NMR-based metabonomic studies reveal changes in the biochemical profile of plasma and urine from pigs fed high-fibre rye bread

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This study presents an NMR-based metabonomic approach to elucidate the overall endogenous biochemical effects of a wholegrain diet. Two diets with similar levels of dietary fibre and macronutrients, but with contrasting levels of wholegrain ingredients, were prepared from wholegrain rye (wholegrain diet (WGD)) and non-wholegrain wheat (non-wholegrain diet (NWD)) and fed to four pigs in a crossover design. Plasma samples were collected after 7 d on each diet, and 1H NMR spectra were acquired on these. Partial least squares regression discriminant analysis (PLS-DA) on spectra obtained for plasma samples revealed that the spectral region at 3.25 parts per million dominates the differentiation between the two diets, as the WGD is associated with higher spectral intensity in this region. Spiking experiments and LC–MS analyses of the plasma verified that this spectral difference could be ascribed to a significantly higher content of betaine in WGD plasma samples compared with NWD samples. In an identical study with the same diets, urine samples were collected, and 1H NMR spectra were acquired on these. PLS-DA on spectra obtained for urine samples revealed changes in the intensities of spectral regions, which could be ascribed to differences in the content of betaine and creatine/creatinine between the two diets, and LC–MS analyses verified a significantly lower content of creatinine in WGD urine samples compared with NWD urine samples. In conclusion, using an explorative approach, the present studies disclosed biochemical effects of a wholegrain diet on plasma betaine content and excretion of betaine and creatinine.

NMR: Betaine: Creatinine: Wholegrain: Metabolomics

Epidemiological studies have revealed a connection between the consumption of diets rich in wholegrain cereals and decreased incidences of several degenerative ‘Western’ diseases, and there is strong evidence that wholegrain foods protect against CVD and certain cancers (Anderson et al. 2000; Levi et al. 2000; Slavin et al. 2001; Truswell, 2002), and increasing data are in support of a protective effect against CVD and certain cancers (Anderson et al. 2000; Levi et al. 2000; Slavin et al. 2001; Truswell, 2002), and there is strong evidence that wholegrain foods protect against CVD and certain cancers (Anderson et al. 2000; Levi et al. 2000; Slavin et al. 2001; Truswell, 2002), and increasing data are in support of a protective effect against type II diabetes (Fung et al. 2002; McKeown et al. 2002; Pereira et al. 2002). The food factor responsible for the preventive effects of wholegrain is still unknown. However, wholegrain cereals are rich in dietary fibre, vitamins and minerals, and bioactive compounds such as phytochemicals and micronutrients with potential effects on metabolism and cancer development (Slavin et al. 1999).

NMR is a non-destructive and non-selective technique that can be applied on a variety of materials. Since the first demonstration of the use of 1H NMR spectroscopy on whole blood and blood plasma (Nicholson et al. 1983), 1H NMR spectroscopy of biofluids has proven an excellent tool for investigation of biochemical changes associated with drug toxicity in animal models (Holmes et al. 1998; Lenz et al. 2000; Waters et al. 2001; Beckwith-Hall et al. 2002; Keun et al. 2002; Beckonert et al. 2003; Coen et al. 2003; Ebbels et al. 2003). Promisingly, the technique has recently also been applied to study the biochemical effects of the bioactive compound epicatechin in a rat model, which allowed a demonstration of the overall endogenous metabolic effects of epicatechin consumption (Solanky et al. 2003a). Moreover, use of 1H NMR spectroscopy on biofluids to screen the biochemical effects of dietary soya isoflavones in man has also been reported (Solanky et al. 2003b). Accordingly, 1H NMR spectroscopy of biofluids can be considered an emerging and promising science with a level of information that spans the traditional approach for elucidating the biochemical response to a factor that can be ascribed to the fact that the method enables an entire endogenous metabolic profiling, i.e. a metabolomic fingerprinting. In the present study, the use of high-resolution 1H NMR spectroscopy was for the first time introduced for elucidating the biochemical effects of rye- and wheat-based diets using the pig as a model for human subjects. The study was carried out in two series of experiments with catheterized pigs, and 1H NMR spectroscopy was performed...
on plasma samples drawn from the hepatic portal vein and the mesenteric artery as well as on urine samples.

Materials and methods

Breads and diets

The diets were made of wheat soft and crisp bread or rye soft and crisp bread, and are referred to as non-wholegrain diet (NWD) and wholegrain diet (WGD). Wheat and rye soft breads were produced at Nordmills (Nordmills, Cerealia AB, Malmo, Sweden) and wheat and rye crisp bread were produced at Wasa Bread (Wasa Bread AB, Sweden). The rye crisp bread contained wholegrain rye flour, rye bran (Wasa T2, Sweden), fat and salt as the main ingredients and the corresponding wheat crisp bread white wheat flour, purified wheat fibre essentially as cellulose (Vitacel WF 600; Rettemair and Söhne, Holzmühle, Germany), sugar (sucrose), salt and dry malt. The soft rye bread contained white wheat flour, rye bran (B3-fin; Nordmills), baker’s yeast, fat, salt and sugar and the corresponding soft wheat bread white wheat flour, vitacel, baker’s yeast, fat, salt and sugar. Immediately after production, the soft bread was frozen at −20°C until consumption, while crisp bread was stored dry. The diets prepared from soft and crisp bread were fortified with vitamins and minerals and provided approximately 19, 15 and 66 % energy from fat, protein and carbohydrates, respectively (see Table 1 for diet ingredients and composition). Compared with diets for optimal growth of pigs at the present physiological state, the diets provided approximately 45 % of the recommended concentration of lysine (first limited amino acid) but approximately 260 % of the concentration of lysine for maintenance.

Table 1. Ingredients and chemical composition of the experimental diets (units/kg DM)*

<table>
<thead>
<tr>
<th>Ingredients</th>
<th>NWD</th>
<th>WGD</th>
</tr>
</thead>
<tbody>
<tr>
<td>Wheat crisp bread (g)</td>
<td>602</td>
<td>661</td>
</tr>
<tr>
<td>Wheat soft bread (g)</td>
<td>370</td>
<td>349</td>
</tr>
<tr>
<td>Rye crisp bread (g)</td>
<td>593</td>
<td>568</td>
</tr>
<tr>
<td>Rye soft bread (g)</td>
<td>379</td>
<td>349</td>
</tr>
<tr>
<td>Vitamins/minerals (g)</td>
<td>28</td>
<td>28</td>
</tr>
<tr>
<td>Ash (g)</td>
<td>33</td>
<td>56</td>
</tr>
<tr>
<td>Protein (N × 6.25) (g)</td>
<td>119</td>
<td>127</td>
</tr>
<tr>
<td>Fat (g)</td>
<td>68</td>
<td>73</td>
</tr>
<tr>
<td>Total carbohydrates (g)</td>
<td>777</td>
<td>717</td>
</tr>
<tr>
<td>Sugars (g)</td>
<td>24</td>
<td>40</td>
</tr>
<tr>
<td>Fructans (g)</td>
<td>4</td>
<td>18</td>
</tr>
<tr>
<td>Starch (g)</td>
<td>529</td>
<td>455</td>
</tr>
<tr>
<td>Total NSP and soluble NSP (g)</td>
<td>220</td>
<td>235</td>
</tr>
<tr>
<td>Klasion lignin (g)</td>
<td>10</td>
<td>32</td>
</tr>
<tr>
<td>Dietary fibre (NSP + lignin) (g)</td>
<td>230</td>
<td>235</td>
</tr>
<tr>
<td>Total lignins (mg)</td>
<td>3-1</td>
<td>35-2</td>
</tr>
<tr>
<td>Alkylresorcinols (mg)</td>
<td>0</td>
<td>1-2</td>
</tr>
<tr>
<td>Betaine (g)</td>
<td>0.96</td>
<td>1.92</td>
</tr>
</tbody>
</table>

*NWD, non-wholegrain diet; WGD, wholegrain diet.

Chemical analyses of diets were performed with classical methods as described in the papers by Bach Knudsen et al. (2003, 2005), Saarinen et al. (2001) and Linko et al. (2006).

Experimental design and pigs

The pigs used in both studies were from the Danish Institute of Agricultural Sciences swine herd, Foulum, Denmark.

Study 1. The study was a crossover design with four male castrated pigs (body weight 44.6 (sd 2.4) kg) fed the two diets for 1 week. Each pig was surgically fitted with two catheters, one in the portal vein (1.2 mm i.d.) and the second in the mesenteric artery (1.0 mm i.d.), and with an ultrasonic blood flow probe (14 mm, Transonic System, Ithaca, NY, USA) around the portal vein. The pigs were given Streptocillin for up to 4 d after surgery. After 10 d post-surgery recovery, the pigs were gradually introduced to the two experimental diets and fed 1250 g DM/d. The bread was cut into pieces, mixed 1:2.5 (w/w) with water and fed in equal amounts three times daily, at 07.00, 15.00 and 22.00 hours. For the present study we used portal and arterial blood samples taken 60 min after the morning feeding on day 7. The blood was collected in heparinized plastic tubes and centrifuged (3000 g at 8°C for 10 min) to separate the erythrocytes from plasma. The plasma was kept frozen at −20°C until analysis.

Study 2. Six female pigs with an initial body weight of 38.7 (sd 2.7) kg were used for the study. The study was designed as a crossover experiment with washout periods before, between and after the dietary interventions. The pigs were fed 1 week with a semi-synthetic diet (Bach Knudsen et al. 2003) in periods 1, 3 and 5, whereas NWD and WGD were fed for 2 weeks in periods 2 and 4. The breads were treated as in study 1 and fed in equal amounts twice daily (09.30 and 21.30 hours) together with a fixed amount of water (1:2.5). In the diet intervention periods (periods 2 and 4), the pigs were equipped with a urine bladder catheter for urine collection (5-0 mm i.d.). All urine produced for 3 d was collected in benzoic acid via catheters. The total amount of urine collected for each pig was registered. No significant effect of diet on the amount of urine was found.

The animal experiments complied with the guidelines of the Danish Ministry of Justice with regard to animal experimentation and care of animals under study.

NMR measurements

The NMR measurements were performed at 300 K on a Bruker Avance 400 NMR spectrometer (Bruker BioSpin GmbH, Rheinstetten, Germany), operating at a 1H frequency of 400-13 MHz, and equipped with a standard 5 mm HX inverse probe.

Prior to the measurements, portal and arterial plasma samples (n 16) were thawed and 200 μl aliquots were mixed with 400 μl D2O. Sodium trimethylsilyl-[2,2,3,3-2H4]-1-proline was added as an internal chemical shift standard (0.7 mg/ml). 1H NMR spectra of plasma samples were obtained using a Carr–Purcell–Meiboom–Gill delay added in order to attenuate broad signals from high molecular weight components. The total Carr–Purcell–Meiboom–Gill delay was 40 ms and the spin-echo delay was 200 μs. Urine samples (n 12) were thawed and 400 μl aliquots were mixed...
with 200 μl D2O, and sodium trimethylsilyl-[2,2,3,3-2H4]-1-propionate was added as an internal chemical shift standard (0.7 mg/ml). NMR measurements on plasma samples were also performed upon spiking of plasma samples with betaine (Sigma Chemical Co., St Louis, MO, USA), glycerophosphorylcholine (Sigma Chemical Co.), enterolactone (Labmaster Ltd, Turku, Finland), and alkylresorcinol C15:0 (Kamal-Eldin et al. 2000), respectively. 1H NMR spectra of urine samples were obtained using a single 90° pulse experiment. Both on plasma and urine samples water suppression was achieved by irradiating the water peak during the relaxation delay of 2 s. A total of 128 transients of 8 K data points spanning a spectral width of 24.03 parts per million (ppm) were collected. An exponential line-broadening function of 3 Hz was applied to the free induction decay prior to Fourier transformation effects.

Post-processing of NMR data

The mean-normalized 1H NMR spectra in the region 9.0–5.1 ppm and the region 4.6–0.5 ppm were used for further data analysis. The spectra were segmented into regions of 0.04 ppm width and the integral of each region was calculated. The reduced spectra consisting of 195 integrated regions were normalized to the whole spectrum to remove any concentration effects.

LC–MS analyses

Portal and arterial plasma samples, taken after 7 d of feeding with the two diets, from two pigs (n 8) were analysed for betaine content, and all urine samples (n 12) were analysed for creatinine content using LC–MS. Original plasma sample (200 μl) was pipetted and transferred into a Whatman vial (Chromtech GmbH, Idstein, Germany); 500 μl acetonitrile (containing 0.1 % formic acid) were added. After spontaneous flocculation of the proteins the plunger was inserted into the vial and pressed down, thus forcing the liquid upward through the membrane. Urine samples of 400 μl were prepared using the same procedure but without addition of acetonitrile to the samples. Analyses were performed on an Agilent (Agilent, Waldbronn, Germany) HPLC–diode-array detector–MS station equipped with an HPLC series 1100 comprising a model G1312A binary pump, a model G1379A micro vacuum degasser, a model G1327A thermostat column compartment, a model G1315B diode-array detector and a model G2707DA LC/MSD SL detector fitted with a model G1948A API-electrospray source. The station was controlled and the results were monitored with Agilent’s ChemStation software (Rev. A.10.02).

Sample separations were carried out on a Varian (Lake Forest, CA, USA) OmniSpher C18 column (100 × 2.0 mm i.d., 3 μm particle size) operated at a temperature of 35°C and with a 0.2 ml/min flow. The solvents used were: (A) aqueous 1% formic acid (formic acid of analytical-reagent grade; Riedel-de-Haën, Seelze, Germany; (B) 1% formic acid in HPLC-grade acetonitrile, using a gradient programme as follows: 0% B isocratic (3 min), 0% B to 98% B (4 min), 98% B isocratic (6 min). The injection volume was 1 μl.

UV spectra were recorded at 214 and 254 nm at a rate of 1.25 scans/s.

MS spectra of samples were recorded simultaneously in SCAN (from m/z 50 to 500) and SIM (ions with m/z 114 and 118) positive modes. The acquisition parameters were as follows: fragmentor 160 V, gain 5 EMV, stepsize 0.10. N was used as drying gas at a flow of 11 l/min and as nebulizing gas at a pressure of 54 psig and a temperature of 280°C. A potential of 3000 V was used on the capillary.

Optimization of ionization and fragmentation parameters was carried out by running a fluoroimmunoassay with standard betaine and creatinine (Sigma Chemical Co).

Statistics

Statistical analysis of LC–MS data was performed by the Statistical Analysis Systems package version 8.2 (SAS Institute, Cary, NC, USA) using the MIXED procedure. A model including the fixed effect of diet (WGD v. NWD) and the random effect of animal was applied. Significant differences are indicated, and the P value is given.

Multivariate data analysis was performed using the Unscrambler software version 8.0 (Camno, Oslo, Norway). Principal component analysis was applied to the centred data to explore any clustering behaviour of the samples, and partial least square regression discriminant analysis (PLS-DA) was performed to explore intrinsic biochemical dissimilarities between predefined sample classes (WGD diet v. NWD diet). During all regressions, Martens uncertainty test (Martens & Dardenne, 1998) was used for jack-knifing, and all models were validated using full cross-validation.

Results

Plasma samples

Principal component analysis, which is an unsupervised method, was performed on the pre-processed 1H NMR spectra, and the resulting plot of score 2 v. score 3 for mean-centred data is shown in Fig. 1 with each spectrum represented by a
single data point. A clear tendency to clustering according to diet is observed. PLS-DA, which is more focused on discriminating variation between pre-defined classes than the unsupervised principal component analysis approach, was performed on the $^1$H NMR spectra to investigate the metabolic differences in plasma profile between WGD and NWD. The PLS-DA score plots of the first and second component show a clear separation of NWD and WGD plasma (Fig. 2(A)). The regions of the NMR spectrum that most strongly influence separation between WGD and NWD plasma are evident in the regression coefficients (Fig. 2(B)). A positive value indicates a relatively greater concentration of metabolite in WGD samples, and a negative value indicates a relatively lower concentration compared with NWD samples. The spectral region at 3.25 ppm dominates the differentiation between the two diets. Jack-knifing reveals that three regions influence the discrimination between WGD and NWD samples significantly: 3.54, 3.25 and 1.28 ppm. Of these regions, the concentration of the compound at 3.54 ppm is lower and the concentration of the compound at 3.25 and 1.28 ppm is higher in WGD plasma samples compared with NWD samples. The major regions showing differences between WGD and NWD plasma are summarized along with a tentative metabolite assignment in Table 2. The concentration of betaine was confirmed and quantified in plasma samples from two pigs after the intake of both rye- and wheat-based diet using LC–MS. A significantly higher content of betaine ($P = 0.0012$) was found in WGD plasma samples (5.53 (SD 0.46) μg/ml) compared with NWD plasma samples (2.10 (SD 0.46) μg/ml).

**Urine samples.** Principal component analysis was performed on the pre-processed $^1$H NMR spectra, and the resulting plot of score 1 vs. score 2 for mean-centred data is shown in Fig. 3. A clear tendency to clustering according to diet is observed along the first score. PLS-DA was performed on the $^1$H NMR spectra to investigate the metabolic differences in urine profile between WGD and NWD. The PLS-DA score plots of the first and second component show a clear separation of NWD and WGD plasma (Fig. 4(A)). The regions of the NMR spectrum that most strongly influence separation between WGD and NWD plasma are evident in the regression coefficients (Fig. 4(B)). A positive value indicates a relatively higher concentration of metabolite in WGD samples, and a negative value indicates a relatively lower concentration compared with NWD samples. The spectral region at 3.25 ppm dominates the differentiation between the two diets. The major regions showing differences between WGD and NWD urine are summarized along with a tentative metabolite assignment in Table 2. The concentration of creatinine in the urine samples was confirmed and quantified using LC–MS. A significantly higher content of creatinine ($P = 0.03$) was found in NWD urine samples (69.95 (SD 5.86) μg/ml) compared with WGD urine samples (48.71 (SD 5.86) μg/ml).

**Spiking experiments**

$^1$H NMR spectra were obtained on plasma after spiking with betaine, glycerophosphorylcholine, enterolactone and alkylresorcinol, respectively, and Table 3 summarizes the results. Betaine intensified two resonances at chemical shift values (3.25 and 3.92 ppm) identical to those resonances in the plasma. In contrast, glycerophosphorylcholine showed three resonances at chemical shift values (3.20, 3.62 and 4.15 ppm) where no resonances otherwise are clearly evident in the plasma. Finally, enterolactone resulted in an additional resonance at 3.56 ppm, while spiking with alkylresorcinol resulted in an enhancement in resonances at 0.9 ppm, 1.3 ppm, probably originating from the methyl and methylene protons in the alkyl part, and at 3.21 ppm, probably originating from protons in the resorcinol part.

**Discussion**

It is well established that a relationship exists between the consumption of wholegrain cereals and incidences of ‘Western’ diseases, CVD, certain types of cancers and diabetes (Anderson et al. 2000; Levi et al. 2000; Slavin et al. 2001; Fung et al. 2002; McKeeown et al. 2002; Pereira et al. 2002; Truswell, 2002). However, the exact physiological mechanisms responsible for this relationship are still not understood. Pigs are recognized to be excellent models for human nutrition because of similarities with regard to digestive anatomy and many aspects of physiology (Miller & Ulrey, 1987). The present study is the first to report the use of $^1$H NMR spectroscopy on plasma and urine samples to study the metabolic differences between pigs fed a rye-based diet (WGD), rich in wholegrain and bran, and a wheat-based diet (NWD) with similar levels of dietary fibres, but with
contrasting dietary fibre characteristics (Bach Knudsen et al. 2005) and phytochemicals (Bach Knudsen et al. 2003; Linko et al. 2006) and where we, with conventional approaches, have identified marked differences in the plasma profiles of mammalian lignans, alkylresorcinols and SCFA but not in the levels of glucose, lactate and insulin (Bach Knudsen et al. 2003; Linko et al. 2006). Principal component analysis on spectra acquired on both plasma and urine samples revealed a clear separation of WGD and NWD samples (Figs 1 and 3), implying a significant effect on the metabolite plasma and urine profile of the two diets. In order to investigate the differences in the plasma metabolite profile of WGD and NWD plasma samples, PLS-DA, which is a supervised method, was performed on spectra obtained with WGD and NWD plasma samples (Fig. 2(A,B)).

The regression coefficients revealed that PLS-DA could discriminate between WGD and NWD plasma samples mainly because of increases in spectral intensities of regions around 3·25 ppm and around 1·20–1·30 ppm in WGD plasma samples compared with NWD plasma samples; 3·25 ppm corresponds to the chemical shift value reported for different N(CH3)3 groups (Lindon et al. 1999), and to investigate if this response should be ascribed to betaine, spiking of plasma with betaine was investigated, which confirmed that betaine has a resonance at 3·25 ppm (Table 3). Accordingly, it appears that the spectral difference at 3·25 ppm between the two diets should be ascribed to a higher level of betaine in WGD plasma compared with NWD plasma. This was confirmed by LC–MS analyses of the plasma samples, which verified a significantly higher content of betaine in WGD samples compared with NWD samples. Moreover, spiking of plasma with lignans (enterolactone) and alkylresorcinol (C15 : 0) were also scrutinized, which revealed that the protons of the resorcinol part give rise to a resonance at 3·22 ppm, which is relatively close to the main difference observed at 3·25 ppm. Because of the fact that resonances from the alkyl chain (approximately 1·3 ppm) were also observed upon spiking with alkylresorcinol (Table 3), and that this resonance also was found to play a role

Table 2. Summary of the major differences between non-wholegrain diet (NWD) and wholegrain diet (WGD) samples*

<table>
<thead>
<tr>
<th>NMR spectral region (ppm)</th>
<th>Assignment†</th>
<th>Plasma‡</th>
<th>Urine‡</th>
</tr>
</thead>
<tbody>
<tr>
<td>approximately 1·2–1·4</td>
<td>(CH3)3 in fatty acids</td>
<td>↑</td>
<td>–</td>
</tr>
<tr>
<td>3·03</td>
<td>N-CH3 in creatinine/creatinine</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>3·25</td>
<td>N-CH3 in betaine</td>
<td>↑</td>
<td>↑</td>
</tr>
<tr>
<td>approximately 3·5–4·5</td>
<td>CH4 in glucose and amino acids</td>
<td>↑</td>
<td>↑</td>
</tr>
<tr>
<td>3·90–3·94</td>
<td>CH3 in betaine</td>
<td>↑</td>
<td>–</td>
</tr>
<tr>
<td>4·06–4·10</td>
<td>N-CH3 in creatinine</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>5·69–5·81</td>
<td>NH2 in urea</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>7·54</td>
<td>CH(CH3)2 in hippurate</td>
<td>–</td>
<td>↑</td>
</tr>
<tr>
<td>7·83</td>
<td>CH2(CH3)6 in hippurate</td>
<td>–</td>
<td>↑</td>
</tr>
</tbody>
</table>

* For details of procedures and diets, see p. 956 and Table 1.
† Based on Lindon et al. (1999) and Griffin et al. (2004).
‡ The symbols ↑ and ↓ indicate higher and lower concentrations in WGD samples compared with NWD samples, respectively; – indicates no effect of diet.

Fig. 3. Principal component analysis score plot showing the two first principal components for non-wholegrain diet (●) and wholegrain diet (○) urine sample groups. For details of procedures, see p. 956.

Fig. 4. (A), Partial least squares regression discriminant analysis (PLS-DA) scores plot from analysis of NMR spectra obtained on non-wholegrain diet (●) and wholegrain diet urine samples (○). For details of procedures, see p. 956. (B), The regression coefficients of the PLS-DA shown in (A).
in the discrimination of the two diets (Fig. 2(B)), it cannot be completely excluded that alkylresorcinol also contributes to the main difference at 3·25 ppm. However, taking into consideration that plasma alkylresorcinol concentrations are reported to be in the nanomolar range (Linko et al. 2006), which most probably is below the limit of NMR detection (Nicholson & Wilson, 1989), and about 1000 times lower than the plasma concentration of betaine found in the present study, it seems most likely the main difference at 3·25 ppm primarily should be ascribed to betaine. Previous studies have shown that dietary betaine is absorbed and causes increases in serum concentrations (Frontiera et al. 1994; McGregor et al. 2002; Schwab et al. 2002; Schwahn et al. 2003), which supports the present findings of betaine absorption. Betaine has three reactive methyl groups, and works as a methyl donor in many biochemical pathways and participates in the methionine cycle (Anon, 2003). Insufficient dietary intake of methyl groups leads to hypomethylation, which may contribute to various diseases, including coronary, cerebral, hepatic and vascular diseases (Craig, 2004). In addition, betaine also has other physiological functions, is suggested to protect against protein denaturation and has been termed a ‘chemical chaperone’ (Caldas et al. 1999), and it helps maintain cell volume under osmotic stress (Craig, 2004). Several positive effects of betaine have been reported, as betaine may protect internal organs (Caldas et al. 1999) and improve vascular risk factors (Borsook & Borsook, 1951; Schwab et al. 2002; Olthof et al. 2003; Steenge et al. 2003). Moreover, it has been demonstrated that betaine supplementation improved athletic performance (Armstrong et al. 2003). The present study is the first to report that a wholegrain diet affects serum betaine levels. In view of the physiological significance of betaine, the present findings must be considered extremely important. Further studies are needed to elucidate the precise role of betaine in promoting health.

Urine samples were also analysed to elucidate the biochemical effects of the two diets. Analysis of the regression coefficients revealed that PLS-DA could discriminate between WGD and NWD urine samples because of increases in spectral intensities of betaine and hippurate, and decreases in spectral intensities of creatinine and urea and in WGD urine samples compared with NWD samples (Table 2). The lower content of creatinine in WGD urine samples compared with NWD urine samples was verified by LC–MS analyses. Possibly the lower creatinine excretion in WGD samples is an indicator of alterations in protein turnover and metabolism in general. However, the reverse relationship between betaine concentrations in plasma and urinary creatinine excretion in WGD samples may also be related. Even though several studies have investigated the various effects of betaine (Borsook & Borsook, 1951; Keshavarz & Fuller, 1971; Burg, 1995; Matthews et al. 2001; McGregor et al. 2002; Schwab et al. 2002; Armstrong et al. 2003; Kim et al. 2003; Ozturk et al. 2003; Schwahn et al. 2003; Slow et al. 2004; Wray-Cahen et al. 2004), to the authors’ knowledge, a latent relationship between plasma betaine content and urinary excretion of creatinine has not been elucidated, but might be a result of three different mechanisms. First, betaine is a well-known osmoregulatory biomolecule (Craig, 2004), and therefore might simply reduce creatinine clearance to the urine. Second, another explanation might be ascribed to the fact that methyl donation is required in the biosynthesis of creatine, and creatinine is the metabolic end-product of creatine catabolism. Accordingly, an interaction between plasma betaine content and creatinine excretion is likely, potentially through homocysteine metabolism. A higher content of muscle creatine upon betaine supplementation has been demonstrated in chicks (Chamruspollert et al. 2002). However, it is not understood why a betaine-induced increase in muscle creatine should reduce the urinary excretion of creatinine. Third, it has been hypothesized that biomolecules with an electrophilic methyl group, e.g. as found in betaine, may represent an important control system for in vivo redox balance (Ghyczy & Boros, 2001), and it has been shown that betaine treatment decreases oxidative stress in the liver of ethanol-treated guinea pigs (Balkan et al. 2004), and in children with increased vulnerability to oxidative stress as a consequence of autism (James et al. 2004). Considering that betaine improves the redox balance in the rye-fed pigs, resulting in decreased oxidative stress, this might be consistent with the lower urine creatinine excretion in these animals, as oxidative stress has been reported to be inversely correlated to creatinine clearance (Antolini et al. 2004). No matter whether the content of plasma betaine and the level of urine creatinine is independent or a result of some of the hypotheses mentioned earlier, these aspects need to be studied further to exploit the biological consequences of the two diets further.

In conclusion, using NMR-based metabolomics as an explorative approach on the pig as a human model, the present study disclosed metabolic effects of a wholegrain diet on the content of betaine in plasma and excretion of betaine and creatinine, which may contribute to the health benefits of a high dietary intake of wholegrain. Further studies emphasizing the beneficial role of betaine and its potential connection with creatinine excretion in the health-promoting effect of wholegrain cereals are needed.

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