Maternal protein restriction and fetal growth: lack of evidence of a role for homocysteine in fetal programming

Simon C. Langley-Evans*, Christina Lilley and Sarah McMullen
School of Biosciences, University of Nottingham, Sutton Bonington, Loughborough LE12 5RD, UK
(Received 31 October 2005 – Revised 15 March 2006 – Accepted 27 March 2006)

The disease-programming effects of a maternal low-protein (MLP) diet in rat pregnancy have been suggested to be attributable of hyperhomocysteinaemia. The aim of the present study was to determine whether MLP feeding impacted upon maternal and day 20 fetal homocysteine concentrations, with ensuing effects upon oxidant/antioxidant status. Sixty-four pregnant rats were fed either MLP diet or control diet before termination of pregnancy at days 4, 10, 18 or 20 gestation (full-term gestation 22 d). Maternal plasma homocysteine concentrations were similar in control and MLP-fed dams at all points in gestation. Fetal plasma homocysteine was similarly unaffected by MLP feeding at day 20 gestation. Activities of superoxide dismutase and glutathione peroxidase were similar in livers of mothers and fetuses in the two groups. Whilst catalase activity was not influenced by diet in maternal liver, MLP exposure increased catalase activity in fetal liver at day 20. Oxidative injury (protein carbonyl concentration) was lower in the livers of MLP-fed animals at day 18 gestation (P<0.05), but significantly greater at day 20. Hepatic expression of methionine synthase was similar in control and MLP-fed dams at all stages of gestation. Expression of DNA methyltransferase 1 in fetal liver was altered by maternal diet in a sex- and gestational age-specific manner. In conclusion, MLP feeding does not impact upon maternal or fetal homocysteine concentrations prior to day 20 gestation in the rat. There was no evidence of increased oxidative injury in fetal tissue that might explain the long-term programming effects of the diet.

Programming: Pregnancy: Homocysteine: Antioxidants

Epidemiological associations between risk of heart disease and diabetes in adult life and indices of fetal growth have prompted the suggestion that early life events programme physiological and metabolic processes (Gluckman & Hanson, 2004). Experimental studies of rodents using a wide variety of nutritional manipulations in pregnancy confirm that nutritional programming plays an important role in the aetiology of disease (Langley-Evans et al. 1996b; Gambling et al. 2003; Khan et al. 2003). One such approach has been the feeding of a maternal low-protein (MLP) diet in rat pregnancy. Offspring of MLP-fed rat dams exhibit hypertension (Langley & Jackson, 1994), disturbed glucose handling (Langley et al. 1994), impaired renal development (Langley-Evans et al. 1999) and have a shorter lifespan (Sayer et al. 2001).

Studies with the MLP diet have generally used casein as the sole protein source, and as such have required additional dietary methionine to provide requirements for sulphur. Rees (2002) suggests that the level of methionine provided exceeds the capacity of the maternal liver to convert it to cysteine via the trans-sulphuration pathway. This leads to accumulation of toxic homocysteine which will have the capacity to cause oxidative injury unless recycled to methionine via methionine synthase or homocysteine-betaine methyltransferase activities (Young et al. 2004). Disturbances of the methionine–homocysteine cycle may impact upon the provision of methyl donors for methylation of DNA, which is proposed as a mechanism for heritable programming of gene function (Waterland & Jirtle, 2003; McMullen & Robinson, 2005).

This view is supported by evidence from a number of studies. Studies in which the MLP diet is supplemented with folic acid (Lillicrap et al. 2005) or with glycine (Jackson et al. 2003) generally show that the effects of low-protein feeding can be reversed. Lillicrap et al. (2005) have shown that MLP feeding impacted upon both the expression and methylation status of specific genes. However, in contrast, Bogdarina et al. (2004) found no evidence that maternal protein restriction impacted on the methylation of the promoter for hepatic glucokinase, despite programmed changes in gene expression. The case for a gross imbalance in methionine–homocysteine metabolism, whilst supported indirectly by the observed effects of folate or glycine supplementation, is yet to be confirmed. Indeed Brawley et al. (2004) reported no differences in maternal homocysteine concentrations in low-protein-fed pregnant rats compared to controls.

The present study aims to assess the potential contribution of elevated plasma homocysteine to programming of physiological and metabolic processes by focusing primarily on the fetal period. This avoids the potential contribution of secondary mechanisms that may act to influence development in the postnatal period. In addition to considering the impact of MLP feeding on maternal homocysteine, the study assesses the possibility that the pro-oxidant effects of homocysteine

Abbreviations: DNMT1, DNA methyltransferase 1; GPx, glutathione peroxidase; MLP, maternal low protein; SOD, superoxide dismutase.

*Corresponding author: Dr Simon C. Langley-Evans, fax +44 (0)115 951 6122, email Simon.Langley-Evans@Nottingham.ac.uk
impact upon oxidant–antioxidant balance within both maternal and fetal tissue. Our hypothesis was that MLP feeding would elevate homocysteine, increase oxidative injury in both the maternal and fetal compartment, and impact upon expression of DNA methyltransferase 1 (DNMT1), which is one of the enzymes responsible for the maintenance of DNA methylation patterns during development (Newell-Price et al. 2000). DNMT1 was selected for study as the other DNA methyltransferases (3L, 3a and 3b) are expressed only in the embryonic period and are responsible for de novo methylation. Their role in the maintenance of imprinted methylation patterns is of little significance.

Materials and methods

Chemicals and reagents

All chemicals and reagents were of reagent grade and purchased from Fisher Scientific (Loughborough, UK).

Animals

Experiments described in the present paper were performed under licence from the Home Office in accordance with the 1986 Animals Act. Rats were housed in plastic boxes on a 12 h light cycle, at a temperature of 20 ± 2°C. The rats had free access to food and water at all times. Sixty-four virgin female Wistar rats (Harlan Ltd, Belton, UK) were mated at weights between 180 and 220 g. Upon confirmation of mating by the appearance of a semen plug on the cage floor, the rats were allocated to be fed either a control diet (180 g casein/kg diet, n 32) or a low-protein diet (MLP; 90 g casein/kg diet, n 32), as described previously (Langley-Evans et al. 1994). The full composition of the diets is described elsewhere (Langley-Evans et al. 1994), but essentially each diet was based on casein mixed with starch–sucrose (2:1, w/w), cellulose, maize oil, and minerals and vitamins formulated to AIN76 specification. The diets were isonenergetic, the difference in energy between the control and MLP diets being made up with additional carbohydrate (starch–sucrose 2:1, w/w). The rats were fed the semi-synthetic diets until pregnancy was terminated at days 4, 10, 18 or 20 gestation. At these time-points the pregnant rats were killed using CO2 asphyxia and cervical dislocation. Maternal blood was collected by cardiac puncture and transferred to heparin tubes. Blood was centrifuged at 13 000 rpm and 4°C for 10 min and the plasma was retained. The maternal liver was rapidly dissected and snap-frozen in liquid nitrogen. Liver and plasma were stored at −80°C for up to 6 months until used for further analyses. Six of the MLP-fed and two control dams failed to carry their pregnancy to the time of cull.

At days 18 and 20 gestation, the uterus was removed and each horn was opened. Fetuses were removed and separated from the placentas and, following weighing, were killed by decapitation for collection of blood samples. Fetal blood samples were pooled from all the animals in the litter. The placentas were trimmed, blotted and weighed. Each fetus had its sex determined by genito-anal distance, and the position in the horn was noted along with the sex of the neighbouring fetuses. The number of resorption sites in the uterus was also noted. The liver, heart and kidneys were dissected from each fetus, weighed to the nearest 0·1 mg and then snap-frozen in liquid nitrogen. Fetal blood was centrifuged at 13 000 rpm and 4°C for 10 min and the plasma was retained. Organs and plasma were stored at −80°C until used for further analyses.

Plasma homocysteine determination

Homocysteine concentrations were measured in day 20 fetal and in all maternal plasma samples using a commercial kit (Diazyme, La Jolla, CA, USA). Insufficient sample was obtained for assay of fetal homocysteine at day 18 gestation. For sample extraction, 20 μl plasma sample or calibrator (2–60 μmol/l S-adenosyl-L-homocysteine) were added to eppendorf tubes containing 150 μl of a mixture of reagents (130 μl adenosine, Tris (2-carboxyethyl)-phosphine hydrochloride, 10 μl Tris buffer, 50 mM-phosphate buffer and 10 μl recombinant S-adenosyl-L-homocysteine hydrolase, phosphate buffer and glycerol) prepared according to the manufacturer’s instructions. After thorough mixing and incubation at 37°C for 30 min, 100 μl adenosine analogue in phosphate buffer were then added to each tube. Tubes were mixed well and incubated at room temperature (18–25°C) for 10 min. Adenosine deaminase (50 μl) was added and tubes were left to incubate at room temperature for a further 15 min. Finally, 100 μl DEAE-Sephadex was added to the tubes, the solution was mixed and incubated at room temperature for 10 min.

The pretreated samples or calibrators (25 μl) were loaded in duplicate on to avidin-coated microtitre plate strips. Diazyme™-12A-biotin conjugate (50 μl) was then added to each microtitre well and the plate was left to incubate for 5 min at room temperature. Horseradish peroxidase–S-adenosyl-L-homocysteine (25 μl) was added to each well and was left to incubate for a further 30 min in the dark, at room temperature. The plate was then decanted and blotted on paper towel. The wells were washed three times with 400 μl wash per well. Tetramethylbenzidine solution (100 μl) was added to each well and left to incubate at room temperature for 10 min. Finally, 100 μl 1 M-phosphoric acid (stop solution) were added to the wells. The plate was then read at 450 nm using a plate reader (Tecan Sunrise, Männedorf, Switzerland) within 15 min of adding the stop solution. Calibration curves were prepared using a four parameter logistic curve fit. The intra-assay variation for this assay was 7.74 % and all samples fell within the dynamic range of the assay.

Determination of mRNA expression

Total RNA was isolated from snap-frozen livers using the TRIzol method (Invitrogen, Paisley, UK). The RNA was treated with DNase (Promega, UK) and subjected to phenol–chloroform extraction and ethanol precipitation. Total RNA (0.5 μg) was reverse-transcribed using MMLV RT (Promega, Southampton, UK). Real-time RT–PCR was performed using an ABI Prism 7700 sequence detection system (Applied Biosystems, Foster City, CA, USA). A template-specific primer pair and an oligonucleotide probe (Sigma-Genosys, Poole, UK) specific to each of methionine synthase, DNMT1 and the housekeeping gene β-actin were designed using Primer Express version 1.5 (Applied Biosystems). The full sequences of the primers and probes are shown in
Table 1. Primer sequences

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward primer</th>
<th>Reverse primer</th>
<th>Probe</th>
</tr>
</thead>
<tbody>
<tr>
<td>Methionine synthase, NM_030864 DNA</td>
<td>gatgcgccaggctgacatgt</td>
<td>caagctcgcagcatttg</td>
<td>catccggtgagcaagtgtctgagg</td>
</tr>
<tr>
<td>methylenetetrahydrofolate reductase, NM_00105006 β-Actin, NM_031144</td>
<td>ttcacccacgaccagt</td>
<td>tggtaagcagcaggatacata</td>
<td>tggtagcagcaggctgtg tgcct</td>
</tr>
</tbody>
</table>

TM, melting temperature (°C); GC, percentage of G and C bases.

Table 1. All primer sets were tested under the Taqman PCR conditions using rat genomic DNA as a template. In all cases a single product of the appropriate size was detected by gel electrophoresis (data not shown). A negative template control and relative standard curve were included on every PCR run. The standard curve was prepared from a pool of sample cDNA at relative dilutions of 0.05, 0.1, 0.2, 0.4, 1.0, 2.5 and 5.0. Relative target quantity was calculated from the standard curve and all samples were normalised to β-actin expression.

Antioxidant enzyme activities

The activities of the antioxidant enzymes catalase, glutathione peroxidise (GPx) and superoxide dismutase (SOD) were determined in liver samples. In the present study, SOD activity represents total SOD within the tissue with no separation of the CuZn and Mn forms. Activity of GPx was determined using the spectrophotometric method of Aebi (1984), in which the conversion of H2O2 to H2O and O2 is monitored as a change in absorbance at 240 nm. SOD activity was determined using the method of Levine et al. (1976). As shown in Table 2, the weight gain of the pregnant animals was unaffected by the feeding of the MLP diet. Litter size (not assessed at days 4 or 10) was not affected by maternal diet at day 18 gestation. Litter size was significantly lower in the MLP group than in the control group at day 20 gestation. Maternal plasma homocysteine concentration tended to decline with advancing gestation (Table 2), but this effect was not statistically significant (P=0.118). There was no statistically significant effect of feeding the MLP diet on maternal homocysteine. Fetal plasma homocysteine concentrations (Table 2) at day 20 were similar to those noted in maternal plasma, but no correlation between maternal and fetal concentrations was observed. Methionine synthase is one of the key enzymes catalysing the regeneration of methionine from homocysteine. No difference in the expression of hepatic methionine synthase was noted between control and MLP-fed dams at any stage of gestation (Fig. 1). Methionine synthase expression was normalised against expression of β-actin, which did not vary with gestation or maternal diet.

Statistical analysis

All data are presented as means and their standard errors. Following assessment of the normality of distributions for all variables using a Kolmogorov–Smirnov test, all data were analysed using two- or three-way ANOVA as appropriate, followed by a least significance difference test as a post hoc test. Fetal and placental weight data were adjusted for litter size, the position in the uterine horn and the sex of the neighbouring fetuses in the horn, as these are factors known to impact on fetal growth in the rat. P<0.005 was accepted as statistically significant.

Results

As shown in Table 2, the weight gain of the pregnant animals was unaffected by the feeding of the MLP diet. Litter size (not assessed at days 4 or 10) was not affected by maternal diet at day 18 gestation. Litter size was significantly lower in the MLP group than in the control group at day 20 gestation. Maternal plasma homocysteine concentration tended to decline with advancing gestation (Table 2), but this effect was not statistically significant (P=0.118). There was no statistically significant effect of feeding the MLP diet on maternal homocysteine. Fetal plasma homocysteine concentrations (Table 2) at day 20 were similar to those noted in maternal plasma, but no correlation between maternal and fetal concentrations was observed. Methionine synthase is one of the key enzymes catalysing the regeneration of methionine from homocysteine. No difference in the expression of hepatic methionine synthase was noted between control and MLP-fed dams at any stage of gestation (Fig. 1). Methionine synthase expression was normalised against expression of β-actin, which did not vary with gestation or maternal diet.
The activities of the major antioxidant enzymes in the maternal liver varied considerably with increasing gestation (Table 3). GPx activity increased 6-fold between days 10 and 18 gestation and remained elevated at day 20. ANOVA indicated that this activity increased to a lesser extent in MLP-fed dams than in the control group at day 18 gestation (interaction of gestation and maternal diet, $P=0.011$). Total SOD activity also increased with gestation. A similar pattern was noted for catalase activity. Oxidative damage, as indicated by the presence of protein carbonyls, was observed in all samples and increased with gestation (Table 3). At day 18 MLP-fed rats had significantly lower protein carbonyl concentrations, but by day 20 MLP feeding was associated with significantly more oxidative damage in liver.

As shown in Table 4, the maternal diet had no significant impact upon fetal weight at either day 18 or day 20, after adjustment for litter size, position in horn and the sex of neighbouring fetuses. Placental weight, however, was significantly lower in the MLP-fed animals than controls at day 18 gestation and at day 20 gestation (female fetuses only). Fetal/placental ratio increased significantly between days 18 and 20 gestation. This increase was less marked in female fetuses, but not affected by the maternal diet. Liver and heart weight relative to body weight was constant throughout gestation and similar in control and MLP-treated pregnancies (Table 4). Kidney weight relative to body weight increased between days 18 and 20, with the same magnitude of increase noted in both groups of animals.

DNMT1 is the enzyme responsible for the maintenance of DNA methylation patterns during replication of DNA and hence cell division (Newell-Price et al. 2000). The expression of DNMT1 mRNA was determined in the livers of male and female fetuses at days 18 and 20 gestation (Fig. 2). There was no significant effect of diet, sex or gestation upon expression of this gene, although there was an interactive effect of all three factors ($P=0.027$). This suggested that the maternal diet influenced expression of this gene in a sex- and gestational age-specific manner. Diet effects on DNMT1 expression were confined to day 18 gestation, and here the...
Table 3. Maternal liver antioxidant enzyme activities and protein carbonyl concentrations of rats fed either maternal low-protein (MLP) diet or control diet‡

(Mean values with their standard errors)

| Days gestation | Control (n 8) | MLP (n 6) | Control (n 8) | MLP (n 8) | Control (n 7)§ | MLP (n 7) | Control (n 7) || MLP (n 5)||
|----------------|-------------|----------|-------------|----------|-------------|----------|------------|-------------|
|                | Mean   | SEM     | Mean   | SEM     | Mean   | SEM     | Mean   | SEM     | Mean   | SEM     | Mean   | SEM     |
| GPx (units/mg protein) | 0.259  | 0.015   | 0.221  | 0.009   | 0.276  | 0.092   | 0.242*  | 0.078   | 1.800†  | 0.146   | 1.493†  | 0.060   |
| SOD (units/mg protein)** | 0.681  | 0.063   | 0.604  | 0.063   | 0.661  | 0.039   | 0.608  | 0.039   | 0.893†  | 0.056   | 0.788  | 0.040   |
| Catalase (units/mg protein)†† | 223.7  | 19.7    | 171.3  | 14.1    | 172.7  | 20.2    | 179.1  | 9.2     | 441.5†  | 71.6    | 312.0†  | 13.9    |
| Carbonyls (nmol/mg protein)‡‡ | 0.134  | 0.020   | 0.142  | 0.010   | 0.139  | 0.020   | 0.124  | 0.020   | 0.352†  | 0.040   | 0.250†*  | 0.015   |

GPx, glutathione peroxidase; SOD, superoxide dismutase.
Mean values were significantly different from those of the control group: *P < 0.05.
Mean values were significantly different from those of day 4 gestation within the same maternal dietary group: †P < 0.05.
‡ For details of procedures, see pp. 579–580.
§ n 6 for GPx and SOD.
‖ n 4 for GPx and SOD.
† Activity of GPx was influenced by gestation (P < 0.001) and the interaction of diet and gestation (P < 0.011).
** Total SOD activity was influenced by gestation (P < 0.001).
†† Catalase activity was influenced by gestation (P < 0.001) and the interaction of diet and gestation (P < 0.001).
‡‡ Protein carbonyl concentration was influenced by gestation (P < 0.001) and the interaction of diet and gestation (P < 0.001).
Gestation (d) Maternal diet Sex

Table 4. Fetal and placental weight and fetal organ weight of rats fed either maternal low-protein (MLP) diet or control diet§

<table>
<thead>
<tr>
<th>Gestation (d)</th>
<th>Maternal diet</th>
<th>Sex</th>
<th>n</th>
<th>Fetal wt (g)</th>
<th>Placental wt (g)</th>
<th>Fetal/placental ratio*</th>
<th>Liver (% body wt)</th>
<th>Heart (% body wt)</th>
<th>Kidney (% body wt)</th>
</tr>
</thead>
<tbody>
<tr>
<td>18</td>
<td>Control</td>
<td>Male</td>
<td>49</td>
<td>1.48</td>
<td>0.04</td>
<td>0.50</td>
<td>0.01</td>
<td>0.50</td>
<td>0.01</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Female</td>
<td>52</td>
<td>1.47</td>
<td>0.04</td>
<td>0.49</td>
<td>0.01</td>
<td>0.50</td>
<td>0.01</td>
</tr>
<tr>
<td></td>
<td>MLP</td>
<td>Male</td>
<td>52</td>
<td>1.44</td>
<td>0.04</td>
<td>0.46</td>
<td>0.01</td>
<td>3.25</td>
<td>0.10</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Female</td>
<td>58</td>
<td>1.44</td>
<td>0.03</td>
<td>0.45</td>
<td>0.01</td>
<td>3.24</td>
<td>0.08</td>
</tr>
<tr>
<td>20</td>
<td>Control</td>
<td>Male</td>
<td>60</td>
<td>3.60†</td>
<td>0.08</td>
<td>0.60†</td>
<td>0.02</td>
<td>6.23†</td>
<td>0.20</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Female</td>
<td>50</td>
<td>3.60†</td>
<td>0.09</td>
<td>0.65†</td>
<td>0.02</td>
<td>5.86†</td>
<td>0.24</td>
</tr>
<tr>
<td></td>
<td>MLP</td>
<td>Male</td>
<td>38</td>
<td>3.80†</td>
<td>0.12</td>
<td>0.61†</td>
<td>0.01</td>
<td>6.36†</td>
<td>0.24</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Female</td>
<td>33</td>
<td>3.55†</td>
<td>0.12</td>
<td>0.61†</td>
<td>0.01</td>
<td>5.87†</td>
<td>0.21</td>
</tr>
</tbody>
</table>

Mean values were significantly different from those of the control group at same age: *P<0.05; 0.001.
Mean values were significantly different from those of day 18 gestation within the same maternal dietary group: †P<0.05.
Mean values were significantly different between males and females of same age and dietary group: ‡P<0.05.
Mean values were significantly different from those of day 18 gestation within the same maternal dietary group: †P<0.05.
§ For details of procedures, see pp. 579–580. Fetal and placental weight data are adjusted for litter size and position in the uterine horn and analysed using litter of origin as a covariate.

Fig. 2. Fetal DNA methyltransferase 1 (DNMT1) mRNA expression. Data are presented as the ratio of DNMT1 mRNA expression to β-actin mRNA expression (arbitrary units) for males (A) and females (B). For details of procedures, see pp. 579–580. Values are means with their standard errors depicted by vertical bars (four to eight observations per group). ANOVA indicated an interaction of gestational age, fetal sex and maternal diet (P=0.027). Mean value was significantly different from that of the control group: *P<0.05. (A) Control; (B) maternal low-protein diet.

Discussion

The feeding of the MLP diet to pregnant rats is an established protocol for the induction of hypertension and renal deficits in later life (Langley-Evans, 2001). As a well-characterised system for studying nutritional programming, it is important to be able to define the mechanism or mechanisms through which MLP feeding exerts its long-term effects. Rees (2002) has proposed that the methionine content of the MLP diet is in excess of requirements, producing an accumulation of homocysteine. This may be envisaged as having toxic actions within maternal and fetal tissues by virtue of its oxidant properties, but may also impact on DNA methylation, since S-adenosylhomocysteine inhibits DNMT1 (Chiang, 1998). One of the most important findings of the present study was the observation that neither maternal nor fetal plasma homocysteine concentrations differed in MLP pregnancies compared to controls. This is in contrast to the findings of Petrie et al. (2002) who reported that low-protein feeding raised maternal homocysteine at day 4 but not day 17 gestation. Consistent with the present study, Brawley et al. (2004) found no difference in homocysteine at day 18 gestation in rats fed MLP diet. The data do not therefore support the hypothesis that MLP feeding promotes gross disturbance in the methionine–homocysteine cycle.

Mean values were significantly different from those of the control group at same age: *P<0.05; 0.001.
Mean values were significantly different from those of day 18 gestation within the same maternal dietary group: †P<0.05.
Mean values were significantly different between males and females of same age and dietary group: ‡P<0.05.
Mean values were significantly different from those of day 18 gestation within the same maternal dietary group: †P<0.05.
§ For details of procedures, see pp. 579–580. Fetal and placental weight data are adjusted for litter size and position in the uterine horn and analysed using litter of origin as a covariate.

Trends for effects on males and females appeared to go in opposite directions (P=0.002 for diet × sex interaction at day 18). At day 18, in females, DNMT1 mRNA expression was elevated by MLP exposure (P<0.005), whilst in males MLP tended to lower expression, although this was not statistically significant (P=0.085). Expression of DNMT1 mRNA was normalised against β-actin mRNA expression, which did not vary between groups.

GPx and SOD activities in the livers of fetuses at days 18 and 20 showed no significant effects of maternal diet (Table 5). Catalase activity increased by 166–317% between days 18 and 20, but GPx and total SOD activities were similar at both time-points. At day 20, MLP-exposed fetuses had significantly greater hepatic catalase activity than controls. No differences in protein carbonyl concentration were noted between the two dietary groups, but oxidative injury was greater at day 20 than at day 18.

Discussion

The feeding of the MLP diet to pregnant rats is an established protocol for the induction of hypertension and renal deficits in later life (Langley-Evans, 2001). As a well-characterised system for studying nutritional programming, it is important to be able to define the mechanism or mechanisms through which MLP feeding exerts its long-term effects. Rees (2002) has proposed that the methionine content of the MLP diet is in excess of requirements, producing an accumulation of homocysteine. This may be envisaged as having toxic actions within maternal and fetal tissues by virtue of its oxidant properties, but may also impact on DNA methylation, since S-adenosylhomocysteine inhibits DNMT1 (Chiang, 1998). One of the most important findings of the present study was the observation that neither maternal nor fetal plasma homocysteine concentrations differed in MLP pregnancies compared to controls. This is in contrast to the findings of Petrie et al. (2002) who reported that low-protein feeding raised maternal homocysteine at day 4 but not day 17 gestation. Consistent with the present study, Brawley et al. (2004) found no difference in homocysteine at day 18 gestation in rats fed MLP diet. The data do not therefore support the hypothesis that MLP feeding promotes gross disturbance in the methionine–homocysteine cycle.
Clearly differences in the finding exist between studies, and this may best be explained by variations in protocol. The present study and those of Petrie et al. (2002) and Brawley et al. (2004) all used identical diets. Whilst we and Brawley et al. (2004) introduced the synthetic diets on day 0 of pregnancy, Petrie et al. (2002) used a habituation protocol whereby dams were fed the diets for 2 weeks prior to mating. The use of a habituation period is known to produce variation in the fetal growth patterns associated with low-protein feeding (Langley-Evans & Nwagwu, 1998), so may also elicit metabolic adaptations in the dams that explain the inconsistent evidence that folate (Lillycrop et al. 2005) or glycine (Jackson et al. 2003; Brawley et al. 2004) supplementation of the MLP diet overcomes some of the long-term programming effects. This may stem from direct effects of the diet upon the folate cycle and availability of serine and glycine. With apparent deficits of folate, homocysteine remethylation would be mostly dependent upon the betaine pathway, which would therefore provide the only mechanism for the normalisation of plasma homocysteine concentrations observed in the present study. The alternative to this is that the original premise that the methionine–protein ration is imbalanced (Rees, 2002) is incorrect and the key link between MLP feeding and DNA methylation lies in a different pathway.

DNMT1 is one of a family of DNA methyltransferases and is important during development as it maintains the DNA methylation pattern when DNA replicates during cell division. DNMT1 is essential for normal development and mice deficient for this gene die in utero (Li et al. 1992). Reduced DNA methylation leading to increased expression of specific genes certainly appears to be one mechanism through which MLP feeding programmes physiological processes (Lillycrop et al. 2005). The present study suggests that variation in DNA methylation could arise due to diet-induced changes in the expression of DNMT1. Interestingly, these changes were specific to the day 18 time-point, although could, of course, have been operating at earlier stages of development. Moreover, the effects of MLP exposure on DNMT expression were sex-specific. In females the significant

### Table 5. Fetal liver antioxidant enzyme activities and protein carbonyl concentrations of rats fed either maternal low-protein (MLP) diet or control diet§

<table>
<thead>
<tr>
<th></th>
<th>Days gestation</th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>18</td>
<td>20</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Control (n 14)</td>
<td>MLP (n 14)</td>
<td>Control (n 14)</td>
<td>MLP (n 10)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Mean</td>
<td>SEM</td>
<td>Mean</td>
<td>SEM</td>
<td>Mean</td>
</tr>
<tr>
<td>GPx (units/mg protein)</td>
<td>0.096</td>
<td>0.010</td>
<td>0.103</td>
<td>0.087</td>
<td>0.171</td>
</tr>
<tr>
<td>SOD (units/mg protein)</td>
<td>0.97</td>
<td>0.10</td>
<td>0.86</td>
<td>0.10</td>
<td>1.53</td>
</tr>
<tr>
<td>Catalase (units/mg protein)</td>
<td>41.57</td>
<td>6.44</td>
<td>38.07</td>
<td>2.78</td>
<td>110.73†</td>
</tr>
<tr>
<td>Carboxyls (nmol/mg protein)</td>
<td>0.197</td>
<td>0.026</td>
<td>0.220</td>
<td>0.023</td>
<td>0.372†</td>
</tr>
</tbody>
</table>

GPx, glutathione peroxidase; SOD, superoxide dismutase.

Mean value was significantly different from that of the control group: *P* < 0.05.

Mean values were significantly different from those of day 18 gestation within the same maternal dietary group: †*P* < 0.05.

‡ For details of procedures, see pp. 579–580. Fetal sex had no impact on antioxidant activities or protein carbonyl concentrations and so data are for males and females combined.

§ GPx activity rose significantly with gestation (*P* < 0.001).

† SOD activity rose significantly with gestation (*P* < 0.001).

** Catalase activity rose significantly with gestation (*P* < 0.001) and varied with the interaction of diet and gestation (*P* < 0.020).

†† Protein carbonyl concentration was influenced by gestation (*P* < 0.001).
up-regulation of DNMT1 may actually favour increased methylation, rather than the hypomethylation observed by Lillycrop et al. (2005). The trend for decreased DNMT1 expression in males is, however, consistent with reduced DNA methylation. These effects will require further investigation to explore their contribution to sex-specific programming, which has been observed in a number of systems (McMullen & Langley-Evans, 2005). Lillycrop et al. (2005) made their observations on a mixed population of male and female offspring and so it is unclear whether there were sex differences in DNA methylation patterns. The other DNA methyltransferases (3L, 3a and 3b) were not studied in the present experiment. These enzymes, which are only expressed in the embryo, are responsible for de novo DNA methylation (Newell-Price et al. 2000) and may therefore be likely targets for further study in the context of nutritional programming.

The original hypothesis for the present study was that homocysteine may contribute to programming via its oxidative properties. Low-protein feeding is known to lower antioxidant defences, particularly through depletion of tissue glutathione (Huang & Fwu, 1992). The combination of raised homocysteine and lower antioxidant defence could be expected to promote oxidative injury, apoptosis and hence tissue remodelling in the developing fetus. There were no consistent differences in protein carbonyl concentrations in either maternal or fetal tissues. This may in part be explained by the fact that MLP feeding increased fetal activity of catalase, which shares similar substrates with GPx and may therefore compensate for any deficit in glutathione cycle activities. Oxidative injury did vary with diet in maternal liver, but this did not show a consistent pattern with increasing gestation, or appear to be related to homocysteine concentration. Whilst fetal liver showed no evidence of oxidant/antioxidant imbalance associated with MLP feeding up to day 20 gestation, we have previously noted increased oxidation of hepatic proteins at birth (Langley-Evans & Sculley, 2005). This suggests that oxidative processes mediate some programming effects of MLP feeding over the last 2 d of gestation, a period of very rapid fetal growth and development in the rat. There is no evidence in the present study that this is attributable to the oxidant effects of homocysteine and other mechanisms must therefore account for this process.

Fetal and placental weights were evaluated at two points in late gestation. After accounting for important factors that impact on rates of fetal growth in a litter-bearing species, including transfer of steroid hormones between adjacent male and female fetuses in the uterine horn, we found no significant effects of maternal protein restriction upon fetal weight, the fetal/placental ratio or the size of the major truncal organs. We have previously reported that MLP feeding impacts principally on fetal growth in mid-late gestation, with accelerated growth up to day 20 and slowing of truncal growth over the last 2 d of gestation (Langley-Evans et al. 1996a). These effects appear to depend upon the timing of introduction of MLP feeding and the plane of nutrition prior to pregnancy (Langley-Evans & Nwagwu, 1998). Differences between the current findings and those of earlier studies are most likely explained by the fact the present study has more carefully considered effects of gender and position in the uterine horn. Ultimately the offspring of MLP-fed dams are of normal weight at birth in most studies. The present data therefore reinforce the view that nutritional programming occurs independently of fetal growth retardation. This has important implications for the interpretation of epidemiological data linking lower weight at birth, or disproportion at birth to disease in later life.

In conclusion, the present study has discounted maternal hyperhomocysteinaemia as a potential basis for altered patterns of DNA methylation in rat offspring exposed to a low-protein diet in utero. MLP feeding did not significantly alter plasma homocysteine concentrations in mothers or fetuses and, similarly, did not produce any changes in maternal hepatic methionine synthase activity. Although MLP feeding compromises antioxidant defences in maternal and fetal tissues, there was no evidence that this process was related to homocysteine. There was a strong suggestion that fetal liver expression of DNMT1 was perturbed by MLP feeding, in a gestation- and sex-specific manner and this will require further investigation to assess whether this provides a route to altered DNA methylation.

References


