The atheroprotective roles of heart-protecting musk pills against atherosclerosis development in apolipoprotein E-deficient mice

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Background: Heart-protecting musk pill (HMP), derived from Chinese herbal medicines, has been found to possess protective roles against atherosclerosis-related cardiovascular diseases (CVDs), however, the anti-atherosclerotic mechanisms of HMP are still unclear. Here, we investigated the effects of HMP on alleviating atherosclerotic lesion severity in mice and explored the molecular mechanisms.

Methods: Apolipoprotein E-deficient mice were fed western-type diet supplemented with HMP (25 mg/kg/day) or normal saline gavage for 20 weeks. Then histopathological staining was performed to assess the atheromatous plaque burden. Biochemical kits were used to detect levels of lipid profiles. Moreover, effector factors associated with lipid metabolism in liver and intestinal tissues were investigated by western blot and real-time PCR assays. Levels of signal molecules participating in the mitochondrial-mediated apoptosis pathway were detected by Western blot.

Results: We found that HMP notably reduced atherosclerotic lesion size (P<0.05) and improved plaque stability (P<0.05). HMP treatment decreased circulating TC (P<0.01), LDL-C (P<0.01) and TG (P<0.05) levels and increased HDL-C (P<0.05) content. HMP was found to suppress SREBP2, HMGCR and PCSK9 expressions (P<0.05), yet promote LDLR expression (P<0.05) in hepatocytes. Moreover, HMP was discovered to activate PPARα/CPT-1A cascade (P<0.05) and inhibit contents of SREBP1c and the lipogenic genes FAS and ACCα (P<0.05). The LBK1/AMPK cascade was also activated after HMP administration (P<0.05). Additionally, HMP was found to facilitate transintestinal cholesterol excretion by increasing ABCG5 and ABCG8 levels and reducing NPC1L1 content (P<0.05). In terms of vasoprotective activities, we observed that HMP decreased cleaved caspase-3 content (P<0.05) in the vascular intima, which might be due to inhibition of mitochondrial-related signaling pathway.

Conclusions: Altogether, our study indicates that HMP plays anti-atherosclerotic roles via regulating lipid metabolism and improving vascular intimal injury.

Keywords: Heart-protecting musk pill (HMP); atherosclerosis; lipid metabolism; vascular intimal injury

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Introduction

Atherosclerosis, featured by progressive plaque lesion formation leading to lumen stenosis commonly occurring in large and medium-sized artery walls, is the principal pathological basis of cardiovascular diseases (CVDs) (1). The development of atheroma lesions is associated with a series of chronic processes involving lipid metabolic dysfunction and vascular intimal injury (2,3). Damaged intima facilitates excessive lipid deposition and circulating leukocytes accumulation in the vessel subendothelial space, which are pivotal pathogenic events facilitating atherogenesis (4). It is well-established that dyslipidemia is a crucial risk factor for atherosclerosis and regulation of blood lipid profiles has been given priority in the treatment of atherosclerotic CVDs (2,5). Moreover, therapeutic approaches protecting against vascular intimal impairment have been proved to inhibit atheroma lesion progression (6,7). Currently, several modern drugs such as statins have been used for the treatment of atherosclerosis. However, up to 20% patients are unable to tolerate statin treatment due to the side effects and the residual risk of adverse cardiovascular events is still high in several patients treated with statins (8,9). Accordingly, highly safe and effective agents targeting the atherogenic processes are urgently needed.

Heart-protecting musk pill (HMP), deriving from an ancient traditional Chinese medicine (TCM) Suhexiang Pill, possesses therapeutic properties against several CVDs in clinical application, such as ischemic heart disease, heart failure and hypertension (10). HMP comprises seven medicinal substances: Radix Ginseng (Panax ginseng C.A.Mey. root), Venenum Bufonis, Styrax (Liquidambar orientalis Mill. balsam), Calculus Bovis Artifactus, Cortex Cinnamomi (Cinnamomum cassia Presl. bark), Borneolum Syntheticum (Dryobalanops aromatica C.F.Gaertn. resin) and Artificial Moschus (11). Zhao et al. have indicated that HMP is capable of alleviating hypertensive renal injury by suppressing the TLR4/NF-κB inflammatory pathway (11). Moreover, HMP has been found to be a potent modulator of blood lipid spectrum in the hyperlipidemic state (12). It has been observed that HMP improves cardiac function and reduces damaged area in rats with myocardial infarction via promoting endothelial progenitor cell mobilization and proliferation (13). In addition, findings of other studies show that HMP is capable of alleviating cardiac pathological remodeling through modulating signal transduction of TGF-β/Smads pathway (14). Recently, there is evidence showing that HMP plays roles in inhibiting atherosclerosis development (12,15). However, the exact anti-atherosclerotic mechanisms of HMP are not fully understood. Here, we investigated the atheroprotective effects of HMP using an animal model of atherosclerosis induced by western-type diet (WTD) and determined the potential underlying mechanisms of action.

Methods

Animal procedure

Adult 24 male homozygous apolipoprotein E knockout (apoE<sup>−/−</sup>) mice at 8 weeks with C57BL/6 background were purchased from Vital River Laboratory Animal Technology Co., Ltd. (Beijing, China). Before the experiments, mice were acclimatized for 2 weeks and housed separately in individually ventilated cages. All animals were maintained in a controlled temperature (22±2 °C) under a 12-h light/dark cycle with ad libitum access to standard food and water. Then, apoE<sup>−/−</sup> mice at 10 weeks old were fed a WTD (containing 21% fat and 0.15% cholesterol) and randomly allocated into two groups (n=12 each group). One group (HMP) was administered 25 mg/kg/day HMP (provided by Shanghai Hutchison Pharmaceuticals, Shanghai, China, batch number 181107) suspended in normal saline intragastrically (13,16,17). Another group (WTD) was intragastrically treated with the same volume of normal saline per day. After 20 weeks of drug delivery, all mice were fasted overnight and euthanized by injecting pentobarbital sodium intraperitoneally (Figure 1).

Blood samples were acquired from the orbital veins and centrifuged at 3,000 rpm at 4 °C for 10 min. The top serum layer was collected and stored at −80 °C until use. The residual blood in solid tissues was removed by whole-body perfusion with ice-cold 0.9% normal saline intragastrically (13,16,17). Another group (WTD) was intragastrically treated with the same volume of normal saline per day. After 20 weeks of drug delivery, all mice were fasted overnight and euthanized by injecting pentobarbital sodium intraperitoneally (Figure 1).

Characterization of aortic atherosclerotic plaques

The full-length aorta from the root to the common iliac
artery was rapidly removed and the extent of atherosclerotic lesions in the whole aorta lumen surface was assessed by Oil Red O staining (Sigma, St. Louis, USA). En face of the entire aorta was photographed and the atheroma plaque area was measured using an image analysis program (Image-pro plus 6.0 software, Media Cybernetics, USA). Atherosclerosis severity was determined as the plaque lesion size relative to the inner surface area of the total aorta.

The aorta segments were fixed in 4% paraformaldehyde solution and embedded in paraffin. Afterwards, the aorta tissues were serially cut into 5-μm thick cross-sections at 50-μm intervals. A set of paraffin sections were stained with hematoxylin and eosin (H&E) for morphological analysis of the atherosclerotic lesions. Subsequently, the collagen content within the plaque area was visualized by Masson's trichrome staining. Moreover, immunohistochemical detection of cross-sections was performed to quantify the levels of monocyte macrophages 2 (MOMA-2), α smooth muscle actin (α-SMA) (Abcam, Cambridge, UK) and cleaved caspase-3 (Cell Signaling Technology, MA, USA). Quantitative analysis of the indicators was performed with the Image-pro plus 6.0 software.

**Histomorphometric evaluation of liver and intestine**

Liver and intestine tissues were excised and fixed in 4% paraformaldehyde solution. The tissues were then dehydrated and embedded in paraffin. Serial 5-μm thick cross-sections were harvested and stained with H&E following standard procedures.

The liver tissue samples were embedded in optimal cutting temperature compound and sectioned into 8-μm consecutive sections using a cryostat. The sections were stained with Oil Red O for the evaluation of lipid content in liver tissues.

**Detection of lipid profiles**

Serum concentrations of triglyceride (TG), total cholesterol (TC), low density lipoprotein cholesterol (LDL-C) and high-density lipoprotein cholesterol (HDL-C) were determined with commercially available reagents according to the manufacturer’s protocols (Jiancheng Bioengineering Institute, Nanjing, China).

For quantifying hepatic and fecal lipid levels, the liver tissue and fecal particles were homogenized with ethyl alcohol. The homogenate was then centrifuged and the top liquid was collected for measuring TG and TC contents using biochemical kits (Jiancheng Bioengineering Institute, Nanjing, China).

**Quantification of circulating bioactive factors**

Concentrations of inflammatory cytokines in the serum including TNF-α, IL-1β, IL-6, and CRP were quantified using the ELISA kits (Boster Biological Technology, Wuhan, China). Moreover, the circulating NO content was evaluated by biochemical kits (Jiancheng Bioengineering Institute, Nanjing, China). Briefly, 100 μL of serum sample was mixed uniformly with 300 μL of pre-processing reagent using a vortex finder. Then the mixture was centrifuged at 4,000 rpm for 15 min at room temperature and the supernatant was collected. Afterwards, 160 μL of the obtained supernatant was incubated with 80 μL of the chromogenic agent for 15 min. The absorbance was detected at 560 nm on a microplate reader.

**Western blot assay**

For protein extraction, the liver, intestine and aorta tissues were lysed with RIPA lysis buffer supplemented with protease and phosphatase inhibitors. The concentrations...
of extracted proteins were tested using bicinchoninic acid assay. After denaturation, proteins were loaded on 10% sodium dodecyl sulfate-polyacrylamide electrophoresis gel for separation. The separated proteins were then transferred onto polyvinylidene difluoride membranes, which were sequentially blocked with 5% BSA dissolved in TBS-T for 1 h at room temperature and probed with appropriate primary antibodies overnight at 4 °C. The membranes were then washed with TBS-T and incubated with horseradish peroxidase-conjugated secondary antibodies at room temperature for 1 h. Then protein bands were visualized using an enhanced chemiluminescence detection reagent and the acquired images were analyzed with the Image J software (NIH, USA). The primary antibodies against sterol regulatory element binding protein 2 (SREBP2), 3-hydroxy-3-methylglutaryl-coenzyme A reductase (HMGCR), LDL receptor (LDLR), proprotein convertase subtilisin/kexin type 9 (PCSK9), peroxisome proliferator-activated receptor α (PPARα) and SREBP1c were obtained from Abcam (Cambridge, UK). Antibodies against c-Raf, p-c-Raf, ERK1/2, p-ERK1/2, AMPK, p-AMPK, calcium/calmodulin-dependent kinase kinase β (CaMKKβ), p-CaMKKβ, Bcl-2, Bax, Bad, Apaf-1, caspase-9 and caspase-3 were purchased from Cell Signaling Technology (MA, USA). Anti-liver kinase B1 (LKB1), anti-p-LKB1, anti-Niemann-pick C1-like 1 (NPC1L1) and anti-Bcl-xl were obtained from Santa Cruz Biotechnology (CA, USA). Moreover, anti-p-HMGCR and anti-camitine palmitoyltransferase 1A (CPT-1A) were obtained from Bioss (MA, USA). Anti-ATP-binding cassette transporter G5 (ABCG5) and anti-ABCG8 were offered by Affinity Biosciences (OH, USA). Antibodies against 7alpha-hydroxylase (CYP7A1), fatty acid synthase (FAS), acetyl-CoA carboxylase α (ACCα), p-ACCα, cytochrome-c and GAPDH were purchased from Abclonal (Boston, USA).

**Real-time quantitative PCR**

Total RNA was isolated from the frozen hepatic and intestinal tissues with RNAiso Plus (Takara, Japan). The concentration of extracted RNA was determined using a Beckman Coulter DU730. Complementary DNAs were synthesized from total RNA templates using a PrimeScript™ RT reagent (Takara, Japan). Then mRNA expression level of each gene was detected by quantitative real-time PCR using TB Green kits (Takara, Japan). mRNA contents of target molecules were normalized to that of GAPDH, which was selected as an internal standard. Primers used here were listed in Table 1.

<table>
<thead>
<tr>
<th>Gene</th>
<th>Primer sequence (sense 5’-3’)</th>
<th>Primer sequence (antisense 5’-3’)</th>
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<tr>
<td>HMGCR</td>
<td>CGCTCTTGTGGAAAGCTTGG</td>
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<td>PCSK9</td>
<td>TTTGCTCTGCCCAGACAT</td>
<td>GTGACCCTGCCCCTAATCCTC</td>
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<td>LDLR</td>
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<tr>
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<td>AGCCATGAAAGCCCTCAAA</td>
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<td>TTGGAGCTAAGGAGATG</td>
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<td>GAPDH</td>
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<td>GATGGGCTTCCGGTGTGTA</td>
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Statistical analysis

All data were presented as the mean ± standard deviation and data analysis was performed using the SPSS software version 21.0. The Kolmogorov-Smirnov test and Levene test was used to analyze the normality and homogeneity of variances of data, respectively. Student’s t-test was applied to compare the significance of differences between two groups. P values less than 0.05 (two-tailed) were considered to be statistically significant.

Results

HMP ameliorated the severity of atherosclerotic lesions

The lesion area in the aortic inner surface was quantified by Oil Red O staining and we observed that the atherosclerotic lesion range in three aortic segments of HMP-treated mice was dramatically decreased in comparison with those of the WTD mice (P<0.05 for aortic arch, P<0.05 for thoracic aorta, P<0.01 for abdominal aorta) (Figure 2A). Moreover, the analysis of H&E-stained aortic cross-sections revealed that HMP administration effectively alleviated the atherosclerotic burden, as seen by decrease of the percentage of plaque size to total aortic lumen area (P<0.05) and lesion thickness reduction at the site of atheromatous plaques (P<0.05) (Figure 2B).

We then assessed the effects of HMP on regulating the stability of atherosclerotic lesions. Histopathological detection showed that the plaque lesion in mice with HMP gavage possessed smaller necrotic core (P<0.01) and higher collagen content (P<0.01) compared with vehicle-treated mice, as indicated by H&E staining and Masson’ trichrome staining (Figure 3A,B). Immunohistochemical test discovered that the lesion area from HMP-treated samples had reduced expression of MOMA-2 (P<0.05), which was a specific marker for macrophages, suggesting the decrease of macrophage accumulation in aortic plaques (Figure 3C). Additionally, the content of smooth muscle cell was increased in the plaques of HMP-treated mice (P<0.05), as assessed by α-SMA staining (Figure 3D). In view of above findings, HMP was proved to encumber atherosclerosis progression and strengthen the stability of plaque lesions.

HMP displayed effective effects on mediation of serum lipid spectrum

To analyze the mechanisms by which HMP reduced atherosclerosis progression, we first examined the effect of HMP on blood lipid profiles, given that dyslipidemia was an early pathological event responsible for atherogenesis (2). As showed in Figure 4A, HMP treatment lowered TG (P<0.05), TC (P<0.01) and LDL-C (P<0.01) contents, while elevating serum HDL-C level (P<0.05) when compared with normal saline management.

HMP markedly mitigated the extent of lipid accumulation in the liver

We then evaluated the effects of HMP on WTD-induced lipid disposition in the liver. Oil Red O staining of liver cryosections showed that numerous lipid droplets existed in liver tissues of the WTD-fed mice and HMP intervention significantly decreased the number and size of lipid droplets in hepatic tissues. Moreover, the cytoplasm of hepatocytes from vehicle-treated mice exhibited many fat vacuoles, as evidenced by H&E staining. We discovered that the severity of vesicular steatosis of hepatocytes was alleviated in the HMP-treated mice (Figure 4B). Similar to histological detection results, TG (P<0.05) and TC (P<0.05) contents in liver tissues were dramatically decreased in the HMP-treated mice when compared to those in the WTD-fed mice (Figure 4C).

HMP regulated signal molecules related with lipid metabolism in hepatic tissues

Since the liver was a pivotal organ involved in lipid metabolism (18), we evaluated whether HMP improved lipid profiles via regulating relevant signal molecules in the liver of apoE−/− mice. We observed that HMP administration diminished the expressions of SREBP2 (P<0.01) and downstream molecules including HMGCR (P<0.05) and PCSK9 (P<0.05), which regulated cholesterol synthesis and LDLR degradation, respectively. LDLR responsible for LDL-C uptake by hepatocytes was another SREBP2-modulated downstream effector (19). However, LDLR was found to be upregulated in hepatocytes of the HMP group (P<0.05) (Figure 5A). We further analyzed the Raf/ERK1/2 pathway which regulated LDLR mRNA stability (20). We found that HMP treatment enhanced the activities of Raf and ERK1/2 (P<0.05) (Figure 5B), hinting that HMP might enhance the stability of LDLR mRNA. Moreover, no difference in CYP7A1 level (P>0.05), which regulated cholesterol bioconversion, was found in liver tissues between the two groups (Figure 5C). PPARα/CPT-1A pathway was reported to facilitate β-oxidation of fatty acids (21). Whereas,
Figure 2 Heart-protecting musk pill (HMP) administration attenuated the progression of atherosclerosis in apoE−/− mice. (A) Representative images of plaque lesions in the area of aortic inner surface stained with Oil Red O. The arrows indicated the plaques in the aortic inner surface. (B) H&E staining assessed the extent of plaque-induced luminal stenosis and the degree of lesion hyperplasia in three segments of the entire aorta. The arrows showed the plaques in the cross-section. Data were expressed as the mean ± SD, n=5. *P<0.05, **P<0.01 vs. WTD group. Scale bar =200 μm.
SREBP1c and its target molecules FAS and ACCα were key factors involved in *de novo* lipogenesis (19,22). Here, we discovered that HMP might trigger TG catabolism and weaken TG anabolism, as seen by level elevation of PPARα (P<0.01) and CPT-1A (P<0.05) and content decrement of SREBP1c, FAS and ACCα (P<0.05) (Figure 5D,E).

Furthermore, AMPK, which mediated the up-regulation of PPARα and down-regulation of SREBPs (23), was found to present enhanced activities in hepatic tissues of HMP-treated mice (P<0.05) (Figure 5F). HMGCR and ACC, acting as the phosphorylation substrates of AMPK, presented a noticeable elevation in the phosphorylated level of the HMP group (P<0.05) (Figure 5A,E). Moreover, we observed that HMP strengthened the activity of LKB1

**Figure 3** The effects of heart-protecting musk pill (HMP) on the stability of atherosclerotic lesions in the aortic root. (A) The size of necrotic core in the plaque area was determined by H&E staining. The arrows showed the necrotic core in the plaques. (B) The collagen content was detected with Masson’s Trichrome staining. The arrows indicated the area of collagen. (D) Immunostaining for MOMA-2 (C) and α-SMA (D) in the lesion area. The arrows indicated the representative area. Results were presented as the mean ± SD, n = 5. *P<0.05, **P<0.01 vs. WTD group. Scale bar =200 μm.
responsible for AMPK activation (P<0.05), yet had no impact on the activity of CaMKKβ (P>0.05), which was another upstream AMPK kinase (Figure 5F).

As shown in Figure 5G, HMP reduced HMGCR and PCSK9 mRNA content and raised LDLR mRNA level (P<0.05). Meanwhile, compared with the WTD mice, mRNA expressions of FAS and ACCα (P<0.05) were decreased and levels of CPT-1A and peroxisome proliferator-activated receptor gamma coactivator 1-α (PGC-1α) mRNA (P<0.05) were increased in liver tissues of HMP-administered mice. In addition, there were no significant differences in levels of CYP7A1, sterol-12α-hydroxylase (CYP8B1), acyl CoA oxidase (ACOX) and stearoyl-CoA desaturase 1 (SCD-1) mRNA (P>0.05) between the two groups.

**HMP affected levels of factors involved in lipid transportation in intestinal tissues**

Intestine was found to affect blood lipid profiles via regulating lipid absorption and excretion (24). We found that HMP raised TC content (P<0.05) but did not affect TC level (P>0.05) in feces of apoE−/− mice (Figure 6A). Additionally, no significant differences in food intake and excrement weight were observed between the two groups (P>0.05) (Figure 6B). H&E staining revealed that HMP oral gavage did not alter the architecture and morphology of intestinal tissues (Figure 6C). Then, we discovered that HMP treatment increased the expression of cholesterol transporter ABCG5 and ABCG8, yet reduced both NPC1L1 mRNA and protein levels (P<0.05) (Figure 6D,E), implying that HMP probably elevated fecal TC content via mediating transintestinal cholesterol migration.

**HMP decreased levels of circulating inflammation-promoting cytokines**

Since inflammation was a vital contributor to atherosclerosis initiation and progression by inducing vascular endothelial injury (25,26), we hypothesized that HMP improved the
Figure 5 Heart-protecting musk pill (HMP) affected the expressions and activities of molecules participating in hepatic lipid metabolism. (A) Protein levels of SREBP2 and downstream effectors involved in cholesterol biosynthesis and uptake. (B) Signal transduction of Raf/ERK1/2 pathway was examined by western blot. The protein expression of CYP7A1 (C), PPARα and CPT-1A (D). (E) Protein levels of SREBP1c and downstream factors involved in de novo lipogenesis. (F) Total protein content and corresponding phosphorylated level of AMPK, LKB1 and CaMKKβ. (G) The mRNA expressions of effector genes associated with cholesterol (left part) and fatty acid (right part) metabolism. Results were presented as the mean ± SD, n=6. *P<0.05, **P<0.01 vs. WTD group.
hyper-inflammatory state in vivo. To validate this hypothesis, serum cytokine levels were quantified. We found that levels of serum TNF-α, IL-1β, IL-6 and CRP were dramatically decreased in response to HMP intervention (P<0.05) (Figure 7A).

**HMP bad beneficial roles against vascular intimal injury**

As vascular intimal damage augmented extension of atheroma plaques (27), we assessed the impact of HMP on tunica intima injury. As shown in Figure 7B, there was a content decrement of cleaved caspase-3 in the arterial intima of HMP-administered mice (P<0.05). Then, results of western blot suggested that HMP raised the Bcl-2/Bax ratio and Bcl-xl level (P<0.01) and reduced Bad content (P<0.05) in aortic tissues (Figure 7C). The cytoplasm cytochrome-c, necessary for apoptotic signal transduction, was found to exhibit a lowered content after HMP treatment (P<0.01). Meanwhile, HMP reduced levels of Apaf-1 and decreased contents of cleaved caspase-9 and cleaved caspase-3 (P<0.05) (Figure 7D). These findings suggested that HMP effectively suppressed mitochondrial-mediated apoptotic pathway. Moreover, NO, a typical pro-survival substance inhibiting apoptosis-related pathways, was upregulated by HMP.
Figure 7 Oral gavage of heart-protecting musk pill (HMP) displayed protective effects against vascular wall injury. (A) Concentrations of serum TNF-α, IL-1β, IL-6 and CRP were detected by ELISA kits. (B) The expression of cleaved caspase-3. The arrows showed the positive area of cleaved caspase-3. (C) Protein levels of Bcl-2 family members in the aortic tissue. (D) The contents of signal proteins associated with mitochondrial-dependent apoptosis pathway. (E) The concentration of serum NO. Data were expressed as the mean ± SD, n=6 for (A) and (E), n=4 for (B-D). *P<0.05, **P<0.01 vs. WTD group. Scale bar =50 μm.

Discussion

Pharmacological studies have demonstrated that several kinds of TCMs are effective in suppressing the development of atherosclerotic lesions (28). Consistent with previous studies, HMP, originating from the TCM Suhexiang pill, was found to decrease atheroma plaque burden in apoE<sup>−/−</sup> mice. For the first time, we discovered that atheroprotective mechanisms of HMP might be due to improvement of lipid metabolism and alleviation of vascular intimal injury (Figure 8).

In comparison with plaque enlargement-induced luminal stenosis, the plaque rupture-triggered arterial occlusion is more prone to cause life-threatening clinical events, such as acute myocardial infarction and cerebral stroke (29). Vulnerable plaques are characterized by large lipid-rich necrotic cores, decreased collagen amount, increased macrophage accumulation and smooth muscle cell reduction (30). According to these traits, HMP appeared to effectively enhance the stability of atheromatous lesions, as evidenced
by shrink of necrotic core, escalation of collagen content, attenuation of macrophage infiltration and increase of smooth muscle cell, suggesting that HMP had therapeutic potentials for improving plaque destabilization.

Hypercholesterolemia has been demonstrated to be a pathological condition related to atherogenesis acceleration (2). Our results showed that HMP-treated mice presented TC and LDL-C level reduction in the bloodstream. Thereafter, we further explored the relevant mechanisms required for regulatory actions of HMP on cholesterol metabolism. HMGCR, regarded as a cholesterol synthesis rate-limiting enzyme, is targeted by several cholesterol-lowering drugs. LDLR acts as a membrane glycoprotein in hepatocytes and is responsible for removing circulating LDL-C. Moreover, PCSK9 is a serine protease inducing the lysosomal degradation of LDLR (23,24). It has been illustrated that these molecules are regulated by the same upstream signal protein SREBP2 (19). In this study, we identified that HMP potently lowered SREBP2 expression, followed by HMGCR and PCSK9 content reduction. It should be noted that the LDLR mRNA and protein levels were elevated in the HMP group compared to those in the WTD group. We speculated that this discrepant result might be attributed to HMP-induced activation of ERK1/2 pathway, given that ERK1/2 cascade could affect mRNA 3’-UTR structure which controlled stabilization of LDLR mRNA and several drugs including berberine and triciribine had been proved to increase the stability and content of LDLR mRNA via stimulating ERK1/2 pathway through SREBP2-independent processes (20,31,32). Moreover, as PCSK9 regulated the elimination of LDLR protein (16), level reduction of PCSK9 induced by HMP treatment might lead to decrease in LDLR degradation in the lysosome compartment, thereby increasing the stability and content.
of LDLR protein. Thus, through modulating SREBP2-regulated pathway and ERK1/2 cascade, HMP might inhibit cholesterol production and accelerate cholesterol uptake in hepatocytes, which resulted in TC and LDL-C concentration reduction in the bloodstream, consequently improving risks of atherosclerosis progression.

It has been proved that atheroprone effects of TG overload are largely due to the induction of inflammation, aggregation of endothelial dysfunction and impairment of fibrinolysis (33). Our results showed that HMP had favorable impacts on lowering blood TG level and improving lipid accumulation in liver tissues. Next, relevant mechanisms necessary for regulatory effects of HMP on TG content were assessed. Fatty acid β-oxidation is a biological process that enables TG consumption and nuclear receptor PPARα has pivotal roles in this process. Once stimulated, PPARα promotes the expression of target gene CPT-1A, which catalyzed long-chain acyl-CoA migration from the cytosol to the mitochondrial matrix for oxygenolysis (21). We observed that HMP induced the level increase of PPARα, followed by up-regulation of CPT-1A. It is verified that de novo lipogenesis is an efficient way favoring TG level increase. SREBP1c is a transcriptional factor that possesses crucial roles in de novo lipogenesis via mediating expressions of downstream lipogenic gene FAS and ACCα. There is evidence that high expression of SREBP1c causes circulating TG accumulation and hepatic steatosis (34,35). Our results showed that HMP notably reduced the level of SREBP1c and the expression of FAS and ACCα. Hence, mediation of signal pathways involved in lipid β-oxidation and de novo lipogenesis might contribute to the regulatory roles of HMP in TG concentration of WTD-fed apoE−/− mice.

AMPK is an intracellular energy sensor that exerts key effects on lipid and glucose metabolism (23). When activated, AMPK switches off anabolic activities including cholesterol and fatty acid biosynthesis, and switches on catabolic processes like fatty acid oxidation through regulating relevant signaling factors, such as up-regulation of PPARα and decrease of SREBP2 and SREBP1c level (36). Similar to results reported by other studies, the enhanced activity of AMPK was seen in the hepatocytes of HMP-treated mice with improved lipid profiles. Moreover, HMGCR and ACCα are well identified downstream targets of AMPK and a pivotal mechanism underlying AMPK-induced lipid production inhibition is due to the induction of phosphorylation of HMGCR and ACCα, the inactive form of the two enzymes (23). Our findings showed that, consistent with AMPK modification, the phosphorylated HMGCR and ACCα levels were markedly increased in the HMP group. Regarding the upstream kinases responsible for AMPK activation, LKB1 and CaMKKβ are two major upstream regulators controlling Thr-172 phosphorylation on the α-subunit of AMPK. It has been verified that LKB1 and CaMKKβ mainly activate AMPK in response to energy stress increased intracellular calcium, respectively (36,37). In this study, p-LKB1 rather than p-CaMKKβ showed the same tendency of level changes with p-AMPK in the hepatocytes of HMP-treated mice, indicating that HMP might affect LKB1-mediated AMPK activation to regulate pathways implicated in lipid metabolism, eventually leading to improvement of blood lipid profiles.

Except for the liver, the intestine is regarded as an important organ responsible for lipid metabolism. By modulating dietary lipid absorption and internal lipid excretion, the intestine indirectly impacts the regulation of circulating lipid profiles (24). We found that HMP raised the fecal TC content, without changing TG level. It is reported that cholesterol absorption from the intestinal lumen is controlled by NPC1L1 in the enterocytes. Several atherogenic factors have been found to increase the expression of intestinal NPC1L1 and then elevate the levels of blood cholesterols (24,38). In addition, level reduction of NPC1L1 is reported to be an effective way to inhibit cholesterol absorption, thereby ameliorating dyslipidemia and suppressing atherosclerosis progression (39,40). There is evidence that cholesterol secretion from enterocytes to the gut lumen is regulated by apical transporters ABCG5 and ABCG8 (16,24). Findings from previous researches demonstrate that up-regulation of ABCG5 and ABCG8 is able to exert atheroprotective roles via enhancing cholesterol excretion and weakening its influx, which might partly mitigate the accumulation of cholesterol in the circulation and then improve the development of atherosclerosis.

Upon pathological stimulation, vascular endothelial cells (ECs) produce an apoptotic phenotype, which causes intima injury, followed by progression of atheromatous lesions (7,27). Our results revealed that HMP gavage alleviated ECs apoptosis in the area of aortic wall. As inflammation could induce atherogenesis by causing vascular intimal damage (25,42), we analyzed the effects of HMP on inflammation development and found that HMP reduced...
levels of TNF-α, IL-1β, IL-6, and CRP in the bloodstream. Furthermore, NO, as a bioactive factor responsible for cellular survival (43), was found to be upregulated in the circulation of HMP-treated mice. These data elucidated the vasoprotective properties of HMP. The Bcl-2 family contains regulatory proteins that either trigger or suppress apoptosis. Among them, Bax oligomerizes and forms pores that permeabilize the mitochondrial outer membrane and cause leakage of apoptogenic factors. In contrast, Bcl-2 forms heterodimers with Bax to prevent Bax oligomerization. Moreover, Bcl-xl is capable of repressing Bax activation, whereas the function of Bad is to inhibit activities of anti-apoptotic proteins (44). We discovered that HMP treatment induced increment of Bcl-2/Bax ratio and Bcl-xl content and triggered decrement of Bad level. Once released in the cytosol, the apoptogenic molecules, particularly cytochrome-c, sequentially activate Apaf-1 and downstream caspase-9, followed by caspase-3 cleavage and apoptosis initiation (44). In our study, there was a decrease in levels of cytoplasm cytochrome-c, Apaf-1 and cleaved caspase-9, 3 in the arterial walls of the HMP group. These data verified that HMP possibly displayed anti-atherogenic actions via improving vascular intimal injury by abating mitochondrial-related apoptosis signaling cascade. However, it is not clear whether HMP directly improves vascular damage or not. Moreover, the specific mechanisms by which HMP inhibits the activation of apoptotic pathway in the arterial wall are poorly understood. These deficiencies are needed to be settled using the in vitro model in the near future.

In summary, our study demonstrates that HMP intervention considerably ameliorates the development of atheromatous lesions in apoE−/− mice fed by WTD and the atheroprotective effects are due to the regulation of lipid metabolism as well as inhibition of vascular intimal damage. These data provide novel insights into the molecular mechanisms underlying HMP attenuates atherosclerosis-related diseases.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. All animal procedures were approved by the Institutional Animal Care and Use Committee of Tongji Medical College, Huazhong University Science and Technology, Wuhan, China. All experiment procedures were conformed to the National Institutes of Health Guide for the Care and Use of Laboratory Animals.

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