral estradiol group. The percentage increases in flow above basal compared with 0 weeks averaged 16±12% and 50±18% at 2 and 12 weeks during the low-dose SNP infusion and 28±12% and 64±19% during the high-dose SNP infusion in the oral estradiol group. In the transdermal group, there were no significant changes in total flows during infusion of the low-dose (8.2±0.6 versus 10.2±0.9 versus 9.3±0.7 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) or the high-dose (10.8±1.0 versus 13.1±1.8 versus 13.1±1.2 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) SNP (Figure 3). Similarly, in the placebo group, total flow remained unchanged during infusion of both the low-dose (10.8±1.7 versus 9.1±1.2 versus 10.1±1.2 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) and the high-dose (13.6±2.3 versus 11.7±1.8 versus 12.2±1.7 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) SNP (Figure 3).

To determine the possible causes of enhanced endothelial function during estrogen therapy, simple correlation coefficients (Spearman) were calculated between changes in metabolic parameters and those of endothelium-dependent and -independent vasodilatation. None of the lipid or hormone concentrations correlated with measures of blood flow within the individual groups (data not shown). In the oral estradiol group, the correlation coefficient between the change in SHBG and the change in the blood flow response to the high-dose ACh was 0.60 (P=0.10, NS). Figure 2 shows estradiol and estrone concentrations and the change in endothelium-dependent vasodilatation as a function of time. Neither the time course nor the fold change in endothelium-dependent vasodilatation paralleled the hormone concentrations, implying that short-term effects of estradiol or estrone were not responsible for improved endothelium-dependent vasodilatation in the oral estradiol group.

**Discussion**

The present study is, to the best of our knowledge, the first to compare the effects of oral and transdermal estradiol and of

---

**Figure 2.** Serum estradiol (top) and estrone (middle) concentrations at 0, 2, and 12 weeks. Bottom, % change in endothelium-dependent vasodilatation during infusion of ACh 15 μg/min at 2 and 12 weeks vs 0 weeks. Solid circles represent oral estradiol (ORAL); open circles, transdermal estradiol (TRANS); and open triangles, control (CONT) group. Top and middle: P<0.001 (ANOVA for repeated measures for oral vs transdermal and placebo groups); bottom, P<0.01 (ANOVA for repeated measures for oral vs transdermal and placebo groups). **P<0.05, **P<0.01, and ***P<0.001 for pairwise comparisons between groups.

Total flow during infusion of the high dose of ACh increased from 10.2±1.4 at baseline to 12.3±1.6 mL·dL⁻¹·min⁻¹ at 2 weeks and by 110±37% to 17.9±1.8 mL·dL⁻¹·min⁻¹ at 12 weeks (P<0.01 versus 0 and 2 weeks) (Figure 3). In the oral estradiol group, the increase in flow above basal during infusion of the low dose of ACh averaged 6.0±0.8, 6.9±0.8, and 11.3±1.2 (P<0.01 versus 0 and 2 weeks) mL·dL⁻¹·min⁻¹ at 0, 2, and 12 weeks, and during the high-dose ACh infusion, 8.6±1.3, 10.2±1.5, and 15.1±1.8 (P<0.05 versus 0 weeks) mL·dL⁻¹·min⁻¹, respectively. The percentage increases in flow above basal compared with 0 weeks averaged 21±14% at 2 weeks and 120±34% at 12 weeks during the low-dose ACh infusion and 22±10% and 119±46%, respectively, during the high-dose ACh infusion.

In the transdermal estradiol group, basal flows remained unchanged (1.7±0.2 versus 1.9±0.2 versus 1.8±0.1 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS). In this group, both total blood flows during infusion of the low (9.6±1.7 versus 12.6±2.2 versus 10.7±2.3 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) and the high (11.4±1.5 versus 14.0±2.6 versus 12.4±2.0 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) doses of ACh remained unchanged (Figure 3). The same was true when flow was expressed as increase in flow above basal. In the placebo group, blood flows were comparable at 0, 2, and 12 weeks during infusion of both the low dose of ACh (data not shown) and the high dose of ACh (Figure 3).

In the oral estradiol group, total flow during infusion of the low dose of SNP averaged 8.2±0.9 at 0 and 9.3±0.8 mL·dL⁻¹·min⁻¹ at 2 weeks (P=NS versus 0 weeks) and 11.6±0.7 mL·dL⁻¹·min⁻¹ at 12 weeks (P<0.05 versus 0 weeks). Total flow during infusion of the high dose of ACh averaged 10.3±1.1 at 0 weeks and 12.4±0.9 mL·dL⁻¹·min⁻¹ at 2 weeks (P=NS versus 0 weeks) and 15.5±1.1 mL·dL⁻¹·min⁻¹ at 12 weeks (P<0.01 versus 0 and 2 weeks) (Figure 3). Results were similar when flow was expressed as flow above basal during the low-dose (6.8±0.6 versus 7.6±0.7 versus 9.5±0.8 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P<0.05 for 12 versus 0 weeks) and high-dose (8.8±0.9 versus 10.6±0.8 versus 13.5±1.1 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P<0.05 for 12 versus 0 and 2 weeks) SNP infusions in the oral estradiol group. The percentage increases in flow above basal compared with 0 weeks averaged 16±12% and 50±18% at 2 and 12 weeks during the low-dose SNP infusion and 28±12% and 64±19% during the high-dose SNP infusion in the oral estradiol group. In the transdermal group, there were no significant changes in total flows during infusion of the low-dose (8.2±0.6 versus 10.2±0.9 versus 9.3±0.7 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) or the high-dose (10.8±1.0 versus 13.1±1.8 versus 13.1±1.2 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) SNP (Figure 3). Similarly, in the placebo group, total flow remained unchanged during infusion of both the low-dose (10.8±1.7 versus 9.1±1.2 versus 10.1±1.2 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) and the high-dose (13.6±2.3 versus 11.7±1.8 versus 12.2±1.7 mL·dL⁻¹·min⁻¹, 0 versus 2 versus 12 weeks, P=NS) SNP (Figure 3).
placebo oral and transdermal preparations on endothelial function in vivo. We found that oral and transdermal estradiol have different effects on endothelial function. Oral estradiol markedly improved endothelium-dependent and, to a lesser extent, endothelium-independent blood flow in forearm resistance vessels. In contrast, transdermal estradiol therapy had no significant effects on endothelial function. Oral estradiol also induced favorable and stable lipid changes within 2 weeks, whereas transdermal estradiol had no effect on lipids or lipoproteins.

We chose the clinically most commonly used doses of estradiol, a patch changed twice a week that delivers 50 μg/d and a daily tablet containing 2 mg estradiol. Both preparations produced symptomatic relief, although serum total estradiol concentrations were 2-fold higher in the oral than the transdermal estradiol group. The measured concentrations are consistent with previous data, which generally show higher total estradiol concentrations with oral than with transdermal estradiol, despite similar clinical efficacy. In previous studies, the impact of changes in binding protein concentrations or free hormone concentrations have not been considered. This is surprising, because 2 studies have shown oral but not transdermal estradiol to increase SHBG concentrations, which will decrease free estradiol concentrations. In the present study, consistent with similar clinical efficacy, serum free estradiol concentrations, calculated on the basis of concentrations of serum SHBG, total testosterone, estrone, and estradiol, were identical between the groups. Despite this, endotholium-dependent vasodilatation increased >100% in the oral estradiol group but was virtually unchanged in the transdermal group (Figure 3). These negative data are in keeping with recent studies showing no effect of estradiol patches delivering 5023 or 100 24 μg/d on endothelial function. The latter studies, however, lasted 3 and 8 weeks, respectively, which is intermediate in time between the lack of effect of estradiol in the present study at 2 weeks and the significant effect at 12 weeks. The small, albeit significant, decrease in free testosterone concentrations could also have contributed to the beneficial effects of oral estradiol on endothelial function.

Acute administration of estradiol7–9,26 improves endothelium-dependent vasodilatation in humans. The total estradiol concentrations in the acute studies have been 3 to 4 times those of mean total estradiol concentrations of a normal menstrual cycle and those in the present study, or corresponded to maximal follicular-phase concentrations of premenopausal women. The

### Table 2. Serum Lipid and Lipoprotein Concentrations

<table>
<thead>
<tr>
<th></th>
<th>Oral Estradiol Group</th>
<th>Transdermal Estradiol Group</th>
<th>Placebo Group</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0 Wk</td>
<td>2 Wk</td>
<td>12 Wk</td>
</tr>
<tr>
<td>Total triglycerides,*</td>
<td>1.42±0.24</td>
<td>1.25±0.15</td>
<td>1.64±0.33</td>
</tr>
<tr>
<td>Cholesterol, mmol/L</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>5.61±0.23</td>
<td>5.38±0.15</td>
<td>5.60±0.12</td>
</tr>
<tr>
<td>LDL</td>
<td>3.57±0.17</td>
<td>3.19±0.08‡</td>
<td>3.28±0.14</td>
</tr>
<tr>
<td>HDL*</td>
<td>1.48±0.11</td>
<td>1.74±0.11‡</td>
<td>1.72±0.08</td>
</tr>
<tr>
<td>Lp(a),*†</td>
<td>26 (11, 30)</td>
<td>21§ (6, 23)</td>
<td>14¶ (5, 15)</td>
</tr>
</tbody>
</table>

*Log-transformed for statistical analyses.
†Median, interquartile range (25%, 75%). Other data are shown as mean±SEM.
‡P<0.05 and §P<0.01 baseline vs 2 weeks.
¶P<0.05 and ¶P<0.01 baseline vs 12 weeks.

Figure 3. Forearm blood flows (top) and % changes in flows at 2 and 12 weeks vs 0 weeks during infusions of high doses of ACh and SNP in oral (●), transdermal (○), and placebo (■) groups. *P<0.05 and **P<0.01.
The acute effects of estradiol have been observed in minutes, which makes it likely that the observed dilatatory responses have been due to estradiol per se and not to estradiol metabolites. In the present study, endothelial function was markedly improved by oral estradiol at 12 but not at 2 weeks, whereas total and free estradiol concentrations were maximal already at 2 weeks (Figure 2). This result is in keeping with data demonstrating doubling of postischemic vasodilatation between 1 and 6 months of oral hormone replacement therapy. If acute effects of estradiol alone had been responsible for the improved endothelial function, it should have improved similarly at 2 and 12 weeks. These time-course data are, to the best of our knowledge, novel and support the conclusion that changes in vascular function during estradiol replacement therapy cannot be attributed to acute vascular effects of estradiol.

Antiatherogenic changes in serum lipids by oral but not transdermal estradiol represent another possibility to explain the beneficial effects of oral estradiol on endothelial function. In the present study, oral estradiol decreased LDL cholesterol by 9% and increased HDL cholesterol by 20%. These data are in line with previous studies that reported 6% to 28% reductions in LDL cholesterol and 2% to 19% increases in HDL cholesterol with unopposed estrogen. Also, consistent with previous data, use of transdermal estradiol 100 μg/d for 6 weeks had no effect on lipid concentrations. In the study by Gerhard et al., 400 μg/wk of transdermal estradiol improved endothelium-dependent vasodilatation and also decreased LDL cholesterol by 11%. Lowering of LDL cholesterol by statins improves endothelium-dependent vasodilatation in brachial and coronary arteries. The 9% decrease in LDL cholesterol by oral estradiol in the present study is 30% to 40% of that found in the statin trials. Conversely, estradiol also clearly increased HDL cholesterol and decreased Lp(a) concentrations, both of which might have contributed to the enhancement in endothelial function.

Estradiol may also protect LDL from oxidation, a possibility not explored in the present study. Estradiol may also protect LDL from oxidation, a possibility not explored in the present study. Taken together, it is possible that the several antiatherogenic changes in lipids caused by estradiol were responsible for enhanced endothelial function. A limitation of our study is the relatively small number of subjects studied. Although we did not find statistically significant associations between changes in serum lipids or hormones and endothelial function, such could be found in a larger cohort. Therefore, these data should be interpreted with caution and reproduced.

In conclusion, 2 mg of oral estradiol markedly enhances endothelium-dependent and, to a lesser extent, endothelium-independent vasodilatation, whereas 50 μg/d of transdermal estradiol, which produces a similar increase in free estradiol, has no effect on vascular function in healthy postmenopausal women. The oral estradiol–induced increase in endothelium-dependent vasodilatation cannot be explained by acute estradiol effects. This is because serum estradiol concentrations were maximal at 2 weeks and remained at this concentration until 12 weeks, whereas endothelium-dependent vasodilatation improved significantly and by 119% only after 12 weeks. Thus, long-term effects of oral estradiol, such as the several antiatherogenic changes in lipids and lipoproteins, could be major mediators.

Acknowledgments

This study was supported by grants from the Academy of Finland (Drs Yki-Järvinen and Vehkavaara) and Liv och Hälsa (Dr Virkamäki). The authors gratefully acknowledge Kati Tuomola and Sari Haapanen for excellent technical assistance and Kaj Blomborg, PhD, for calculating serum free estradiol concentrations.

References


Differential Effects of Oral and Transdermal Estrogen Replacement Therapy on Endothelial Function in Postmenopausal Women
Satu Vehkavaara, Tiina Hakala-Ala-Pietilä, Antti Virkamäki, Robert Bergholm, Christian Ehnholm, Outi Hovatta, Marja-Riitta Taskinen and Hannele Yki-Järvinen

_Circulation._ 2000;102:2687-2693
doi: 10.1161/01.CIR.102.22.2687

_Circulation_ is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
Copyright © 2000 American Heart Association, Inc. All rights reserved.
Print ISSN: 0009-7322. Online ISSN: 1524-4539

The online version of this article, along with updated information and services, is located on the World Wide Web at:
http://circ.ahajournals.org/content/102/22/2687

Permissions: Requests for permissions to reproduce figures, tables, or portions of articles originally published in _Circulation_ can be obtained via RightsLink, a service of the Copyright Clearance Center, not the Editorial Office. Once the online version of the published article for which permission is being requested is located, click Request Permissions in the middle column of the Web page under Services. Further information about this process is available in the Permissions and Rights Question and Answer document.

Reprints: Information about reprints can be found online at:
http://www.lww.com/reprints

Subscriptions: Information about subscribing to _Circulation_ is online at:
http://circ.ahajournals.org/subscriptions/