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Hyperlipoproteinemia Enhances Susceptibility to Acute Disseminated Candida albicans Infection in Low-Density-Lipoprotein-Receptor-Deficient Mice

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Recent studies have suggested the use of lipoproteins as an adjuvant treatment of lethal gram-negative infections. However, other important microorganisms for the etiology of sepsis, such as Candida species, grow better in lipid-rich environments. We investigated the effect of hyperlipoproteinemia on systemic candidiasis in low-density-lipoprotein-receptor-deficient (LDLR−/−) mice, in which the loss of the receptor results in a seven- to ninefold-higher plasma LDL level than that in their wild-type littermates (C57BL/6J). LDLR−/− mice died earlier, and the outgrowth of Candida albicans in the kidneys and livers of LDLR−/− mice was significantly higher compared with that of controls. After infection, circulating cytokine concentrations were significantly higher in LDLR−/− mice. In vitro, C. albicans grew better in plasma samples of LDLR−/− mice than in control plasma samples and peritoneal macrophages of LDLR−/− mice challenged with heat-killed C. albicans produced more cytokines than did those of controls. This latter phenomenon was probably due to increased binding of yeast cells to macrophages of LDLR−/− mice. These data suggest that hyperlipoproteinemia is deleterious in systemic candidiasis.

Acute disseminated candidiasis is a life-threatening condition that occurs predominantly in immunocompromised hosts. The mortality rate associated with disseminated candidiasis is high (31), and the incidence of this disease has increased in recent years (1, 3). Candida albicans ranks fourth among the organisms most frequently isolated from blood cultures in the United States (1). Clinically, systemic candidiasis sometimes mimics gram-negative sepsis. Viable Candida cells and cell wall constituents are able to induce the synthesis of proinflammatory cytokines in vitro (13, 14, 18), similar to gram-negative bacteria and their lipopolysaccharide (LPS) component (23). However, the role of these cytokines in systemic candidiasis is probably beneficial rather than deleterious (20, 26). In contrast, the induction of proinflammatory cytokines, such as interleukin-1α (IL-1α) and IL-1β and tumor necrosis factor alpha (TNF-α), is merely a deleterious event in gram-negative sepsis (4) and treatment aimed at blocking cytokine action has merely a deleterious event in gram-negative sepsis.

In vivo infusion of lipoproteins in rats protected the animals against mortality due to gram-negative bacterial sepsis in a model of cecal ligation and puncture (24). Taken together, these experiments demonstrate the capacity of lipoproteins to neutralize LPS and support their potential use as adjuvants in the therapy of sepsis.

On the other hand, earlier in vitro studies of lipid-containing parenteral solutions introduced for clinical use suggested that certain microorganisms, such as C. albicans and Staphylococcus aureus, grew better in a lipid-rich environment (5, 9, 16). Therefore, it is important to know the effect of hyperlipoproteinemia on the outcome of a systemic infection with these organisms before suggesting the application of infusion with lipoproteins as an adjuvant therapy in sepsis. In the present study, we assessed the influence of hyperlipoproteinemia on experimental systemic candidiasis in LDLR−/− mice.

MATERIALS AND METHODS

Animals. Homozygous C57BL/6 LDLR−/− mice and wild-type littermates were obtained from Jackson Laboratory (Bar Harbor, Maine) as mating pairs and bred in our local facility. For experiments, 6- to 8-week-old mice, weighing 20 to 25 g, were used. The animals were fed standard laboratory chow (Hope Farms, Woerden, The Netherlands) and housed under specific-pathogen-free conditions. The experiments were approved by the ethical committee for animal experiments at the Catholic University Nijmegen.

C. albicans infection. C. albicans (strain UCS20), maintained on agar slants at 4°C, was inoculated into 100 ml of Sabouraud broth and cultured for 24 h at 37°C. After three washes with pyrogen-free saline by centrifugation at 1,500 × g, the number of yeast cells was counted in a hemacytometer; occasional strings of two or more yeast cells were counted as 1 C. albicans CFU. The suspension was diluted to the appropriate concentration with pyrogen-free saline. The viability of yeast cells was at least 95%, as confirmed by plating serial dilutions on Sabouraud dextrose agar plates. Mice were injected intravenously (i.v.) in the retro-orbital plexus with 1010 or 108 CFU of C. albicans. Survival was assessed daily for 14 days in groups of at least 15 animals. In separate groups, after 4 h and 1 and 3 days, subgroups of five mice were killed by cervical dislocation and blood samples were collected for the measurement of plasma cytokine concentrations.
The outgrowth of microorganisms from the livers, spleens, and kidneys of animals was quantified on days 1 and 3 after 
C. albicans infection. For this purpose, the organs were removed aseptically, weighed, and homogenized in sterile saline in a 
tissue homogenizer. The number of viable C. albicans cells in tissue samples was determined by plating serial dilutions on Sabouraud dextrose agar plates as 
described previously (15), and CFU were counted after overnight incubation at 37°C. The results were expressed as the log CFU per gram of tissue.

RESULTS

C. albicans infection. Plasma cholesterol concentrations decreased significantly during 
C. albicans infection in both mouse strains (P < 0.05) but remained three to four times higher in 
LDLR-/- mice than in control mice (P < 0.01) (Table 1). The initial plasma triglyceride levels in LDLR-/- mice were two 
times higher than those of controls (P < 0.01). Four hours after infection, triglyceride levels decreased significantly in 
LDLR-/- mice (P < 0.05) and only marginally in control animals (P > 0.05) (Table 1). The triglyceride levels 4 h after 
C. albicans infection did not differ significantly between the two mouse strains (P > 0.05) (Table 1). After i.v. injection of 
either 10^6 or 10^7 CFU of C. albicans, LDLR-/- mice died significantly earlier than did control animals (P < 0.05) (Fig. 1).

One day after infection with 10^6 CFU of C. albicans, yeast 
outgrowth in the kidneys of LDLR-/- mice was significantly higher compared with that of controls; 3 days after infection, 
outgrowth of C. albicans was increased in both the kidneys and livers of LDLR-/- mice (Table 2). No difference in the out­ 
growth of C. albicans in the spleen was detected between the two mouse strains (Table 2).

Four hours after infection, plasma TNF-a concentrations were below the detection limit and IL-1a concentrations were 
higher in LDLR mice than in controls (140 ± 22 versus 78 ± 48 pg/ml, respectively; P < 0.05). Circulating IL-1b concentrations were similar in both strains (56 ± 29 versus 37 ± 28 pg/ml, respectively; P > 0.05). Plasma TNF-a concentrations were significantly higher in LDLR-/- mice compared with 
those of control animals at both 24 (42 ± 5 versus 18 ± 4 pg/ml, respectively; P < 0.01) and 72 (302 ± 237 versus 98 ± 88 pg/ml, respectively; P < 0.02) h after infection (Fig. 2). No differences in plasma IL-1a and IL-1b concentrations were observed at these time points (Fig. 2).

In vitro cytokine production. We investigated the capacity of 
peritoneal macrophages of both mouse strains to produce cy­ 
tokines when stimulated in vitro with heat-killed C. albicans. Compared with those of controls, the TNF-a concentrations in 
supernatants from macrophages of LDLR-/- mice were signifi­ 
cantly higher (Fig. 3a). The IL-1a and IL-1b concentrations were similar in both strains (56 ± 29 versus 37 ± 28 
pg/ml, respectively; P > 0.05). Circulating IL-1b concentrations were significantly higher in LDLR-/- mice compared with 
those of control animals at both 24 (42 ± 5 versus 18 ± 4 
pg/ml, respectively; P < 0.01) and 72 (302 ± 237 versus 98 ± 88 pg/ml, respectively; P < 0.02) h after infection (Fig. 2). No differences in plasma IL-1a and IL-1b concentrations were observed at these time points (Fig. 2).

### Table 1. Plasma cholesterol and triglyceride concentrations before and after C. albicans infection in LDLR-/- and C57BL/6J mice

<table>
<thead>
<tr>
<th>Mouse strain</th>
<th>Time</th>
<th>Mean conen (mmol/liter) ± SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>LDLR-/-</td>
<td>Before</td>
<td>9.55 ± 1.11**</td>
</tr>
<tr>
<td>LDLR-/-</td>
<td>After</td>
<td>6.76 ± 0.92**, 0.58 ± 0.05**</td>
</tr>
<tr>
<td>C57BL/6J</td>
<td>Before</td>
<td>2.25 ± 0.45</td>
</tr>
<tr>
<td>C57BL/6J</td>
<td>After</td>
<td>1.84 ± 0.09**, 0.49 ± 0.11</td>
</tr>
</tbody>
</table>

* Mice were infected i.v. with 10^6 CFU of C. albicans. Each group consisted of five mice.

** P < 0.05 for comparison between results for LDLR-/- and control mice; 
*** P < 0.05 for comparison between results before and after C. albicans infection.

FIG. 1. Survival during C. albicans infection. LDLR-/- mice (closed symbols) infected with either 10^6 (triangles) or 10^7 (circles) CFU of C. albicans died significantly earlier than did C57BL/6J mice (open symbols) (P < 0.05; Kaplan-Meier log rank test). The data are pooled results of two experiments with at least 15 animals per group.
IL-1β concentrations did not differ between the two mouse strains (Fig. 3b).

**C. albicans binding.** To investigate the total binding of *C. albicans* to macrophages from the two mouse strains, we incubated radiiodinated *C. albicans* cells with macrophages (5:1 ratio) and determined the amounts bound to macrophages after 5, 10, and 30 min of incubation. The binding of yeast cells to macrophages of LDLR mice was increased and more rapid compared to that of control macrophages (Fig. 4).

**Growth of *C. albicans* in vitro.** To investigate whether the outgrowth of *C. albicans* in the organs of LDLR −/− mice is due to enhanced growth of yeast cells in a lipoprotein-rich environment, we compared the in vitro growth of 10⁵ CFU of *C. albicans* in plasma samples (diluted 1:1 with Sabouraud medium) of LDLR −/− mice and controls. The capacity of LDLR −/− plasma to inhibit the outgrowth of *C. albicans* was decreased compared with that of control plasma, as shown by the growth of *C. albicans* in the two types of plasma samples after 12 [(7.2 ± 2.4) × 10⁴ versus (2.4 ± 1.5) × 10⁴ CFU/ml, respectively; *P < 0.05] and 24 [(4.2 ± 1.6) × 10⁴ versus (1.5 ± 1.0) × 10⁴ CFU/ml, respectively; *P < 0.05] h of incubation.

**DISCUSSION**

The main conclusion from the present study is that hyperlipoproteinemia has deleterious effects on the outcome of severe *C. albicans* infection, in contrast to gram-negative bacterial infections. We have shown that LDLR −/− mice, with seven- to nine-times-higher LDL levels, are more susceptible to *C. albicans* infection than are their wild-type littermates. The earlier mortality of LDLR −/− mice was associated with increased outgrowth of *C. albicans* in their organs, and these mice produced significantly more proinflammatory cytokines than did control mice.

In general, mortality after infection may be due to lethal cytokinemia or to functional impairment by the growth of *C. albicans*. It may be hypothesized that the enhanced outgrowth of *C. albicans* in the organs of LDLR −/− mice is due to elevated lipoprotein concentrations. Normal serum has a candidicidal property (22), whereas anti-TNF antibodies (26) or pharmacologic inhibition of proinflammatory cytokines proved to be deleterious during severe *C. albicans* infection (20). Thus, proinflammatory cytokines seem to play a beneficial rather than a deleterious role in the defense against *C. albicans*. It should be noted that despite the greater cytokine response, LDLR −/− mice were not protected against *C. albicans* infection, probably due to the overwhelming outgrowth of yeast cells in their organs.

It may be hypothesized that the enhanced outgrowth of *C. albicans* in the organs of LDLR −/− mice is due to elevated lipoprotein concentrations. Normal serum has a candidicidal effect (22), and as shown by in vitro growth experiments, this property was significantly decreased in plasma samples from LDLR −/− mice. This effect may be due to the use of lipoproteins as a nutrition factor by *C. albicans*, as has been suggested by earlier studies showing increased growth of *C. albicans* in lipid-containing parenteral solutions compared with that in formulations without lipid contents (5, 9, 16). Another possible mechanism by which lipoproteins could influence *C. albicans* growth is interaction with other plasma factors. We cannot

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**TABLE 2. Outgrowth of *C. albicans* in the organs of LDLR −/− and C57BL/6J mice after infection with 10⁶ CFU of *C. albicans***

<table>
<thead>
<tr>
<th>Mouse strain</th>
<th>Organ</th>
<th>Day</th>
<th>Log CFU/g of tissue</th>
</tr>
</thead>
<tbody>
<tr>
<td>LDLR −/−</td>
<td>Kidney</td>
<td>1</td>
<td>6.3 ± 0.5*</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3</td>
<td>7.4 ± 0.3**</td>
</tr>
<tr>
<td></td>
<td>Liver</td>
<td>1</td>
<td>4.8 ± 0.3**</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3</td>
<td>5.2 ± 0.2**</td>
</tr>
<tr>
<td></td>
<td>Spleen</td>
<td>1</td>
<td>4.9 ± 0.2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3</td>
<td>5.1 ± 0.5</td>
</tr>
<tr>
<td>C57BL/6J</td>
<td>Kidney</td>
<td>1</td>
<td>5.9 ± 0.2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3</td>
<td>6.8 ± 0.3</td>
</tr>
<tr>
<td></td>
<td>Liver</td>
<td>1</td>
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<tr>
<td></td>
<td>Spleen</td>
<td>1</td>
<td>5.1 ± 0.2</td>
</tr>
<tr>
<td></td>
<td></td>
<td>3</td>
<td>5.0 ± 0.2</td>
</tr>
</tbody>
</table>

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* Data are means ± standard deviations of pooled data from two experiments with 10 animals per group. *, *P < 0.01; **, *P < 0.05.
exclude the possibility that plasma candidicidal factors, such as platelet microbicidal protein (32) and the calprotectin complex (19), are bound and inactivated by lipoproteins.

The higher cytokine concentrations during infection in LDLR<sup>−/−</sup> mice, compared with those of controls, were probably at least in part a response of the host against enhanced C. albicans outgrowth in the organs of LDLR<sup>−/−</sup> mice. However, surprisingly, stimulated in vitro with heat-killed C. albicans, macrophages of LDLR<sup>−/−</sup> mice produced significantly more TNF-α and IL-1α than macrophages of control mice did. Most likely, this was due to the observed increased binding of C. albicans to macrophages of LDLR<sup>−/−</sup> mice compared with that of control macrophages. This phenomenon may be explained by the influence of constitutively increased lipoprotein concentrations in LDLR<sup>−/−</sup> mice on the Candida-binding proteins on macrophage. It has been shown previously that hypercholesterolemia is able to modify the number and clustering of other receptors, such as the LPS receptor CD14 (7, 25). Earlier, we observed similar higher binding of radiolabelled LPS to macrophages of LDLR<sup>−/−</sup> mice, followed by higher cytokine production (21). Thus, similar changes in the number and/or clustering of Candida-binding proteins may facilitate the binding of C. albicans to macrophages of LDLR<sup>−/−</sup> mice, with a subsequent increase in cytokine production. Hyperlipoproteinemia could also modify the hydrophobicity of cells, which may also influence the adherence of C. albicans to macrophages, as has been shown for endothelial cells (10). Which of these mechanisms is responsible for the observed increase in cytokine production by macrophages of LDLR<sup>−/−</sup> mice is under study.

An alternative desirable experiment to our model in order to investigate the influence of hyperlipoproteinemia in C. albicans infection would have been to infuse lipoproteins into animals before and during infection. However, an infusion of lipoproteins into mice is not possible and other models of lipoprotein infusion in rabbits (11) and rats (24) are short-term models that are not suitable for sustained lipid infusion during systemic candidiasis. Therefore, the genetically modified mouse model is a good alternative for studying the in vivo effects of hyperlipoproteinemia in models of sustained infection.

In conclusion, hyperlipoproteinemia has deleterious effects on the course of acute disseminated C. albicans infection, in contrast to its beneficial effect in gram-negative infection. Although no epidemiological studies have been done to show a relationship between hyperlipoproteinemia and increased susceptibility to C. albicans, an infusion of lipoproteins into a patient with disseminated candidiasis under the presumptive diagnosis of gram-negative sepsis may prove deleterious. These divergent effects of hyperlipoproteinemia should be taken into account when the use of lipoproteins as an adjuvant treatment of sepsis is considered.

ACKNOWLEDGMENTS

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REFERENCES

CANDIDIASIS AND HYPERLIPROTEINEMIA


