Maternal dietary betaine supplementation modifies hepatic expression of cholesterol metabolic genes via epigenetic mechanisms in newborn piglets

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Abstract

To elucidate the effects of maternal dietary betaine supplementation on hepatic expression of cholesterol metabolic genes in newborn piglets and the involved epigenetic mechanisms, we fed gestational sows with control or betaine-supplemented diets (3 g/kg) throughout pregnancy. Neonatal piglets born to betaine-supplemented sows had higher serum methionine concentration and hepatic content of betaine, which was associated with significantly up-regulated hepatic expression of glycine N-methyltransferase. Prenatal betaine exposure increased hepatic cholesterol content and modified the hepatic expression of cholesterol metabolic genes in neonatal piglets. Sterol regulatory element-binding protein 2 was down-regulated at both mRNA and protein levels, while 3-hydroxy-3-methylglutaryl-CoA reductase (HMGCR) was down-regulated at the mRNA level, but up-regulated at the protein level, in betaine-exposed piglets. The transcriptional repression of HMGCR was associated with CpG island hypermethylation and higher repressive histone mark H3K27me3 (histone H3 lysine 27 trimethylation) on the promoter, whereas increased HMGCR protein content was associated with significantly decreased expression of miR-497. Furthermore, LDL receptor was significantly down-regulated at both mRNA and protein levels in the liver of betaine-exposed piglets, which was associated with promoter CpG hypermethylation. In addition, the expression of cholesterol-7α-hydroxylase (CYP7α1) was up-regulated at both mRNA and protein levels, while the expression of cholesterol-7α-hydroxylase (CYP7α1) was increased at the mRNA level, but unchanged at the protein level associated with increased expression of miR-181. These results indicate that maternal betaine supplementation increases hepatic cholesterol content in neonatal piglets through epigenetic regulations of cholesterol metabolic genes, which involve alterations in DNA and histone methylation and in the expression of microRNA targeting these genes.

Key words: Betaine; Epigenetic regulation; Cholesterol metabolism; Maternal diet; Piglets

Cholesterol is an essential component of cell membranes and also serves as a precursor for life-sustaining steroid hormones and bile acids. It is well known that deregulation of cholesterol metabolism contributes to obesity, diabetes and CVD. Moreover, cholesterol is particularly essential for embryogenesis, and low plasma cholesterol level is usually correlated with low body weight at birth. Hepatic cholesterol homeostasis is maintained through the coordinated regulation of three relevant processes: biosynthesis; transportation; transformation. In particular, sterol regulatory element-binding protein-2 (SREBP2) and 3-hydroxy-3-methylglutaryl-CoA reductase (HMGCR) are key factors/enzymes for cholesterol biosynthesis, LDL receptor (LDLR) and HDL receptor (scavenger receptor class B type I (SR-BI)) are responsible for transportation, while cholesterol-7α-hydroxylase (CYP7α1) and cholesterol-27α-hydroxylase (CYP27α1) are main enzymes catalysing transformation to bile acids. It has been shown that hepatic cholesterol homeostasis in offspring is highly vulnerable to maternal nutrition status, predominantly through epigenetic mechanisms such as DNA methylation, histone modification and microRNA-mediated post-transcriptional regulation. Methyl donors, such as methionine or folic acid, are able to reverse the epigenetic modifications and thereby restore the behavioural or metabolic disorders in offspring caused by prenatal or neonatal insults.

Betaine, also known as betaine anhydrous or trimethylglycine, functions as a substrate for the formation of methionine, which can then be further converted to S-adenosylmethionine (SAM). In the methionine metabolic cycle, SAM is further converted to S-adenosylhomocysteine by glycine N-methyltransferase (GNMT), thereby donating the methyl group for DNA and protein methylation catalysed by DNMT (DNA methyltransferases) and histone methyltransferases, respectively.
Consequently, betaine supplementation may modulate gene expression through modifying epigenetic marks such as DNA and histone methylation \( ^{(20,21)} \).

Betaine deficiency results in a series of metabolic abnormalities \( ^{(19)} \), while betaine supplementation in the diet can prevent obesogenic diet-induced hepatic steatosis in rodents \( ^{(22)} \) and improve growth performance and carcass characteristics in domestic animals \( ^{(23,24)} \). It has been observed that betaine supplementation could change cholesterol production in human subjects and animals \( ^{(25–26)} \), yet the mechanisms remain unclear. Furthermore, although betaine is known to be critical for embryonic and fetal development \( ^{(19)} \), the effects of betaine supplementation in maternal diet during gestation on hepatic cholesterol metabolism in neonatal offspring and the underlying mechanisms have not been investigated.

In the present study, we aimed to investigate the effects of feeding gestational sows with a betaine-supplemented diet on hepatic cholesterol metabolism in newborn piglets. To explore the potential epigenetic mechanisms underlying such effects, we studied hepatic expression of genes involved in cholesterol and methionine metabolism, and determined the status of DNA and histone methylation on the promoter of cholesterol metabolic genes, as well as the expression of microRNA targeting these genes, in the liver of neonatal piglets.

Materials and methods

Animals and sampling

A total of sixteen second-parity cross-bred sows (Landrace × Yorkshire) at the age of approximately 8 months were artificially inseminated, at the observation of oestrus, with a mixture of Duroc semen samples obtained from two littermate boars. The sows were randomly divided into control and betaine groups (eight per group) after 1 week of the artificial insemination. The sows in the control group were fed a basal diet not supplemented with betaine, while those in the betaine group received a betaine-supplemented (3 g/kg) diet throughout pregnancy. The composition of the two experimental diets is shown in Table 1. Betaine (Skystone Feed Company Limited) was added in the form of betaine hydrochloride with 98% purity. All the sows were fed three times per d at 05.00, 10.00 and 17.00 hours, and had free access to water. Immediately after parturition, newborn piglets were fed a basal diet not supplemented with betaine, while those in the betaine group received a betaine-supplemented (3 g/kg) diet throughout pregnancy. The slaughter and sampling procedures complied with the ‘Guidelines on Ethical Treatment of Experimental Animals’ (2006) No. 398 set by the Ministry of Science and Technology, China.

| Table 1. Composition and nutrient content of the experimental diets |
|---------------------------------|-----------------|-----------------|
| Ingredients (g/kg)             | Control         | Betaine         |
| Maize                          | 370             | 370             |
| Wheat                          | 300             | 300             |
| Bran                           | 80              | 80              |
| Soyabean meal                  | 170             | 170             |
| Lignocelluloses                | 30              | 30              |
| CaHPO₄                          | 20              | 20              |
| Soyabean oil                   | 8               | 8               |
| Premix*                        | 20              | 20              |
| Betaine                        | 0               | 3               |
| Digestible energy (MJ/kg)      | 13.1            | 13.1            |
| Calculated composition (%)     |                 |                 |
| Crude protein                  | 15              | 15              |
| Crude fibre                    | 4.5             | 4.5             |
| Ca                             | 0.84            | 0.84            |
| P                              | 0.65            | 0.65            |

* The premix contained the following vitamins and minerals (per kg of diet): vitamin A, 72 mg; vitamin D₃, 1.5 mg; vitamin E, 648 mg; vitamin K₃, 30 mg; vitamin B₆, 30 mg; vitamin B₁₂, 1.2 mg; vitamin B₉, 60 mg; vitamin B₂, 360 mg; niacin, 600 mg; pantothenic acid, 300 mg; folic acid, 6 mg; manganese sulphate, 1 g; zinc oxide, 2.5 g; ferrous sulphate, 40 g; copper sulphate, 4.0 g; sodium selenite, 6 mg; Ca, 150 g; P, 15 g; NaCl, 40 g.

Determination of hepatic betaine and S-adenosylmethionine content and serum methionine concentration

For the determination of betaine concentrations, 1 g of frozen liver samples was prepared as described previously \( ^{(29)} \). Betaine concentrations in the liver samples were measured with a liquid chromatography (Agilent 1200; Agilent Technologies)–MS (API 5000TM; AB Sciex) system.

Hepatic SAM content was measured using the HCB Quantitative Porcine Competitive ELISA kit (S200FC; Hermes Criterion Biotechnology) following the manufacturer’s instructions. Serum concentration of methionine was determined in duplicate with an automatic amino acid analyser (L-8900; Hitachi).

Measurement of cholesterol and bile acid concentrations

Serum concentration of total cholesterol was measured using a commercial cholesterol assay kit (E1005; Applygen Technologies, Inc.). Serum concentrations of LDL-cholesterol (LDL-C) and HDL-cholesterol were measured with assay kits (006340 and 006328, respectively; Beijing BHKT Clinical Reagent Company Limited). Hepatic total cholesterol concentration was measured using a tissue total cholesterol assay kit (E1015; Applygen Technologies, Inc.). Serum concentrations of LDL-cholesterol (LDL-C) and HDL-cholesterol were measured with assay kits (006340 and 006328, respectively; Beijing BHKT Clinical Reagent Company Limited). Hepatic total cholesterol concentration was measured using a tissue total cholesterol assay kit (E1015; Applygen Technologies, Inc.) following the manufacturer’s instructions. Hepatic total bile acid concentrations were determined by enzymatic colorimetric methods using a commercial kit (E003; Nanjing Jiancheng Bioengineering Institute).

Real-time PCR for mRNA quantification

For RNA extraction, approximately 200 mg of liver samples were homogenised in 1 ml of TRIzol reagent (Invitrogen), according to the manufacturer’s protocol. RNA samples (2 μg) were digested with DNase and reverse transcribed to complementary DNA using random hexamer primers.
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Proteins.

H1 (H1) was selected as the loading control for nuclear the loading controls for total cellular protein, while histone primary antibodies used in Western blot are listed in online protocols provided by the manufacturer. The sources of Western blot analysis of target proteins was carried out according to the protocols provided by the manufacturer. The sources of primary antibodies used in Western blot are listed in online supplementary Table S2. GAPDH or β-actin was selected as the loading controls for total cellular protein, while histone H1 (H1) was selected as the loading control for nuclear proteins.

Protein extraction and Western blot analysis

Total cellular protein and nuclear protein were extracted from 200 mg of frozen liver samples as described previously. Protein concentrations were measured with a Pierce BCA Protein Assay kit (no. 23225; Thermo Scientific). Western blot analysis of target proteins was carried out according to the protocols provided by the manufacturer. The sources of primary antibodies used in Western blot are listed in online supplementary Table S3. GAPDH or β-actin was selected as the loading controls for total cellular protein, while histone H1 (H1) was selected as the loading control for nuclear proteins.

DNA methylation analysis – methylated DNA immunoprecipitation

Genomic DNA extracted from the liver samples was sonicated to produce small fragments ranging from 300 to 1000 bp. The fragmented DNA (2 μg) was heat-denatured to produce single-stranded DNA. A mouse monoclonal antibody against 5-methyl cytidine (ab10805; Abcam) was used to immunoprecipitate the methylated DNA fragments. Pretreated protein G agarose beads (40 μl, 50 % slurry, sc-2003; Santa Cruz Biotechnology) were used to capture the immune complexes. The beads bound to immune complexes were washed to eliminate non-specific binding and resuspended in 250 μl digestion buffer containing proteinase K to purify the immunoprecipitated methylated DNA. A small aliquot of immunoprecipitated methylated DNA was used to amplify the proximal promoter sequences of target genes by real-time PCR. A pair of negative control primers was used to amplify a promoter region absent of CpG sites as the internal control. The results of methylated DNA immunoprecipitation were calculated relative to the internal control, and expressed as the fold change relative to the mean value of the control group. The specific and negative control primers are listed in online supplementary Table S2.

Chromatin immunoprecipitation assay

Approximately 200 mg of frozen liver samples were ground in liquid N2 and resuspended with PBS containing protease inhibitor cocktail (no. 11697498001; Roche). Cross-linking of protein and DNA was performed by adding formaldehyde to a final concentration of 1 %, and then the reaction was terminated with glycine (2.5 mol/l) at room temperature. The reaction mix was centrifuged and the pellets were rinsed with PBS and homogenised in a SDS lysis buffer containing protease inhibitors. Crude chromatin preparations were sonicated to an average length ranging from 200 to 500 bp and precleared with salmon sperm DNA-treated protein G agarose beads (40 μl, 50 % slurry, sc-2003; Santa Cruz Biotechnology). The mixture of precleared chromatin preparations and 2 μg of specific primary antibody were incubated overnight at 4°C (for details of antibodies, see online supplementary Table S2). A negative control was included with normal rat IgG. Protein G agarose beads (40 μl, 50 % slurry, sc-2003; Santa Cruz Biotechnology) were added to capture the immunoprecipitated chromatin complexes. Finally, DNA fragments were released from the immunoprecipitated complexes by reverse cross-linking at 65°C for 1 h, and quantitative real-time PCR was used to quantify the fragments of target gene promoters with specific primers (see online supplementary Table S2) using purified immunoprecipitated DNA as the template.

Quantification of microRNA

Total RNA (2 μg) treated with RNase-free DNase I (Promega) were polyadenylated by poly(A) polymerase using a poly(A) tailing kit (AM1350; Applied Biosystems), according to the manufacturer’s instructions. Polyadenylated RNA was then dissolved and reverse transcribed using a poly(T) adapter. Real-time PCR was performed with SYBR Green qPCR master mix reagent (Takara) in triplicate using a microRNA (miRNA)-specific forward primer and a universal reverse primer complementary to part of the poly(T) adapter sequence. U6 small-nuclear RNA (U6 snRNA) was used as a reference gene to normalise the expression of miRNA. The sequences of all porcine miRNA were acquired from miRBase (http://www.mirbase.org/). miRNA targeting HMGCR, LDLR, CYP7a1 and CYP27a1 were predicted with an online miRNA prediction tool. Among all the predicted miRNA, four miRNA targeting HMGCR, ten targeting LDLR, five targeting CYP7a1 and twelve targeting CYP27a1 were predicted by real-time PCR. The primer sequences used for miRNA analysis are listed in online supplementary Table S4.

Functional validation of microRNA

Genomic sequences of porcine miR-497 and miR-181 precursors (see online supplementary Table S5) were synthesised and inserted into the pSilencer 3.0-H1 small-interfering RNA expression vector by Invitrogen. The 3′-UTR (untranslated regions) of the porcine HMGCR gene containing a conserved motif for miR-497 targeting was amplified by PCR using the primers 5′-GGCCGGGGGGGACACGCTGTCCCTCTA-3′ and 5′-GGAGAGAAGACGCTGGAGG-3′. The 3′-UTR of the CYP7a1 gene containing a conserved motif for miR-181 targeting was amplified using the primers 5′-TGCAAGACGGAGAAATACATTAAACAGTGGCTTA-3′ and 5′-TAACATTAAAATATATATTTA-3′. The PCR products were then cloned downstream to the pGL3-Control luciferase reporter vector (Promega) and the plasmids pGL3-Control/HMGCR and pGL3-Control/CYP7a1, respectively.

Hela cells were cultured in Dulbecco’s modified Eagle’s medium containing 10 % fetal bovine serum at 37°C in
Table 2. Body and liver weights and hepatic total cholesterol and bile acid content, and serum cholesterol concentration in newborn piglets (Mean values with their standard errors, n = 8)

<table>
<thead>
<tr>
<th>Variables</th>
<th>Control group</th>
<th>Betaine group</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean (n = 8)</td>
<td>SEM (n = 8)</td>
</tr>
<tr>
<td>Body weight (kg)</td>
<td>1.65</td>
<td>0.04</td>
</tr>
<tr>
<td>Liver weight (g)</td>
<td>46.08</td>
<td>1.89</td>
</tr>
<tr>
<td>Hepatic TBA (µg/g)</td>
<td>0.94</td>
<td>0.07</td>
</tr>
<tr>
<td>Serum Tch (mmol/l)</td>
<td>1.08</td>
<td>0.11</td>
</tr>
<tr>
<td>Serum LDL-C (mmol/l)</td>
<td>0.50</td>
<td>0.06</td>
</tr>
<tr>
<td>Serum HDL-C (mmol/l)</td>
<td>0.50</td>
<td>0.03</td>
</tr>
<tr>
<td>LDL-C:HDL-C</td>
<td>0.99</td>
<td>0.09</td>
</tr>
</tbody>
</table>

Tch, total cholesterol; TBA, total bile acids; LDL-C, LDL-cholesterol; HDL-C, HDL-cholesterol.

Hepatic betaine and methionine metabolism

Piglets born to betaine-supplemented sows had significantly higher hepatic betaine content (P < 0.05; Fig. 1(a)) and serum methionine concentration (P < 0.05; Fig. 1(b)). Moreover, hepatic SAM content tended to be higher (P = 0.09) in piglets born to betaine-supplemented sows (Fig. 1(c)). These alterations were associated with significantly up-regulated mRNA (P < 0.05; Fig. 1(d)) and protein (P < 0.05; Fig. 1(e) and (f)) expression of GNMT. However, no significant alteration was observed for the mRNA or protein expression of DNMT1, DNMT3a (P = 0.06) or DNMT3b in the liver of newborn piglets born to betaine-supplemented mothers (Fig. 1(d)–(f)).

Results

Reproductive performance of sows and body weight and liver weight of piglets

Maternal betaine supplementation did not affect the litter size or litter weight (see online supplementary Table S1). Moreover, newborn piglets born to betaine-supplemented sows exhibited comparable body weight and liver weight to their control counterparts (Table 2).
Serum cholesterol concentrations and hepatic total cholesterol content

No significant alterations were observed in the serum concentrations of LDL-C- and HDL-cholesterol or in the ratio of LDL-C:HDL-cholesterol. However, hepatic content of total cholesterol was significantly increased (P<0.05) in piglets born to betaine-supplemented sows (Table 2).

Hepatic expression of cholesterol metabolic genes

Among the genes involved in cholesterol de novo synthesis, HMGCR was significantly reduced (P<0.05) by 45% at the mRNA level in piglets born to betaine-supplemented sows (Fig. 2(a)), which was accompanied by a significant down-regulation (P<0.05) of SREBP2 at both mRNA and nuclear protein content levels (Fig. 2(a)–(c)). However, the hepatic protein content of HMGCR was significantly higher (P<0.05) in piglets born to betaine-supplemented sows (Fig. 2(b) and (c)).

Among the genes involved in cholesterol transport and uptake, although maternal betaine supplementation did not alter the serum concentrations of LDL-C or HDL-cholesterol in neonatal offspring (Table 2), hepatic LDLR was significantly down-regulated (P<0.05) at both mRNA (Fig. 2(d)) and protein (Fig. 2(e) and (f)) levels. In contrast, neither the mRNA nor the protein expression of SR-BI was altered in betaine-exposed piglets (Fig. 2(d)–(f)).

Of the genes involved in cholesterol transformation, hepatic CYP7α1 mRNA was significantly up-regulated (P<0.05) by 87% in newborn piglets born to betaine-supplemented sows (Fig. 2(g)), but no alteration was observed at the protein level (Fig. 2(h) and (i)). In contrast, the expression of CYP27α1 was up-regulated (P<0.05), at both mRNA and protein levels, in the liver of piglets born to betaine-supplemented sows (Fig. 2(g)–(i)).

Epigenetic modifications of altered cholesterol metabolic genes

The level of CpG methylation on the HMGCR promoter was 2.34-fold higher (P<0.05) in the liver of piglets born to betaine-supplemented sows (Fig. 3(a)), which was in parallel with a 40% increment (P<0.05) of the repressive
histone mark H3K27me3 (histone H3 lysine 27 trimethylation; Fig. 3(b)). Significant DNA hypermethylation (P < 0.05; Fig. 3(c)) and a trend of more enriched H3K27me3 mark (P = 0.06; Fig. 3(d)) was also detected in the promoter of the \(LDLR\) gene in the liver of piglets born to betaine-supplemented sows. Because no CpG islands were found within the 5'-flanking sequence of the porcine \(CYP7a1\), \(CYP27a1\) or \(SREBP2\) genes, and the \(SR-BI\) mRNA level was not changed in betaine-exposed piglets, we excluded \(CYP7a1\), \(CYP27a1\), \(SREBP2\) and \(SR-BI\) genes from methylated DNA immunoprecipitation and chromatin immunoprecipitation analyses in the present study.

### Hepatic expression of microRNA targeting altered cholesterol metabolic genes

Piglets born to betaine-supplemented sows demonstrated a significant down-regulation (P < 0.05) in the hepatic expression of miR-497 (Fig. 4(a)), which is predicted to target \(HMGCR\). Moreover, \(LDLR\)-targeting miRNA, including miR-1285, miR-138, miR-4334-5p and miR-7144-5p, were all up-regulated (P < 0.05) in the liver of betaine-exposed piglets (Fig. 4(b)). Furthermore, miR-181 predicted to target \(CYP7a1\) was remarkably up-regulated (P < 0.05; Fig. 4(c)), while miR-17-3p, miR-202-3p, miR-30-3p, miR-361-3p and miR-7143-3p predicted to target \(CYP27a1\) were significantly down-regulated (P < 0.05, Fig. 4(d)) in the liver of piglets born to betaine-supplemented sows.

### Functional validation of miR-497 and miR-181

Ectopic expression of miR-497 significantly suppressed (P < 0.05) the luciferase activity of HeLa cells co-transfected with the pGL3-Control/\(HMGCR\) luciferase reporter plasmid at both 24 and 48 h (Fig. 5(a)). Similarly, the luciferase activity of HeLa cells transfected with the pGL3-Control/\(CYP7a1\) reporter plasmid was significantly decreased by co-transfection of pSilencer 3.1 H1-neo miR-181 (P < 0.05) at both 24 and 48 h (Fig. 5(b)).

### Discussion

In the present study, maternal betaine supplementation during gestation caused a significant increase in hepatic betaine content in neonatal offspring. In mammals, betaine is used as a substrate for methionine synthesis (32). Dietary betaine supplementation has been shown to increase serum methionine concentrations in human subjects (33). In agreement with these previous findings, we found elevated serum methionine concentrations associated with higher hepatic betaine content in betaine-exposed piglets. Previous observation indicates that excessive methionine leads to an accumulation of SAM and
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Fig. 4. MicroRNA (miRNA) predicted to target 3′-UTR of 3-hydroxy-3-methylglutaryl CoA reductase (HMGCR) (a), LDL receptor (LDLR) (b), cholesterol-7α-hydroxylase (CYP7a1) (c) and cholesterol-27α-hydroxylase (CYP27α1) (d) in the liver of newborn piglets. Values are means (n 8), with their standard errors represented by vertical bars. * Mean value was significantly different from that of the control group (P<0.05).

The activation of GNMT\(^{(34)}\). Indeed, betaine-exposed piglets demonstrated a higher hepatic SAM content and activated GNMT expression. GNMT is known to provide methyl groups for DNA and protein methylation through the conversion of SAM to S-adenosylhomocysteine\(^{(35)}\). Therefore, maternal betaine supplementation during gestation may modify the methylation of DNA and proteins, especially histones, in the liver of betaine-exposed piglets.

Dietary methionine supplementation has been shown to elevate cholesterol concentrations in the liver and plasma of rats\(^{(36)}\). In the present study, neonates born to betaine-supplemented dams had 37.1% higher hepatic cholesterol content, associated with altered expression of cholesterol metabolic genes involved in cholesterol biosynthesis, transformation and transportation. To our surprise, we found that mRNA abundances of \textit{SREBP2}, \textit{LDLR} and \textit{CYP7a1} were in accordance with their respective protein content, while \textit{HMGCR} and \textit{CYP7a1} demonstrated dissociated mRNA and protein levels. The regulation of gene expression is known to occur at different levels, namely transcription, post-transcription and translation. For some genes, the regulation occurs predominantly at the level of transcription. In this case, mRNA abundance usually agrees with the protein content. However, many other genes are also subjected to post-transcriptional and/or translational regulations, and in which case, mRNA abundance and protein content can be uncoupled. mRNA abundance reported herein represent the steady-state mRNA level resulting from a balance between transcription and degradation. Proteins are considered to be the executors of biological functions. However, many proteins, especially enzymes, have to be modified post-translationally to gain biological activities. In the present study, neither mRNA abundance nor protein content is directly relevant to biological functions. The results presented herein only demonstrate that cholesterol metabolic genes are differently regulated at both transcription and/or post-transcription levels in the liver of betaine-exposed piglets.

DNA methylation and histone modifications are recognised as crucial mechanisms involved in the transcriptional regulation of cholesterol metabolic genes. For instance, maternal dietary protein restriction increased hepatic cholesterol content in rat offspring through modulating histone modifications on the \textit{CYP7a1} promoter and thus altering the mRNA expression of \textit{CYP7a1}\(^{(37)}\). Moreover, dietary protein restriction in sows increased hepatic \textit{HMGCR} expression, which was associated with DNA hypomethylation and lower H3K27me3 on the \textit{HMGCR} gene promoter\(^{(38)}\). In the present study, the promoters of \textit{HMGCR} and \textit{LDLR} genes were found to be hypermethylated in the liver of piglets prenatally exposed to betaine, which coincided with the down-regulation of these two genes at the mRNA level. Furthermore, H3K27me3 is a repressive histone mark that negatively regulates transcription by promoting a compact chromatin structure\(^{(37)}\). The diminished \textit{HMGCR} and \textit{LDLR} gene expression was associated with more enriched H3K27me3 on the respective gene promoters. Therefore, it appears that increased DNA
methylation and repressive histone mark play a role in the transcriptional regulation of HMGCR and LDLR genes in the liver of piglets in response to prenatal betaine exposure.

It is well known that SREBP-2 is a major transcriptional factor that activates the expression of HMGCR and LDLR genes\(^{10,80}\). We found a significant down-regulation of SREBP-2 at both mRNA and protein levels in the liver of the betaine-exposed group, which appears to be the direct cause for repressed transcription of HMGCR and LDLR genes. It should be mentioned that the CpG islands on HMGCR and LDLR gene promoters detected in the present study are predicted to contain binding sites for a number of specific transcriptional factors, thereby regulating the transcriptional level of these genes.

Another interesting finding of the present study is the incongruity between the mRNA and protein levels for HMGCR and CYP7a1 genes. HMGCR was decreased at the mRNA level but remained unchanged at the protein level. The dissociation of mRNA abundance and protein content implies the possible involvement of post-transcriptional regulation. MicroRNA are known to participate predominantly in the post-transcriptional regulation through target mRNA degradation and/or translational repression. Previous studies have indicated that miRNA predominantly promotes the cleavage of mRNA in plants\(^{39,40}\), while miRNA acts mainly through translational repression in animals\(^{41}\). Moreover, it has been reported that in animal models and cell lines, miRNA targets, in principal, protein translation rather than mRNA degradation\(^{42,43}\). Therefore, when transcriptional regulation and miRNA-mediated translational repression are not synchronised, mRNA and protein levels can be uncoupled. In the present study, the down-regulated expression of miR-497 was associated with increased protein content of HMGCR, and the up-regulated expression of miR-181 was associated with the unchanged protein level of CYP7a1, in the liver of betaine-exposed piglets. To verify the targeting sites of miR-497 and miR-181 on 3’-UTR of HMGCR and CYP7a1 transcripts, respectively, we conducted luciferase reporter gene assay and confirmed that miR-497 and miR-181 inhibited luciferase activity through targeting 3’-UTR of HMGCR and CYP7a1, respectively.

Betaine insufficiency is associated with the metabolic syndrome including lipid disorder and diabetes, while betaine supplementation increases serum total cholesterol and LDL-C concentrations\(^{25-26,44}\). Betaine, as a methyl donor, is known to be critical for human embryonic and fetal development\(^{19}\), and plasma betaine concentration has been reported to be correlated with total cholesterol concentrations in an epidemiological study in human subjects\(^{45}\). Owing to their similarities to humans in anatomy, body size, physiology, metabolism and omnivorous habits, pigs are better-suited for human metabolic studies when compared with rodents\(^{46,47}\).

Although mostly descriptive, the results presented herein provide the first evidence that maternal dietary betaine supplementation during gestation causes complex and different changes in the hepatic expression of cholesterol metabolic genes in neonatal piglets, with the involvement of epigenetic modifications including DNA and histone methylation, as well as miRNA expression. These findings may help understand the role of maternal betaine supplementation in fetal programming of cholesterol metabolism in humans.

Obviously, the present study has some limitations. First, epigenetic regulation is only one of the possible mechanisms underlying the altered hepatic expression of cholesterol metabolic genes. The participation of other regulatory mechanisms, such as transcriptional factors and their interaction with epigenetic mechanisms, cannot be ruled out. Second, changes in hepatic cholesterol metabolism in neonates may persist to adulthood, causing long-term consequences in cholesterol homeostasis later in adult life. Follow-up studies are necessary to address the long-term effects on adult health and disease in response to the fetal programming of hepatic cholesterol metabolism caused by maternal betaine supplementation.
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Supplementary material
To view supplementary material for this article, please visit https://dx.doi.org/10.1017/S0007114514002402

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The authors’ contributions were as follows: D. C. performed the experiments and measurements of serum biochemical parameters, analysed and interpreted the results, and drafted the manuscript; Y. J. and J. L. determined the serum hormone and amino acid levels; M. Y., S. S. and H. S. contributed to the experimental design; R. Z. contributed to the experimental concepts and design, provided scientific direction, analysed and interpreted the results, and finalised the manuscript. All authors read and approved the final manuscript.

There are no conflicts of interest to declare.

References


