



factors (Tsai et al., 2014), especially high fat intake (Rizvi et al., 2003). Many reports have demonstrated that obesity is linked to an obvious oxidative stress (Furukawa et al., 2004; Matsuzawa-Nagata et al., 2008). Charradi et al. (2013) found that obesity is strongly related to systemic oxidative stress and was in particular concerned with the relationship between altered glutathione homeostasis and organ lipotoxicity associated with diet-induced obesity. Moreover, antioxidant supplementation showed beneficial effects in preventing dyslipidemia disease caused by a high-fat diet (HFD) (Gorinstein et al., 2006). Oxidative stress is an imbalance between oxidants and antioxidants (Sies, 1997). It can result in DNA hydroxylation, protein denaturation, lipid per-oxidation, apoptosis, tissue damage (Valko et al., 2006), and a disruption in the redox signaling pathway (Kamata and Hirata, 1999).

Several strategies have been used to prevent obesity, including diet control, exercise, behavior therapy and medication. Probiotics, which are defined as “live microorganisms that, when administered in adequate amounts, confer a health benefit on the host” (Araya et al., 2002), could modulate the intestinal flora and have some anti-oxidative and lipid-lowering abilities (Endo et al., 2013; Everard et al., 2014).

*Lactobacillus* is an intestinal microorganism that may grow in the gut and consume oxygen to maintain an anaerobic environment for the prevention or therapy of gastrointestinal disorders (Hu et al., 2014). In addition, *Lactobacillus* is preferred due to its higher resistance to harsh environments and capacity for long-term storage at ambient temperature (Hong et al., 2005). Previously, we have reported that

*Lactobacillus* B10 possessed positive effects on the anti-oxidative defense of cells (Li et al., 2013) and those findings lead us to further investigate the regulatory effect of *Lactobacillus* B10 on the anti-oxidative and lipid-lowering capacities of HFD-induced obese mice.

## 2 Materials and methods

### 2.1 Bacterial strain

B10 powder ( $10^8$  colony-forming unit (CFU)/g) was prepared by Microbiology and

Genetic Engineering Laboratory, Institute of Feed Sciences, Zhejiang University, China. B10 strain was cultured on Luria-Bertani media, kept at 37 °C for 24 h and shaken at 180 r/min. Pure bacterial cells were collected after centrifugation at 5000 rpm for 10 min at 4 °C and cooling. Thereafter, these cells were washed twice with sterile 0.85% (8.5 g/L) sodium chloride solution. Ultimately, the culture purity and identification were constantly checked by the spreading plate method (Nikoskelainen et al., 2003).

### 2.2 Animals and diets

Thirty male ICR mice (7–8 weeks old,  $n=15$  per group) were obtained from the Institutional Animal Care and Use Committee of Zhejiang University, China. All mice were housed in standard plastic cages (three mice per cage) and maintained under a 12-h light-dark cycle at constant temperature and humidity ( $(23\pm 1)$  °C and  $(55\pm 5)\%$ , respectively). One group of mice was fed with an HFD (92.8% normal diet, 2% cholesterol, 5% lard, and 0.2% cholate) and the other was kept on an HFD supplemented with 0.1% (w/w) B10 powder for 30 d. The normal diet was provided by Shanghai SLAC Laboratory Animal Co., Ltd. (Shanghai, China) and its nutrient components level is listed in a supplemental material Table S1. During the preparation of the B10 powder, starch was used to dilute B10 and the same amount of starch was also added to the HFD group to compensate for the difference in nutrient composition of the diets. All the mice were fed *ad libitum* and food intake was recorded. The experiment was approved by and performed in accordance with the guidelines of the local ethics committee.

### 2.3 Sample collection

After 30 d, mice ( $n=6$ ) from each group (HFD and B10) fasted for 12 h, and then were anesthetized with diethyl ether. Mice were killed by cervical dislocation and the blood samples were taken from the inferior vena cava. After incubation at 4 °C for 30 min and centrifugation at 2000 rpm for 20 min, the serum was obtained (Centrifuge 5804R, Eppendorf, Germany). Liver samples were dissected out and placed in liquid nitrogen. Samples were finally frozen and kept at  $-80$  °C for no more than 2 months until further analysis.

## 2.4 Biochemical assays of serum

The contents of glucose, triglyceride (TG), total cholesterol (TC), low-density lipoprotein cholesterol (LDL-C), high-density lipoprotein cholesterol (HDL-C), blood urea nitrogen (BUN), glutamic oxaloacetic transaminase (GOT), and glutamic pyruvic transaminase (GPT) in serum were determined by spectrophotometry according to the kit manufacturer's instructions. The methods of those kits are presented in the supplemental material Data S1. The materials of glucose measurement were purchased from Rongsheng Bio-pharmaceutical Co., Ltd., Shanghai, China. Kits for serum TG and TC were purchased from Saike Bio-technology Co., Ltd., Ningbo, China. Kits for serum BUN, GOT, and GPT were bought from Nanjing Jiancheng Bioengineering Institute, Nanjing, China. Materials for serum LDL-C and HDL-C were purchased from Beihuakangtai Clinical Reagent Co., Beijing, China.

## 2.5 Biochemical assays of the liver

Liver samples were homogenized with ice-cold physiologic saline (1:10, v/v) and centrifuged at 2000 for 10 min (Centrifuge 5804R, Eppendorf, Germany). Supernatant was collected for determination of the total anti-oxidant capability (T-AOC), anti-superoxide anion activity (ASOA), concentrations of TC, TG, LDL-C, HDL-C, 8-hydroxy-2'-deoxyguanosine (8-OHdG), glutathione (GSH), and thioredoxin reductase (TrxR), and the activities of superoxide dismutase (SOD), catalase (CAT), glutathione peroxidase (GSH-Px), and xanthine oxidase (XO). Kits for hepatic TC and TG were obtained from Saike Biological Technology Co., Ltd., Ningbo, China. Kits for hepatic LDL-C and HDL-C were purchased from Beihuakangtai Clinical Reagent Co., Beijing, China. Kits for SOD, CAT, GSH-Px, XO, T-AOC, ASOA, and GSH were purchased from Nanjing Jiancheng Bioengineering Institute, Nanjing, China. All the above parameters were determined by spectrophotometry according to the manufacturers' instructions. Enzyme-linked immunosorbent assay (ELISA) kits for 8-OHdG and TrxR were purchased from Bioleaf Biological Co., Ltd., Shanghai, China.

## 2.6 RNA extraction and quantitative real-time polymerase chain reaction (PCR)

Liver tissues of approximately 100 mg were used for RNA extraction. RNA was extracted using

the RNAiso plus method (TaKaRa, Dalian, China) according to the instructions of the manufacturer. Quantitative and qualitative analyses of isolated RNA were assessed from the ratio of absorbance at 260 and 280 nm by NanoDrop spectrophotometer (Thermo Scientific, Wilmington, DE, USA) and agarose gel electrophoresis (Sangon Biotech, Shanghai, China). Complementary DNA (cDNA) was synthesized from 1 µg of total RNA using M-MLV reverse transcriptase (TaKaRa, Dalian, China). Transcriptional changes were then identified by quantitative PCR (qPCR), which was performed using the Premix Ex Taq™ with SYBR Green (TaKaRa, Dalian, China) as per the manufacturer's instructions, and the ABI 7500 Fast Real-Time PCR system (Applied Biosystems, Carlsbad, CA, USA). The thermocycle protocol lasted for 30 s

**Table 1 Primers used for real-time PCR**

Gene	Primer sequence
$\beta$	F: 5'-GTTGTCTCCTGCGACTTCA-3'
	R: 5'-GCCCCCTCCTGTTATTATGG-3'
	F: 5'-CCTGTACGCCAACACAGTGC-3'
	R: 5'-ATACTCCTGCTTGCTGATCC-3'
	F: 5'-CAAGAGGTTGATAGAGCGTTT-3'
	R: 5'-TTGTTCTCGGGACTTTGGA-3'
$\alpha$	F: 5'-GGCATCCTAACCTCCAACCA-3'
	R: 5'-TGCTCATAACCCATCCACA-3'
$\beta$	F: 5'-CTTTACATCAGTGTTTCCGTCAG-3'
	R: 5'-TCAAATTGCCACCGTTCTT-3'
$\gamma$	F: 5'-GGTGACCCTCCTCAAGTATGG-3'
	R: 5'-GGTGTCTGGATGGCTTCTA-3'
	F: 5'-AGAACCTCGGACTCCACCTTA-3'
	R: 5'-CCACAGACTCGGCACTCAAT-3'
	F: 5'-TACAAGAAGACCCATAAGAATC-3'
	R: 5'-CCACAGACTCGGCACTCAAT-3'
	F: 5'-GGACCGTGCTCCCTTCTTAG-3'
	R: 5'-GAGCTCTTCGTGGGTTCTGT-3'
F: 5'-GGAAACCAGAAGCCGAACA-3'	
R: 5'-GACAAGGAATAGAAGGGACAGAT-3'	
F: 5'-GTAGGAAGCCCACCACGAC-3'	

at 95 °C, followed by 40 cycles of 5 s denaturation at 95 °C, 34 s annealing/extension at 60 °C, and then a final melting curve analysis to monitor purity of the PCR product. Primer sequences for the mouse genes were designed and selected by Primer 5.0 and Oligo 7.0 softwares and the sequences are presented in Table 1. The  $2^{-\Delta\Delta q}$  method was used to estimate mRNA abundance.  $\Delta q$  is  $q_{target} - q_{reference}$  and  $\Delta\Delta q$  is  $\Delta q_{treatment} - \Delta q_{control}$ . Relative gene expression levels were normalized to those of the eukaryotic reference genes glyceraldehyde-3-phosphate dehydrogenase ( $\beta$ ) and  $\beta$ .

### 2.7 Statistical analysis

Data are expressed as mean±standard deviation (SD) and were analyzed using the one-way analysis of variance (ANOVA) procedure of SPSS 16.0 (SPSS Inc., Chicago, IL, USA) for Windows by the Tukey test and the variable used is the administration of B10. Differences were considered statistically significant at  $P < 0.05$ .

## 3 Results

### 3.1 Effects of dietary supplementation of *Bacillus subtilis* B10 on body weight and food intake of mice

While the final body weight and average daily weight gain for HFD+B10 group significantly ( $P < 0.05$ ) decreased compared with the HFD group,

no significant differences in daily food intake were observed between the two groups (Table 2).

### 3.2 Effects of dietary supplementation of *Bacillus subtilis* B10 on lipid profile in the serum and liver of mice

Hepatic TC and TG of HFD-fed mice were effectively ( $P < 0.05$ ) decreased in B10-treated mice. In addition, the TC, LDL-C, and HDL-C levels in serum of the B10 treatment group were significantly ( $P < 0.05$ ) increased (Table 3). Besides, B10 supplementation also exhibited a significant decrease in the serum glucose concentration and GOT and GPT activities compared with the HFD group, while serum BUN was unaffected between the two groups (Table 4).

### 3.3 Effects of dietary supplementation of *Bacillus subtilis* B10 on mice hepatic mRNA expression of genes associated with lipid metabolism of mice

The administration of B10 down-regulated the transcript levels of 3 $\beta$ -hydroxysteroid- $\Delta$ 24 reductase ( $\beta$ ) significantly ( $P < 0.05$ ) and up-regulated that of peroxisome proliferator-activated receptor  $\alpha$  ( $\alpha$ ) and 3-hydroxy-3-methylglutaryl-CoA synthase 2 ( $\beta$ ) remarkably ( $P < 0.05$ ). No significant differences were noted in the gene expression levels of carnitine acyltransferase I ( $\beta$ ), peroxisome proliferator-activated receptor  $\beta$  ( $\beta$ ), and peroxisome proliferator-activated receptor  $\gamma$  ( $\gamma$ ) between the two groups (Fig. 1).

**Table 2** Effects of dietary supplementation of *Bacillus subtilis* B10 on body weight and food intake of mice

Group	Initial body weight (g)	Final body weight (g)	Average daily gain (g)	Average food intake (g)
HFD	35.47±0.58	41.16±0.52 <sup>a</sup>	0.19±0.03 <sup>a</sup>	4.72±0.30
HFD+B10	35.13±1.20	38.51±1.10 <sup>b</sup>	0.11±0.03 <sup>b</sup>	5.02±0.30

Data are expressed as mean±SD ( $n=15$  for initial body weight, final body weight, and average daily gain;  $n=5$  for average food intake). Different letters indicate a statistically significant difference between groups ( $P < 0.05$ ).

**Table 3** Effects of dietary supplementation of *Bacillus subtilis* B10 on lipid profiles in the liver and serum of mice

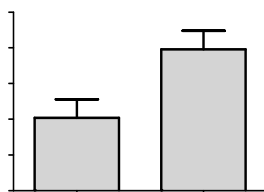
Group	Serum				Liver	
	TG (mmol/L)	TC (mmol/L)	LDL-C (mmol/L)	HDL-C (mmol/L)	TG (mg/g)	TC (mg/g)
HFD	0.96±0.09	3.14±0.12 <sup>b</sup>	0.32±0.08 <sup>b</sup>	1.63±0.11 <sup>b</sup>	20.35±1.38 <sup>a</sup>	6.72±0.63 <sup>a</sup>
HFD+B10	0.94±0.08	4.06±0.17 <sup>a</sup>	0.55±0.11 <sup>a</sup>	2.41±0.58 <sup>a</sup>	13.60±0.60 <sup>b</sup>	4.66±0.17 <sup>b</sup>

Data are expressed as mean±SD ( $n=6$ ). Different letters indicate a statistically significant difference between groups ( $P < 0.05$ ).

**Table 4** Effects of dietary supplementation of *Bacillus subtilis* B10 on serum biochemical parameters of mice

Group	Glucose (mmol/L)	BUN (mmol/L)	GOT (IU/L)	GPT (IU/L)
HFD	8.62±1.00 <sup>a</sup>	5.30±0.39	4.43±0.28 <sup>a</sup>	5.68±0.30 <sup>a</sup>
HFD+B10	5.18±0.54 <sup>b</sup>	5.28±0.22	3.30±0.10 <sup>b</sup>	3.12±0.38 <sup>b</sup>

Data are expressed as mean±SD ( $n=6$ ). Different letters indicate a statistically significant difference between groups ( $P < 0.05$ ).



**Table 5** Effects of dietary supplementation of *Bacillus subtilis* B10 on HFD-induced oxidative stress in the liver of mice

Group	T-AOC (U/mg prot)	TrxR (mg/L)	SOD (U/mg prot)	CAT (U/mg prot)	GSH-Px (U/mg prot)	GSH (mg/g prot)	ASOA (U/g prot)	XO (U/g prot)	8-OHdG (pg/ml)
HFD	0.84±0.04	0.96±0.04	55.32±4.00	0.83±0.06	597.15±31.74 <sup>c</sup>	0.81±0.12	82.18±4.53	16.00±1.26	14.42±1.32 <sup>a</sup>
HFD+B10	1.04±0.17	1.31±0.13	50.53±4.37	1.15±0.13	1441.22±105.98 <sup>a</sup>	0.90±0.08	75.82±4.07	14.45±0.76	10.28±1.02 <sup>b</sup>

Data are expressed as mean±SD (n=6). Different letters indicate a statistically significant difference between groups (P<0.05). prot: protein

### 3.4 Effects of dietary supplementation of *Bacillus subtilis* B10 on hepatic antioxidant biochemical values and DNA damage of mice

As shown in Table 5, the administration of B10 lowered concentrations of hepatic 8-OHdG remarkably (P<0.05), increased T-AOC, GSH, and TrxR levels and the CAT activity slightly, and significantly elevated, by a factor of 1.4, the GSH-Px activity (P<0.05).

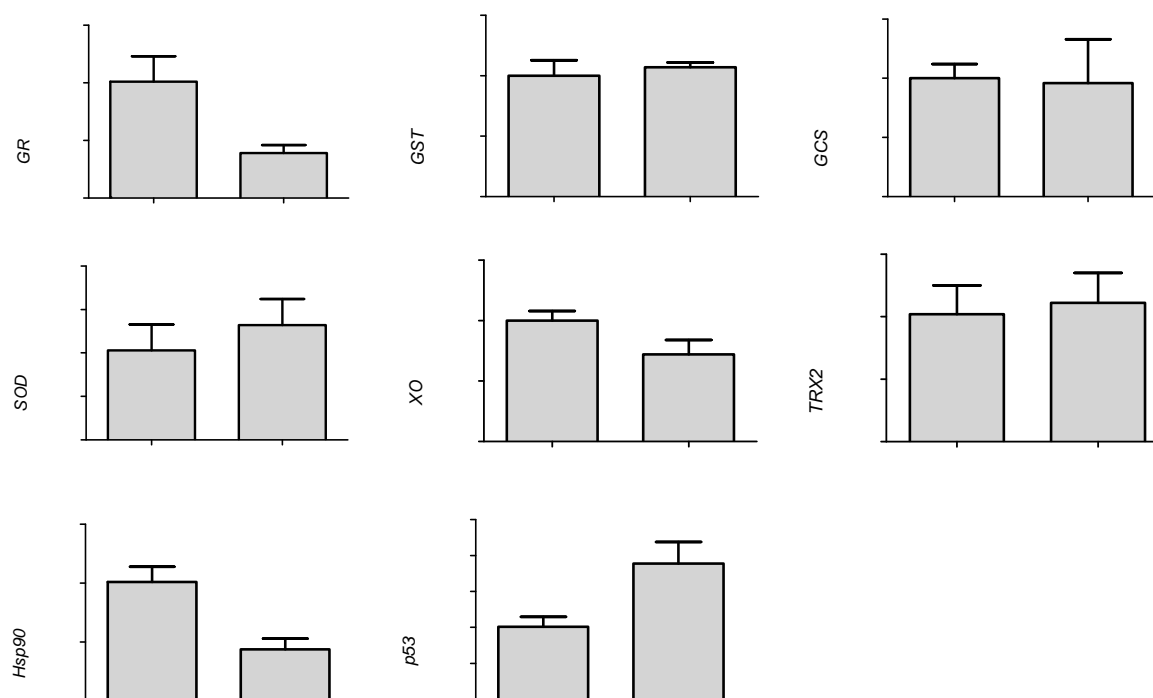
### 3.5 Effects of dietary supplementation of *Bacillus subtilis* B10 on hepatic mRNA expression of genes associated with oxidative stress of mice

It was found that the transcript levels of *GPx1*, *GPx2*, and *GPx3* genes in the liver were much less for the B10 group than for the HFD group (P<0.05). However, an increased hepatic gene expression of *GPx1* was also observed in the B10 treatment group (P<0.05). There was no significant difference between the two groups in the transcript levels of *GPx1*,

glutamylcysteine synthetase (*GPx2*), and thioredoxin reductase 2 (*GPx3*) (Fig. 2).

## 4 Discussion

A 30-d supplementation of B10 is able to decrease mice body weight, although there is no difference in food intake between the two groups (Table 2). The decrease in body weight induced by probiotics is in agreement with the results of other research (Park et al., 2013; Wang et al., 2015). Although we did not compare the digestive capacity and nutrient absorption of mice between the HFD and HFD+B10 groups, previous research showed that the digestive enzyme activity was unchanged between animals fed an HFD or HFD supplemented with probiotics (An et al., 2011). So we conjecture that the lower body weight gain caused by B10 was due to its modulation of lipid metabolism other than because of



**Fig. 2** Effect of dietary supplementation of *Bacillus subtilis* B10 on the expression of mRNA associated with anti-oxidation in mice liver of the HFD (high-fat diet) and HFD+B10 (high-fat diet with 0.1% *Bacillus subtilis* B10) groups

All mRNA quantification data are normalized to the reference gene. Gene expression levels are expressed as values relative to that of the HFD group. Data are expressed as mean±SD (n=6). Different letters indicate a statistically significant difference between groups (p < 0.05)

different food intake or digestion. In recent years, numerous investigations have shown that probiotics can be considered as biotherapeutics in treating abnormal lipid metabolism (Park et al., 2013; Xin et al., 2014). In our study, hepatic TC, TG and serum HDL-C, glucose, activities of GOT and GPT are much less than those of HFD-fed mice (Tables 3 and 4). In fact, GPT is an enzyme related to the liver fat accumulation (Westerbacka et al., 2004). From the point of view of Eckel et al. (2005), once fatty liver disease occurs, widespread lipid will accumulate in muscles and other organs, which facilitates the metabolic syndrome. As GPT could be a sensitive marker of hepatic dysfunction associated with metabolic syndrome, even in the range below the current limit (Jeong et al., 2004), the low GPT activity in the B10 treatment group may imply that B10 could be involved in the alleviation of the susceptibility to metabolic syndrome in HFD-fed mice. Nevertheless, the significant increases of TC and LDL-C in the serum were in contrast to the findings of Panda et al. (2006),

in which serum TC and LDL were reduced by dietary supplementation of B10. However, the study of Novak et al. (2011) also found an enhanced serum LDL-C level. In their opinion, the lowered concentrations of cecal propionic and butyric acids in B10-fed chickens might be the reason for higher levels of LDL compared with the control, because products of bacterial fermentation, specifically short chain fatty acids, may inhibit cholesterol synthesis in the plasma cholesterol in the liver (Rafter, 2002).

Genes of lipid synthesis ( $\gamma$ ,  $\beta$ ), lipolysis ( $\alpha$ ), and energy metabolism ( $\alpha$ ) are involved in lipid metabolism. The obese mice induced by the HFD have association with abnormal lipid metabolism in the liver, including altered acetyl-CoA carboxylase 1 ( $\alpha$ ), fatty acid synthetase ( $\beta$ ), fasting-induced adipose factor ( $\alpha$ ), and  $\gamma$  gene expression (Kang et al., 2013; Xin et al., 2014). In our study, HFD also induced abnormal lipid metabolism compared with

mice fed a normal diet (data not shown), including the up-regulation of lipid synthesis genes and down-regulation of lipolysis. For the HFD+B10 group, the mRNA expression of hepatic  $\alpha$  and is increased while mRNA expression decreases (Fig. 1). Thus, it could be deduced that the positive effect of B10 on lipid reduction may be due to the up-regulation of lipolysis and the down-regulation of lipid synthesis. This is consistent with the study of Kang (2013) and Park (2013), in which the transcript level of  $\alpha$  was also increased and the expression of some lipid synthesis and lipolysis genes was regulated by probiotics.

The abnormal lipid metabolism induced by high fat intake could also affect pro- and anti-oxidative balances in many tissues (Marczuk-Krynicka et al., 2009). The altered lipoprotein profiles enhance the synthesis and accumulation of intramyocellular TG, resulting in an increased synthesis of toxic fatty acid-delivered metabolites, causing enhanced lipid peroxides, hepatic membranes damage, and oxidative stress (Newsholme and Krause, 2014). Gao (2011) demonstrated that HFD-fed ICR mice supplemented with lactic acid bacteria could enhance the activities of SOD and GSH-Px. In the present study, the GSH-Px activity was also significantly up-regulated in HFD+B10 group (Table 5). Moreover, the increased enzyme activity was accompanied by the regulation of genes related to antioxidant enzymes and oxidative stress. XO is an enzyme that catalyzes the oxidation of hypoxanthine to xanthine and could generate reactive oxygen species, leading to oxidative stress (Ardan et al., 2004). On the other hand, the p53, a tumor-suppressor protein, exhibits antioxidant activities to eliminate oxidative stress and ensure cell survival in response to low levels of oxidative stresses through the p53-regulated gene, manganese superoxide dismutase ( ) or other pathways (Sablina et al., 2005; An et al., 2008; Liu and Xu, 2011; Chen et al., 2012; Wu et al., 2014). As we expected, the transcript level of in mice treated with B10 being obviously down-regulated compared with HFD group, expression is markedly up-regulated (Fig. 2). Moreover, as mentioned above, the activity of GPX was significantly increased (Table 5), which means, p53 may exert the antioxidant activity via the increment of GPX activity after B10 treatment. Our findings show that B10 has a

positive regulatory effect on lipid profile, GSH-Px activity and , expression. All these alterations caused by B10 administration result in the attenuation of oxidative stress and DNA damage, which was verified by the decreased expression of antioxidant-related genes ( and ) and lowered 8-OHdG level, a major product of DNA oxidation (de Souza-Pinto et al., 2001). Based on our study, the anti-oxidative activity of B10 is reflected in the increased activity of GSH-Px, the regulation of antioxidant-related genes, like and as well as the lipid profile and may be the modulation of gut microbiota. Accumulating evidences showed a strong correlation between gut microbiota and metabolic syndrome (Araya et al., 2002; Everard et al., 2014) and therefore, further research is needed to clarify the effect of B10 on the modulation of gut microbiota.

## 5 Conclusions

In conclusion, B10 could decrease body weight gain of diet-induced obese mice by improving lipid metabolism and oxidative stress status.

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## Compliance with ethics guidelines

Kai LEI, Ya-li LI, Yang WANG, Jing WEN, Hong-zhao WU, Dong-you YU, and Wei-fen LI declare that they have no conflict of interest.

All institutional and national guidelines for the care and use of laboratory animals were followed.

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## List of electronic supplementary materials

Table S1 Compositions of mice normal diet  
Data S1 Methods for biochemical parameter detection

		<b>B10</b>	
		B10	
		B10	
	B10		$\alpha$
		GSH-Px	
	ICR		
	30		
	8-	8-OHdG	
		PCR	
		B10	
		2	
	3 4		
			1
		B10	2