Supplementary heat-killed *Lactobacillus reuteri* GMNL-263 ameliorates hyperlipidaemic and cardiac apoptosis in high-fat diet-fed hamsters to maintain cardiovascular function

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Obesity and hyperlipidaemia increase the risk of CVD. Some strains of probiotics have been suggested to have potential applications in cardiovascular health by lowering serum LDL-cholesterol. In this work, high-fat diet-induced hyperlipidaemia in hamsters was treated with different doses (5 × 10⁸ and 2.5 × 10⁹ cells/kg per d) of heat-killed *Lactobacillus reuteri* GMNL-263 (Lr263) by oral gavage for 8 weeks. The serum lipid profile analysis showed that LDL-cholesterol and plasma malondialdehyde (P-MDA) were reduced in the GMNL-263 5 × 10⁸ cells/kg per d treatment group. Total cholesterol and P-MDA were reduced in the GMNL-263 2.5 × 10⁹ cells/kg per d treatment group. In terms of heart function, the GMNL-263 2.5 × 10⁹ cells/kg per d treatments improved the ejection fraction from 85-71 to 91-81 % and fractional shortening from 46-95 to 57-92 % in the high-fat diet-fed hamster hearts. Moreover, the GMNL-263-treated, high-fat diet-fed hamster hearts exhibited reduced Fas-induced myocardial apoptosis and a reactivated IGF1R/PI3K/Akt cell survival pathway. Interestingly, the GMNL-263 treatments also enhanced the heat-shock protein 27 expression in a dose-dependent manner, but the mechanism for this increase remains unclear. In conclusion, supplementary heat-killed *L. reuteri* GMNL-263 can slightly reduce serum cholesterol. The anti-hyperlipidaemia effects of GMNL-263 may reactivate the IGF1R/PI3K/Akt cell survival pathway and reduce Fas-induced myocardial apoptosis in high-fat diet-fed hamster hearts.

Key words: Hyperlipidaemia: LDL-cholesterol: *Lactobacillus reuteri* GMNL-263 (Lr263): Heart protection

It was recently proven that cumulative exposure to hyperlipidaemia during early adulthood increases the risk of CVD in a dose-dependent manner. In fact, prolonged exposure to decreased LDL-cholesterol beginning in early life significantly reduces the risk of CVD. Recently, some strains of probiotics, such as multi-strain probiotic capsules (*Streptococcus thermophilus, Lactobacillus plantarum, Lactobacillus acidophilus, Lactobacillus rhamnosus, Bifidobacterium lactis, Bifidobacterium longum and Bifidobacterium breve*), were reported to significantly reduce serum cholesterol, waist circumference and body weight in adults with a BMI > 25 kg/m² during an 8-week treatment. In addition, *Lactobacillus reuteri* NCIMB 30242, a probiotic associated with cardiovascular health, claims to clinically lower LDL-cholesterol levels by 11 % in hyperlipidaemic adults, and it has been approved by Health Canada.

Abbreviations: %FS, fractional shortening; EF, ejection fraction; HSP27, heat-shock protein 27; P-MDA, plasma malondialdehyde.

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Supplementary *L. reuteri* may reduce serum cholesterol because of its high bile salt hydrolase (BSH) activity. Probiotics with high BSH activity promote secondary amino acid conjugates and the deconjugation of bile acids. Finally, probiotics process bile salts and block cholesterol absorption in the gut. Previously, *L. reuteri* GMNL-263 (Lr263), a probiotic strain, was reported to have activity that may improve insulin resistance and ameliorate hepatic steatosis in hamsters fed a high-fructose diet.

However, there are still some doubts about whether probiotics arrive alive in the gut after gastric acid exposure. In 2013, Shinkai et al. performed a randomised, double-blind, placebo-controlled trial that proved that oral intake of the heat-killed *Lactobacillus pentosus* strain b240 had similar immunoprotective effects as probiotics. In the current study, heart-protective and anti-hyperlipidaemic effects were investigated using a high-fat diet to induce hyperlipidaemia in hamsters. The hamsters were treated with different doses of heat-killed *L. reuteri* GMNL-263 (Lr263) via oral gavage for 8 weeks.

**Methods**

**Preparation of probiotic suspensions**

Heat-killed *L. reuteri* GMNL-263 was provided by GenMont Biotech Inc. Two concentrations of *L. reuteri* GMNL-263 (5 × 10⁹ and 2.5 × 10⁹ cells/ml) were prepared in PBS for oral gavage treatments.

**Animal model**

In all, twenty-four male golden Syrian hamsters without spontaneous cardiomyopathy (*Mesocricetus auratus*, 6 weeks old) were purchased from the National Laboratory Animal Center and divided into four groups (n = 6 each). The experimental protocol used in this study was approved by the Institutional Animal Care and Use Committee of China Medical University (No.100-4-B). The normal control group hamsters were fed a normal diet with water. An HF group and are presented as the group mean values and standard deviations. A one-way ANOVA was used to indicate an overall statistical significance from the means of the four experimental groups. A *P* value <0.05 was considered significant. Statistical analyses were performed using the SigmaPlot v.10.0 software.
Results

Serum lipid profile

After 8 weeks, the TAG, TC, LDL-cholesterol, HDL-cholesterol and P-MDA serum levels were significantly increased in the hamsters fed a high-fat diet. These results are shown in Table 1. In the hamsters treated with GMNL-263 $5 \times 10^8$ cells/kg per d and fed the high-fat diet for 8 weeks, LDL-C and P-MDA were reduced. Further, after 8 weeks, the TC and P-MDA serum levels were further reduced in the hamsters treated with GMNL-263 $2.5 \times 10^8$ cells/kg per d and fed the high-fat diet. There was no difference in the ratio of LDL-cholesterol/HDL-cholesterol between the HF and HKL groups (GMNL-263 $5 \times 10^8$ cells/kg per d treatment group). However, the LDL-cholesterol/HDL-cholesterol ratio was reduced in the HKL group (GMNL-263 $2.5 \times 10^8$ cells/kg per d treatment group) compared with the HF group.

Echocardiography

After 8 weeks, the EF of hamster hearts was reduced from 89-79 (so 0·72) % in the control group to 85-71 (so 2·36) % in the HF group. These results are shown in Table 2. The %FS of hamster hearts was reduced from 53-80 (so 2·17) % in the control group to 46-93 (so 5·04) % in the HF group. In the hamsters treated with GMNL-263 $5 \times 10^8$ cells/kg per d and fed the high-fat diet, the EF was slightly increased to 87-18 (so 2·41) %, and the %FS increased to 51-31 (so 3·15) %. Furthermore, in the group treated with GMNL-263 $2.5 \times 10^8$ cells/kg per d and fed the HKL diet, the EF improved up to 91·81 (so 0·71) % and the %FS also improved up to 57·92 (sd 1·30) %.

Myocardial biopsy

H&E staining was used for the hearts from each group. After 8 weeks, the cardiomyocytes from the HF group were in disarray and had more space between the cells (Fig. 1). In the group treated with GMNL-263 $5 \times 10^8$ cells/kg per d and fed the high-fat diet, the myocardial disarray and space between the cells were slightly improved. In the group treated with GMNL-263 $2.5 \times 10^8$ cells/kg per d and fed the high-fat diet, the myocardial disarray and space between the cardiomyocytes were significantly reduced.

Table 1. Serum lipid profile

<table>
<thead>
<tr>
<th></th>
<th>N</th>
<th>HF</th>
<th>HKL</th>
<th>HKH</th>
</tr>
</thead>
<tbody>
<tr>
<td>TAG (mmol/l)</td>
<td>0.89***</td>
<td>0.221</td>
<td>3.961</td>
<td>0.328</td>
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<tr>
<td>TC (mmol/l)</td>
<td>3.310***</td>
<td>0.194</td>
<td>7.433</td>
<td>1.046</td>
</tr>
<tr>
<td>LDL-cholesterol</td>
<td>0.355***</td>
<td>0.065</td>
<td>2.971</td>
<td>0.865</td>
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<tr>
<td>HDL-cholesterol</td>
<td>1.878*</td>
<td>0.186</td>
<td>2.440</td>
<td>0.394</td>
</tr>
<tr>
<td>LDP-cholesterol/HDL-cholesterol</td>
<td>0.2***</td>
<td>0.04</td>
<td>1.2</td>
<td>0.35</td>
</tr>
<tr>
<td>P-MDA (μmol/l)</td>
<td>3.96*</td>
<td>0.04</td>
<td>6.59</td>
<td>0.53</td>
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</tbody>
</table>

N, normal control; HF, group of hamsters fed a high-fat diet only with normal water; HKL, group of hamsters fed the high-fat diet with normal water and heat-killed *Lactobacillus reuteri* GMNL-263 $5 \times 10^8$ cells/kg per d via oral gavage; HKH, group of hamsters fed the high-fat diet with normal water and heat-killed *L. reuteri* GMNL-263 $2.5 \times 10^8$ cells/kg per d via oral gavage; TC, total cholesterol; P-MDA, plasma malondialdehyde. * P < 0·05, ** P < 0·01 compared with HF group.

Table 2. Echocardiography

<table>
<thead>
<tr>
<th>Parameters</th>
<th>N</th>
<th>HF</th>
<th>HKL</th>
<th>HKH</th>
</tr>
</thead>
<tbody>
<tr>
<td>IVSd (mm)</td>
<td>0.89</td>
<td>0.02</td>
<td>0.90</td>
<td>0.02</td>
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<td>LVdDd (mm)</td>
<td>4.79</td>
<td>0.37</td>
<td>4.84</td>
<td>0.32</td>
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<td>LVPWd (mm)</td>
<td>0.99</td>
<td>0.01</td>
<td>0.99</td>
<td>0.01</td>
</tr>
<tr>
<td>IVSs (mm)</td>
<td>1.97</td>
<td>0.05</td>
<td>1.71***</td>
<td>0.06</td>
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<tr>
<td>LVIDs (mm)</td>
<td>2.46</td>
<td>0.23</td>
<td>1.98**</td>
<td>0.02</td>
</tr>
<tr>
<td>LVWs (mm)</td>
<td>1.81</td>
<td>0.06</td>
<td>2.17**</td>
<td>0.15*</td>
</tr>
<tr>
<td>EDV (ml)</td>
<td>0.27</td>
<td>0.06</td>
<td>0.20</td>
<td>0.02</td>
</tr>
<tr>
<td>ESV (ml)</td>
<td>0.03</td>
<td>0.01</td>
<td>0.04</td>
<td>0.02</td>
</tr>
<tr>
<td>EF (%)</td>
<td>89.79</td>
<td>0.72</td>
<td>85.71*</td>
<td>2.36</td>
</tr>
<tr>
<td>%FS (%)</td>
<td>53.80</td>
<td>2.17</td>
<td>46.93</td>
<td>5.04</td>
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<tr>
<td>LV mass (g)</td>
<td>0.83</td>
<td>0.03</td>
<td>0.75***</td>
<td>0.01</td>
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</table>

N, normal control; HF, group of hamsters fed a high-fat diet only with normal water; HKL, group of hamsters fed the high-fat diet with normal water and heat-killed *Lactobacillus reuteri* GMNL-263 $5 \times 10^8$ cells/kg per d via oral gavage; HKH, group of hamsters fed the high-fat diet with normal water and heat-killed *L. reuteri* GMNL-263 $2.5 \times 10^8$ cells/kg per d via oral gavage; IVSd, interventricular septum diastolic; LVdDd, left ventricular dimension diastolic; IVSs, interventricular septum systolic; LVIDs, left ventricular dimension systolic; LVWs, left ventricular posterior wall systolic; EDV, end-diastolic velocity; ESV, end-systolic velocity; EF, ejection fraction; %FS, fractional shortening; LV mass, left ventricular mass. * P < 0·05 compared with the normal group, ** P < 0·01 compared with the normal group, *** P < 0·001 compared with the normal group, **** P < 0·05 compared with the HF group, ***** P < 0·01 compared with the HF group, ****** P < 0·001 compared with the HF group.
Myocardial apoptosis signalling analysis

In the protein analysis, the Fas ligand protein level was higher in the HF hamster hearts compared with those of the control group (Fig. 2). The proteins downstream of Fas ligand, especially caspase-8, were more highly expressed and had higher levels of cleaved active form in the HF hamster hearts compared with the controls. In the GMNL-263 5 × 10⁸ cells/kg per d treatment plus HKL and the GMNL-263 2.5 × 10⁹ cells/kg per d treatment plus HKH hamster hearts, Fas ligand expression was reduced. The protein levels of active caspase-8 and AIF were also reduced. Further, HSP27 was increased in both the GMNL-263 5 × 10⁸ cells/kg per d treatment plus HKL and the GMNL-263 2.5 × 10⁹ cells/kg per d treatment plus HKH hamster hearts.

A DAPI and TUNEL dual staining assay was used to evaluate myocardial apoptosis in the hamster heart sections from each group. Some apoptotic cardiomyocytes were labelled by green TUNEL stain in the high-fat diet-only hamster hearts (Fig. 3).
After 8 weeks, the number of apoptotic cardiomyocytes in the GMNL-263 5 × 10⁸ cells/kg per d treatment plus HKL hamster hearts was reduced. Moreover, apoptosis of the cardiomyocytes in the GMNL-263 2·5 × 10⁹ cells/kg per d treatment plus the HKH hamster heart group was inhibited.

**Myocardial survival signalling analysis**

There were no significant differences in the p-IGF1R and downstream p-PI3K, p-Akt, Bcl-2 and p-Bad protein levels between the control and the HF hamster hearts (Fig. 4). After 8 weeks, the expression levels of p-IGF1R downstream proteins p-PI3K, p-Akt and p-Bad were significantly increased in the GMNL-263 5 × 10⁸ cells/kg per d-treated HKL group and GMNL-263 2·5 × 10⁹ cells/kg per d-treated HKH group. The Bcl-2 expression level was increased in the HKL group, but not significantly in the HKH group.

**Discussion**

Dynamic microbial communities significantly affect human health, including the cardiovascular system. In CVD, the
most investigated application for probiotic therapy is the reduction of serum LDL-cholesterol. Increased LDL-cholesterol levels is a major risk factor for CVD(14). Previously, Shi et al.(15) used heat-inactivated Lactobacillus gasseri treatments and reduced metabolic syndrome symptoms in high-fat and high-salt diet-fed Sprague–Dawley rats. Similarly, L. reuteri can reduce serum cholesterol by interrupting lipid acid and bile acid conjugation. In addition, using heat-killed L. reuteri GMNL-263 treatments still can slightly decrease serum LDL-cholesterol in high-fat diet-fed hamsters, as shown in Table 1. The BSH activity of heat-killed L. reuteri GMNL-263 remains unclear. The mammalian intestine harbours a great number of bacteria (approximately 10^{14} bacteria)(16). Supplementing a specific amount of probiotics, such as GMNL-263 bacteria, might change the gut microbiota environment and affect serum cholesterol.

Serum LDL-cholesterol was only somewhat reduced in the GMNL-263 treatment plus high-fat diet groups, but the heart functions were improved and are shown in Table 1. In 8 weeks, the GMNL-263 2.5 x 10^6 cells/kg per d treatment improved the EF from 85.71 % in the high-fat diet-only hamster hearts to 91.81 % and the %FS from 46.93 to 57.92 %. In the heart section investigations, the GMNL-263 treatments also improved the myocardial disarray (Fig. 1).

Our previous work indicated that increased expression of Fas ligand and its receptor Fas leads to cardiomyocyte apoptosis through the release and activation of caspase-8 from FADD(17). In this study, the GMNL-263 treatments reduced the Fas ligand expression and downstream apoptosis-inducing signalling proteins (Fig. 2 and 3). Further, HSP27 is thought to be inhibitory against Fas-induced apoptosis(18,20). Interestingly, the GMNL-263 treatments in the high-fat diet-fed hamsters increased HSP27 in a dose-dependent manner. This result is similar to the other strains of probiotic bacteria, such as Lactobacillus plantarum, Lactobacillus brevis and Bacillus subtilis, in their host(21-24).

Recently, a study revealed that selenium-enriched probiotics might increase the HSP27 and HSP70 mRNA levels(25). However, the mechanism behind the GMNL-263 treatments increasing the HSP27 expression levels and the role of HSP27 in hyperlipidaemia need further investigation.

After the 8-week experiment, there was no significant difference in the IGF1R-associated cell survival signalling pathway between the control group and the HF group (Fig. 4). However, serum cholesterol-lowering effects might increase the p-IGF1R and downstream p-P3K, p-Akt, Bcl-2 and p-Bad reactivation in the GMNL-263 treatment groups.

In conclusion, supplementary heat-killed L. reuteri GMNL-263 can slightly reduce serum cholesterol. The moderate anti-hyperlipidaemia effects of GMNL-263 may reactivate the IGF1R/P3K/Akt cell survival pathway and reduce Fas-induced myocardial apoptosis in high-fat diet-fed hamster hearts.

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The authors declare no conflicts of interest.

References