Nutrient Metabolism

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

There is increasing interest in the development of compounds that antagonize the effects of female sex steroids in the prevention and treatment of estrogen-dependent cancers. Several such compounds, including a variety of selective estrogen receptor modulators (SERMs),2 were developed in the past years; some of these compounds display, along with their strong antiestrogenic action in cancer-prone tissues, interesting estrogen-like actions on bone mineral density, body fat accretion, and the plasma lipid profile (1–5). Of particular interest in this context is acoflifene (ACOL, also known as EM-652), a 4th generation SERM with potent and pure antiestrogenic properties in the mammary gland and uterus. ACOL was shown to reduce bone mineral mass loss and body fat accretion, as well as to improve the lipoprotein profile in rodents (6).

ACOL has consistently been found to dramatically reduce circulating cholesterol in rats (7–10). The fact that in these studies rats were fed a cholesterol-free diet strongly points to mechanisms related to hepatic cholesterol metabolism. Recently, we found (10) that ACOL upregulates the abundance of 2 key hepatic receptors responsible for a large fraction of cholesterol uptake by the liver: the scavenger receptor, class B, type 1 (SR-B1), which selectively takes up cholesterol from HDL (11–13) and plays a central role in reverse cholesterol transport; and the LDL receptor (LDLr), which internalizes mainly VLDL remnants, LDL particles, and apolipoprotein (apo)E-rich HDL (14–16). ACOL did not affect pathways of cholesterol synthesis (10), supporting the involvement of the clearance-related receptors in its hypocholesterolemic action.

As stated above, the hypocholesterolemic effect of ACOL was observed in rats fed a purified high carbohydrate diet virtually devoid of cholesterol (8–10). The downregulation of the LDLr by dietary cholesterol is well established (17,18), and a recent study demonstrated an identical effect of dietary cholesterol on the SR-B1 (19). These considerations raise the question whether ACOL retains its capacity to upregulate...
 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 acclimatized 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 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 1 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. 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 1 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.

Table 1: Composition of the reference diet

<table>
<thead>
<tr>
<th>Nutrient</th>
<th>Amount (g/kg diet)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cornstarch</td>
<td>265</td>
</tr>
<tr>
<td>Sucrose</td>
<td>265</td>
</tr>
<tr>
<td>Soybean oil</td>
<td>40</td>
</tr>
<tr>
<td>Beef tallow</td>
<td>100</td>
</tr>
<tr>
<td>Casein</td>
<td>215</td>
</tr>
<tr>
<td>DL-Methionine</td>
<td>4</td>
</tr>
<tr>
<td>Vitamin mix</td>
<td>11</td>
</tr>
<tr>
<td>Mineral mix</td>
<td>5</td>
</tr>
<tr>
<td>Cellulose</td>
<td>50</td>
</tr>
</tbody>
</table>

1 The C-diet had the same composition and was supplemented with 20 g/kg cholesterol.
2 Teklad no. 40060 Vitamin Mix (Teklad Test Diets), Composition (g/kg mix): p-aminobenzoic acid, 11.0; ascorbic acid, coated (97.5%), 101.7; biotin, 0.04; vitamin B-12 (0.1% trituration in mannitol), 3.0; calcium pantothenate, 6.6; choline dihydrogen citrate, 349.7; folic acid, 0.2; inositol, 11.0; menadione, 5.0; nacin, 9.9; pyridoxine-HCl, 2.2; riboflavin, 2.2; thiamin-HCl, 2.2; dry vitamin A palmitate (500,000 U/g), 4.0; dry cholecalciferol (500,000 U/g), 0.4; dry vitamin E acetate (500 U/g), 24.2; cornstarch (carrier), 466.7.
3 AlN-95G Mineral Mix (ICN Biochemicals), Composition (g/kg mix): calcium carbonate, 357.0; potassium phosphate, monobasic, 196.0; potassium citrate, monohydrate, 70.8; sodium chloride, 74.0; potassium sulfate, 46.6; magnesium oxide, 24.3; ferric citrate, 6.1; zinc carbonate, 1.6; manganese carbonate, 0.6; cupric carbonate, 0.3; tannic acid, 0.01; sodium selenite, 0.01; ammonium paramolybdate, 0.008; sodium meta-silicate, 1.4, chromium potassium sulfate, 0.3; lithium chloride, 0.02; boric acid, 0.08; sodium fluoride, 0.06; nickel carbonate, hydroxide, tetrahydrate, 0.03; ammonium vanadate, 0.007; sucrose, finely ground (carrier), 220.7.
4 Alphacel (ICN Biochemicals).

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HYPOLCHOLESTEROLEMIC ACTION OF THE SERM ACOLBIFENE

**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 quantitated 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-

**TABLE 2**

<table>
<thead>
<tr>
<th></th>
<th>Purified diet</th>
<th>Purified + 2% Cholesterol diet</th>
<th>ANOVA</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Placebo</td>
<td>Acolbifene</td>
<td></td>
</tr>
<tr>
<td>Body weight, g</td>
<td>238 ± 7</td>
<td>218 ± 6*</td>
<td></td>
</tr>
<tr>
<td>g/21 d</td>
<td>49 ± 5</td>
<td>27 ± 3*</td>
<td></td>
</tr>
<tr>
<td>Food, g/21 d</td>
<td>313 ± 12</td>
<td>253 ± 11*</td>
<td></td>
</tr>
<tr>
<td>Liver, g</td>
<td>3.2 ± 0.3</td>
<td>1.7 ± 0.1*</td>
<td></td>
</tr>
<tr>
<td>Liver cholesterol, μmol/g</td>
<td>5.6 ± 0.2</td>
<td>6.3 ± 0.4</td>
<td></td>
</tr>
</tbody>
</table>

1 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. /nH11005

acolbifene-treated rats for 3 wk. Bars represent the means
cholesterol, treated or not with acolbifene, or untreated and pair-fed to
concentrations in rats fed a purified diet or the same diet containing 2%

tendency for increased hepatic SR-B1 (19,41). Remarkably, in the present
study ACOL was able to blunt by approximately two thirds the 3
fold increase in cholesteryloma brought by cholesterol feeding,
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 normolipemic rats is unknown. Prelimi
nary 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 cholesteryloma 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 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
= 6–7 rats. *Different from the placebo group; †different from the
corresponding drug treatment group fed the purified diet.

expected, because HDL transports more than two thirds of circ
ulating 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 lower
ing 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 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

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LITERATURE CITED


