The Selective Estrogen Receptor Modulator Acolbifene Reduces Cholesterolemia Independently of Its Anorectic Action in Control and Cholesterol-Fed Rats

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ABSTRACT The cancer-preventing selective estrogen receptor modulator (SERM) acolbifene (ACOL) exerts a potent and pure antiestrogenic action in the mammary gland and uterus, yet it displays beneficial, estrogen-like actions on energy and lipid metabolism in rodents. The compound reduces food intake and strongly decreases cholesterolemia in rats fed a cholesterol-free diet. This study was designed to establish whether the anorectic effect of ACOL is involved in its cholesterol-lowering action, and whether the compound retains its ability to lower cholesterol concentrations in rats with diet-induced hypercholesterolemia. Female rats were fed a purified diet devoid of cholesterol (reference diet) or containing 2% cholesterol (C-diet); they were either not treated or treated daily with ACOL or not treated and pair-fed to the ACOL-treated rats. The C-diet did not affect food intake or weight and fat gains. ACOL reduced food intake (16%) and weight gain (45%, mainly fat) similarly in both dietary cohorts. ACOL, but not pair feeding, reduced cholesterolemia by 33% in rats fed the reference diet. As expected, the C-diet raised serum total cholesterol almost 3-fold and this increase was largely prevented by ACOL but not by pair feeding. Cholesterol was reduced by ACOL, mainly in the HDL fraction, in rats fed the reference diet, but only in the non-HDL fraction in those fed the C-diet. In livers of rats fed the reference diet, ACOL, but not pair feeding, increased protein abundance of the scavenger receptor, class B, type 1, and the LDL receptor, thought to be involved in ACOL-mediated cholesterol lowering. These findings demonstrate that the potent hypocholesterolemic action of ACOL is independent of the concomitant reduction in food intake and fat accretion, and that such action occurs in rats with overt diet-induced hypercholesterolemia. J. Nutr. 135: 2225–2229, 2005.

KEY WORDS: • estrogen antagonist • diet-induced hypercholesterolemia • blood cholesterol • scavenger receptor B1 • LDL receptor
these receptors and lower cholesterol in the presence of diet-induced hypercholesterolemia. In addition, estrogen, the SERM tamoxifen, and the steroidal antiestrogen ICI-182,780 affect whole-body energy balance (20–22). In rats, ACOL exerts an estrogen-like effect on food intake, which in turn reduces adipose tissue accretion (9). Although such negative energy balance is more likely to affect triacylglycerol rather than cholesterol metabolism, there is no a priori reason to reject the possibility that the anorectic effect of ACOL may participate in its hypcholesterolemic action.

The present study was designed to address both of these central issues related to the potency of ACOL to reduce cholesterololemia and to its mechanisms of action. Female rats were fed a purified diet without cholesterol or the same diet to which 2% cholesterol was added. Each of these 2 dietary cohorts was subjected to ACOL treatment, or alternatively to an imposed reduction in food intake matching that of the ACOL-treated rats. The paradigm allowed the assessment of the ability of ACOL to counteract the effects of dietary cholesterol as well as the contribution of changes in ingestive behavior to its hypcholesterolemic action.

MATERIALS AND METHODS

Animals and treatments. Female Sprague-Dawley rats (n = 42) initially weighing 175–200 g were purchased from Charles River Laboratories and housed individually in stainless steel cages in a room kept at 23 ± 1°C with a 12-h light:dark cycle (lights on at 1900 h). The rats were cared for and handled in conformity with the Canadian Guide for the Care and Use of Laboratory Animals, and the protocol was approved by our institutional animal care committee. The rats were acclimated to their environment for 1 wk and consumed tap water and a nonpurified rodent diet (Charles River Rodent Diet no. 5075, Ralston Products) ad libitum. Half of the rats were then fed an obesity- and hyperlipidemia-promoting purified diet to maximize the effect of ACOL treatment. The diet provided 50% of energy as carbohydrate, 30% as fat, and 20% as protein; its composition is given in Table 1. The other half of the rats were given the same diet to which 2% (wt:wt) cholesterol (ICN Biochemicals) was added. In each of the 2 dietary cohorts, 7 rats were assigned to the placebo (vehicle) group and another 7 rats were treated with ACOL. The compound was given once daily by oral gavage at a dose of 2.5 mg/kg (vehicle) group and another 7 rats were treated with ACOL. The remaining 7 rats in each dietary cohort were not treated and were pair-fed to the ACOL-treated group in the following manner: each rat was randomly paired with one of the ACOL-treated rats, and the amount of food provided was adjusted to the amount of food ingested the previous day by the ACOL-treated counterpart. To avoid intake of the restricted amount of food in a single large meal, two thirds of the food was provided to the pair-fed rats at the beginning of the dark active period, and the remaining third was given at the beginning of the light period. The placebo and pair-fed groups were administered the methylcellulose vehicle daily by gavage. To induce body weight gain, the remaining 7 rats in each dietary cohort were not treated and were pair-fed to the ACOL-treated rats in the following manner: each rat was randomly paired with one of the ACOL-treated rats, and the amount of food provided was adjusted to the amount of food ingested the previous day by the ACOL-treated counterpart. To avoid intake of the restricted amount of food in a single large meal, two thirds of the food was provided to the pair-fed rats at the beginning of the dark active period, and the remaining third was given at the beginning of the light period. The placebo and pair-fed groups were administered the methylcellulose vehicle daily by gavage.

Blood and tissue collection. At the time of killing, blood was collected from the neck wound and was immediately centrifuged at 1500 × g for 4°C for 15 min. Serum was stored at −80°C for later biochemical measurements. The liver was excised and weighed, and a sample of liver was immediately frozen in liquid nitrogen and stored at −80°C for later determination of cholesterol content. Retropertioneal and inguinal white adipose tissues were excised and weighed.

Serum and tissue measurements. The HDL fraction was isolated by precipitation of apoB-containing lipoproteins with sodium phosphotungstate-magnesium chloride. Total and HDL cholesterol were quantified using a reagent kit (Total Cholesterol) from Wako Diagnostics. Non-HDL cholesterol was obtained by difference. Frozen liver samples were thawed; total lipids were extracted according to the method of Folch et al. (23), solubilized in isopropanol, and liver cholesterol concentration quantified in these lipid extracts using the above-mentioned reagent kits.

Western blot analysis of SR-B1 and LDLr protein. The liver content of these proteins was determined in liver extracts from rats fed the reference, cholesterol-free diet, after electrophoretic separation, by RIA using a commercially available anti-SR-B1 antibody (Novus Biologicals) and an anti-LDLr antibody kindly provided by Dr. Joachim Herz (University of Texas Southwestern Medical Center, Dallas, TX). To extract the SR-B1 protein, a liver sample (~50 mg) was homogenized in buffer A containing 20 mmol/L Tris-HCl (pH 7.5), 2 mmol/L MgCl2, 0.2 mol/L sucrose, 5 mmol/L phenylmethylsulfonyl fluoride (PMSF), 10 μg/mL leupeptin, 20 μg/mL aprotenin, and 5 μg/mL pepstatin A. The crude extract was centrifuged at 10,000 × g for 10 min at 4°C, and the supernatant was then ultracentrifuged at 100,000 × g for 45 min at 4°C. The membrane fraction was resuspended in buffer B containing 62.5 mmol/L Tris-HCl (pH 6.8), 2% (v:v) SDS, 10% (v:v) glycerol and 5% (v:v) β-mercaptoethanol. To extract the LDLr protein, a liver sample (~100 mg) was homogenized in 1 mL of buffer C containing 50 mmol/L Tris (pH 7.5), 2 mmol/L CaCl2, 0.5% Triton X-100, and 1 mmol/L each of leupeptin, PMSE, and phenanthroline. The crude extract was centrifuged at 10,000 × g for 10 min at 4°C and the supernatant was then ultracentrifuged at 100,000 × g for 60 min at 4°C. The supernatant fractions were frozen at −80°C until further processing. Protein concentration of the liver extracts was determined by the method of Lowry et al. (24) SR-B1 (5 μg) and LDLr (10 μg) protein extracts per lane were loaded onto 7.5% polyacrylamide gels and separated by electrophoresis under nonreducing con
dations. The protein bands were transferred onto polyvinylidene fluoride membranes, exposed to 1:1500 anti-SR-B1 or 1:10,000 anti-LDLr, and revealed with 1:10,000 hors eradish peroxidase-linked anti-rabbit IgG (Amersham Biosciences).

**Statistical analysis.** Data are expressed as means ± SEM; except for SR-B1 and LDLr protein concentration, data were analyzed by factorial ANOVA with 2 factors: diet, with 2 levels (cholesterol-free, cholesterol 2%), and a factor termed “drug treatment” for convenience, with 3 levels (placebo, ACOL, pair-fed). Main and interactive effects of treatments were determined and are reported within ANOVA tables. Pairwise comparisons were also carried out to identify individual between-group differences using Fisher’s post-hoc Least Significant Difference test. The liver SR-B1 and LDLr data, which were obtained from the cohort fed the reference diet, were analyzed by one-factor ANOVA. Differences reaching the confidence level of $P < 0.05$ were considered significant.

**RESULTS**

Consumption of the C-diet did not affect final body weight, weight gain, food intake, or adipose tissue weights (Table 2). ACOL had its expected anorectic effect (−16%), with a consequent decrease in body weight (−8%), weight gain (−41%), as well as white adipose tissue weights (−45%). Pair feeding untreated rats had the same effects on these variables as did treatment of rats with ACOL (Table 2). Differences in weight gain were largely explained by changes in white adipose mass ($r = 0.80$, $P < 0.0001$). Liver weight was increased by the C-diet, partly because of cholesterol accumulation (see below), whereas ACOL and pair feeding reduced liver weight in proportion to body weight. The C-diet vastly increased liver cholesterol concentration. In fact, in the placebo-treated groups, the C-diet increased liver total cholesterol content from 15 to 440 mg cholesterol, which explained 28% of the C-diet–induced increase in liver weight. Neither ACOL treatment nor pair feeding affected liver cholesterol concentration.

Diet and ACOL treatment had interactive effects on total and lipoprotein cholesterol, i.e., the drug did not exert the same effect, or did so but to a different extent, depending on diet (Fig. 1). As expected, the C-diet greatly increased (275%) serum total cholesterol concentration (Fig. 1A), which in turn was reduced by ACOL treatment, but relatively more so in the C-fed rats (−56%) than in those fed the reference diet (−33%). Pair feeding did not affect cholesterolemia. The C-diet reduced HDL cholesterol in placebo-treated and pair-fed rats to approximately one third of the level in rats fed the reference cholesterol-free diet, whereas HDL cholesterol was unaffected by diet in ACOL-treated rats (Fig. 1B). Conversely, the C-diet dramatically increased non-HDL cholesterol (19-fold), and ACOL treatment, which did not affect rats fed the reference diet, dampened the diet-induced elevation in non-HDL cholesterol to less than half that in placebo-treated and pair-fed rats (Fig. 1C). In the cohort fed the reference diet, the ACOL-induced reduction in total cholesterol was due almost entirely to a reduction in HDL cholesterol, whereas the decrease in non-HDL cholesterol (Fig. 1C) totally explained the drug-induced decrease in cholesterol in C-fed rats.

To gain insight into the reasons why ACOL reduced cholesterolemia independently of its anorectic action, protein levels of SR-B1 and the LDLr were quantified in livers of rats fed the reference diet. ACOL, but not pair feeding, tended (−54%, $P = 0.1$) to increase liver SR-B1 concentration (Fig. 2A), and increased that of the LDLr by 170% ($P < 0.0001$; Fig. 2B). Serum total and HDL cholesterol concentrations, but not those of non-HDL cholesterol, were inversely correlated with those of liver SR-B1 ($r = 0.56$, $P < 0.02$, and $r = 0.70$, $P < 0.003$, respectively) and the LDLr ($r = 0.59$, $P < 0.005$, and $r = 0.65$, $P < 0.002$, respectively).

**DISCUSSION**

The 2 major goals of the present study were to assess whether the reduction in food intake elicited by treatment with the SERM ACOL plays a role in its hypcholesterolemic action, and whether the latter is maintained in diet-induced hypercholesterolemia. The findings demonstrate that ACOL reduces cholesterol independently of changes in food intake, and that the SERM maintains its hypcholesterolemic potential under cholesterol feeding.

The study confirms previous findings regarding the effects of ACOL on body weight, food intake, and fat deposition (8–10). The decrease in body weight, mainly in the form of fat stores, was associated with a reduction in food intake. The drug therefore shares some actions on energy and fat metabolism with natural estrogen and other antiestrogens (20,21,25–27).

In the cohort fed the cholesterol-free reference diet, the hypcholesterolemic action of ACOL described previously (7–10) was confirmed. Importantly, the 16% decrease in food intake did not contribute to the ACOL-induced reduction in cholesterolemia because the latter was identical in untreated rats that were pair-fed and those that consumed food ad libitum. ACOL therefore acts upon cholesterol metabolism through mechanisms that are independent of ingestive behavior and the effect of the latter on lipid metabolism. As ex-

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**TABLE 2**

<table>
<thead>
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<th>Purified + 2% Cholesterol diet</th>
<th>ANOVA</th>
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<td>Placebo</td>
<td>Acolbifene</td>
<td>Pair-fed</td>
</tr>
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<td>Body weight, g</td>
<td>238 ± 7</td>
<td>218 ± 6*</td>
<td>212 ± 4*</td>
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<td>g/21 d</td>
<td>49 ± 5</td>
<td>27 ± 3*</td>
<td>22 ± 2*</td>
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<td>Food, g/21 d</td>
<td>313 ± 12</td>
<td>253 ± 11*</td>
<td>259 ± 7*</td>
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<td>_white adipose, g</td>
<td>3.2 ± 0.3</td>
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<td>1.8 ± 0.1*</td>
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<td>Liver, g</td>
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<td>5.4 ± 0.2*</td>
<td>5.4 ± 0.2*</td>
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<tr>
<td>Liver cholesterol,</td>
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<td>6.0 ± 0.3</td>
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<td>µmol/g</td>
<td>134 ± 7†</td>
<td>131 ± 13†</td>
<td>137 ± 8†</td>
</tr>
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</table>

*Values are means ± SEM, $n = 6–7$ rats. * Different from placebo group; † different from corresponding drug treatment group fed the purified diet.
corresponding drug treatment group fed the purified diet. ACOLbifene-treated rats for 3 wk. Bars represent the means concentrations in rats fed a purified diet or the same diet containing 2% 

tendency for increased hepatic SR-B1 (ering in rats fed the reference diet was associated with a
decrease explained most of the hypocholesterolemic effect of 
culating cholesterol in rats fed a cholesterol-free diet, its

expected, because HDL transports more than two thirds of 
circulating cholesterol in rats fed a cholesterol-free diet, its
decrease explained most of the hypocholesterolemic effect of 
ACOL in rats fed the reference diet.

Further confirming a previous study (10), cholesterol low-
ering in rats fed the reference diet was associated with a
tendency for increased hepatic SR-B1 (P = 0.1) and a robust
elevation in LDLr protein concentrations. In these rats, most of 
the reduction in cholesterol occurred in the HDL fraction, 
which correlated strongly with both liver SR-B1 and LDLr 
concentrations. The ACOL-induced increase in SR-B1 protein 
levels was more variable and somewhat less robust than in 
our previous study (10); this may have been due to the differ-
ing lipid composition of the diets because dietary lipids were 
shown to modulate SR-B1 expression (28,29). The LDLr may 
have contributed to the small, nonsignificant reduction in the 
non-HDL cholesterol fraction, as well as to the clearance of 
apoE-containing HDL. Interestingly, pharmacologic amounts 
of natural estrogen (mostly estradiol) strongly upregulate liver 
LDLr expression, which explains most of its hypocholester-
olemic effect (30–35); at the same time, however, the hor-
mone completely blunts SR-B1 expression (19,36,37), in clear 
contrast to ACOL. The reasons why ACOL exerts either pro-
or antiestrogenic actions in a target-dependent fashion are 
likely linked to conformational determinants of the ligand-
receptor complex that differentially affect the recruitment of 
coactivators and corepressors (6). Although the involvement of 
the SR-B1 and LDLr in the cholesterol-lowering action of 
ACOL awaits direct confirmation, a mechanism of action 
related to the clearance of plasma cholesterol is highly likely 
because ACOL does not seem to affect hepatic cholesterol 
synthesis (10) and acts efficiently even in the absence of 
dietary cholesterol. It should be noted that in the present 
study, ACOL did not affect liver cholesterol concentration, 

FIGURE 1 Serum total (A), HDL- (B), and non-HDL (C) cholesterol 
concentrations in rats fed a purified diet or the same diet containing 2% 
cholesterol, treated or not with acolbifene, or untreated and pair-fed to 
acolbifene-treated rats for 3 wk. Bars represent the means ± SEM, n = 5–7 rats. *Different from the placebo group; †different from the 
corresponding drug treatment group fed the purified diet.

strongly suggesting that the liver was able to dispose of the 
additional cholesterol cleared from the circulation.

The cholesterol-enriched diet brought about the expected 
shift in cholesterol lipoprotein distribution, the bulk of which 
was found in the non-HDL fraction, along with a decrease in 
the absolute amount of cholesterol transported by HDL. Di-
etary cholesterol exerts these actions by favoring the hepatic 
assembly and secretion of cholesterol-enriched VLDL partic-
ules, with a concomitant increase in intravascular intermediate 
density lipoprotein formation, and by decreasing HDL 
formation through modulation of apolipoprotein production 
(38–40). Cholesterol feeding also lowers the liver abundance 
of the SR-B1 and LDLr (19,41). Remarkably, in the present 
study ACOL was able to blunt by approximately two thirds the 
3-fold increase in cholesterololomia brought by cholesterol feed-
ing, again independently of its effect on food intake. As 
expected, the non-HDL fraction explained all of the ACOL-
induced cholesterol lowering in this cholesterol-fed cohort. It 
should be mentioned that this fraction also contains some 
cholesterol-rich HDL; however, the contribution of the latter 
to the cholesterol content of the fraction termed non-HDL is 
minimal (39). Because livers from C-diet-fed rats were un-
available for receptor quantitation, it remains to be demon-
strated whether ACOL can prevent the reduction in SR-B1 and 
LDLr that would be expected with cholesterol feeding. 

Similarly, whether the hypocholesterolemic action of ACOL 
in diet-induced hypercholesterolemia is exerted through the 
same pathways as in normalipemic rats is unknown. Prelimi-
inary studies suggest that ACOL may modulate the expression 
of the cholesterol transporters ABCG5/8 (C. Lemieux and Y. 
Deshaias, unpublished observations), which may in turn affect 
its intestinal absorption, a contributor of potential importance 
under cholesterol feeding conditions. Further investigations of 
these major modulators of cholesterololomia are therefore clearly 
warranted to identify the precise mechanisms of the hypocho-
lesterolemic action of ACOL in cholesterol-fed rats.

The consequences of the HDL cholesterol-lowering action 
of ACOL in rats fed a cholesterol-free diet must be considered 
in the context of the fundamental differences that exist be-
tween human and rodent lipoprotein metabolism. As stated 
above, HDL is the major cholesterol carrier in rats, and a 
robust lowering of total cholesterol seldom occurs without a 
decrease in this fraction. Gene manipulation studies point to 
clearly beneficial consequences of high SR-B1 and reverse 
cholesterol transport activities (42–44). Therefore, in the 
context of human physiology, the ACOL-mediated tendency

FIGURE 2 Liver concentration of SR-B1 (A) and LDLr (B) protein 
in rats fed a purified diet without cholesterol, treated or not with 
acolbifene, or untreated and pair-fed to acolbifene-treated rats for 3 wk. 
Bars represent the means ± SEM, n = 5–7 rats. *Different from the 
placebo group, P < 0.05.
to increase liver SR-BI abundance, along with the increase in the LDLR, would both be considered to be positive effects in terms of cardiovascular risk associated with lipoprotein metabolism.

In summary, the present findings demonstrate that ACOL exerts its potent hypcholesterolemic action independently of the concomitant reduction in food intake and body fat accretion, possibly because the compound increases the abundance of key liver lipoprotein receptors that are not affected by a reduction in food intake equivalent to that elicited by the compound. The study also highlights the potency of ACOL as a hypcholesterolemic agent, because its action is largely maintained in rats with severe diet-induced hypercholesterolemia.

ACKNOWLEDGMENT

The authors gratefully acknowledge the expert professional assistance of Céline Martel.

LITERATURE CITED


27. Krieger, M. (1999) Charting the fate of the “good cholesterol”: identification of key liver lipoprotein receptors that are not affected by a reduction in food intake equivalent to that elicited by the compound. The study also highlights the potency of ACOL as a hypcholesterolemic agent, because its action is largely maintained in rats with severe diet-induced hypercholesterolemia.