In vitro ileal and caecal fermentation of fibre substrates in the growing pig given a human-type diet

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(Submitted 30 April 2020 – Final revision received 8 August 2020 – Accepted 1 September 2020 – First published online 11 September 2020)

Abstract
This study characterised the in vitro ileal fermentability of different substrates in the growing pig, adopted as an animal model for the adult human, and compared in vitro ileal and caecal fermentation in the pig. Substrates (arabinogalactan (AG), cellulose, fructo-oligosaccharide (FOS), inulin, mucin, citrus pectin and resistant starch) were fermented in vitro (ileal 2 h and caecal 24 h) with an ileal or caecal inoculum prepared from ileal or caecal digesta collected from growing pigs (n 5) fed a human-type diet for 15 d. The organic matter (OM) fermentability and production of organic acids were determined. In general, there was considerable in vitro ileal fermentation of fibre, and the substrates differed (P < 0.001) for both in vitro ileal and caecal OM fermentability and for organic acid production. Pectin had the greatest in vitro ileal OM fermentability (26%) followed by AG, FOS and resistant starch (15% on average), and cellulose, inulin and mucin (3% on average). The fermentation of FOS, inulin and mucin was greater for in vitro caecal fermentation compared with the ileal counterpart (P ≤ 0.05). In general, the organic acid production was higher for in vitro caecal fermentation (P ≤ 0.05). For instance, the in vitro ileal acetic acid production represented 4–66 % of in vitro caecal production. Energy from fibre fermentation of 0.6–11 kJ/g substrate fermented could be expected in the ileum of the pig. In conclusion, substrates are fermented in both the ileum and caecum. The degree of fermentation varies among substrates, especially for in vitro ileal fermentation.

Key words: Fermentability: In vitro ileal and caecal fermentation: Organic acids: Dietary fibre

It is commonly held that dietary fibre fermentation in monogastric animals, including humans, occurs mainly in the large intestine with little fermentation occurring in the foregut (stomach and small intestine). This, however, may not be the case(1–3). A combined in vivo/in vitro ileal fermentation assay has been developed using the growing pig as an animal model for the adult human(4). The assay is based on collecting ileal digesta as this part of the small intestine has a high population of microbes and is associated with a longer digesta retention time when compared with other foregut regions. In the assay, ileal digesta are collected for the preparation of an inoculum, which is used to ferment in vitro the substrate collected from the lower jejunum. The combined in vivo/in vitro fermentation assay has been applied to estimate the amount of organic matter (OM) fermented for a human-type diet in the ileum and caecum(5). There appears to be considerable ileal fermentation for a typical human diet in the growing pig, with ileal-fermented OM potentially being 1.5-fold greater than caecal-fermented OM. This work underlines the possible importance of ileal fermentation in the pig, but little is known about the ileal fermentation of different types of fibre substrates.

Dietary fibres have different monomeric units with specific links (e.g. α-1,4-linked glucose for resistant starch, β-2,1-linked fructose for fructo-oligosaccharide, FOS). These structural differences can affect their rate and extent of fermentation by the gastrointestinal (GIT) microbial population(6–8). For instance, it has been reported that one-third of purified citrus pectin was fermented in vitro with an ileal inoculum from growing pigs(9). Due to structural differences in fibre types, it is hypothesised that ileal fermentation may differ among fibre substrates leading to a different production of organic acids. Thus, different energy contributions are expected across fibre substrates during ileal fermentation.

Differences in the rate and extent of fermentation of dietary fibres can also be influenced by the composition of the microbial population(10). For example, the production of acetic, propionic

Abbreviations: AG, arabinogalactan; FOS, fructo-oligosaccharide; GIT, gastrointestinal tract; OM, organic matter.
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and butyric acids differed after in vitro fermentation of FOS, sorghum arabinoxylan and maize arabinoxylan with *Prevotella* spp. and *Bacteroides* spp. predominant faecal bacteria\(^{[11]}\). Considering the differences in microbial populations between the ileum and large intestine\(^{[12,13]}\), the ileal fermentation of dietary fibre sources may differ from caecal fermentation. However, there are no reported studies that have determined both ileal and caecal fermentation across a range of fibre substrates.

In this study, the ileal fermentation of several fibre substrates was characterised in terms of OM fermentability and organic acid production, using an optimised in vitro ileal fermentation assay\(^{[8]}\) and compared with an optimised in vitro caecal fermentation assay\(^{[14]}\). For ileal and caecal inocula, ileal and caecal digesta, respectively, were collected from growing pigs fed a human-type diet. The growing pig was used as an animal model for in vitro ileal fermentability in the adult human due to similarities between species in foregut digestion\(^{[15,16]}\). Similar viable counts of lactobacilli, streptococci and coliforms have been found in the ileal digesta of humans and pigs\(^{[17]}\). Purified fibre substrates for in vitro fermentation were characterised in terms of OM fermentability and organic acid production, using an optimised in vitro ileal fermentation assay\(^{[14]}\). For ileal and caecal fermentation assays optimised for different parameters such as incubation time, amount of digesta and pH were used in this study.

### Experimental methods

#### Animals and dietary treatment

Ethics approval for the animal trial was obtained from the Animal Ethics Committee, Massey University, Palmerston North, New Zealand. Five entire male pigs of 9 weeks of age (Hampshire × (Landrace × Large white) and 23 (so 1·6) kg body weight) were housed individually in metabolism pens (1·5 m × 0·5 m) in a room maintained at 24 (so 2·4)°C with a 10 h light–14 h dark cycle. The pigs were fed a high-fibre human-type diet for 15 d (Table 1). The diet was formulated to meet the nutrient requirements of the growing pig\(^{[18]}\) as described in detail elsewhere\(^{[6]}\), with ingredients commonly used in Western diets.

#### Substrates for in vitro fermentation

Different purified substrates (arabinogalactan (AG), cellulose, FOS, inulin, mlin, citrus pectin and resistant starch) were used for both the in vitro ileal and caecal fermentations. Cellulose, pectin, AG (a model hemicellulose) and resistant starch were selected as they represent important dietary fibre components of a human diet. FOS and inulin were selected to determine if structural differences (i.e. chain length) affect OM fermentability and the production of organic acids. Mucin was selected as it is one of the major contributors to SCFA produced in the hindgut\(^{[19]}\). All substrates, except for resistant starch, were purchased from Sigma. Resistant starch (native high-amylose maize starch) was obtained from Megazyme.

<table>
<thead>
<tr>
<th>Ingredient</th>
<th>g/kg DM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cooked carrot (minced)</td>
<td>100</td>
</tr>
<tr>
<td>Canned beans (minced)</td>
<td>150</td>
</tr>
<tr>
<td>Peeled hard-boiled egg (minced)</td>
<td>120</td>
</tr>
<tr>
<td>Cooked white rice (minced)</td>
<td>550</td>
</tr>
<tr>
<td>Raw apple (minced)</td>
<td>48·5</td>
</tr>
<tr>
<td>Premix of vitamins and minerals†</td>
<td>5</td>
</tr>
<tr>
<td>Sodium chloride</td>
<td>3</td>
</tr>
<tr>
<td>Titanium dioxide</td>
<td>3</td>
</tr>
<tr>
<td>Calcium carbonate</td>
<td>0·5</td>
</tr>
<tr>
<td>Dicalcium phosphate</td>
<td>20</td>
</tr>
<tr>
<td>Nutrient (g/kg DM)</td>
<td></td>
</tr>
<tr>
<td>Crude protein</td>
<td>143</td>
</tr>
<tr>
<td>Total lipid</td>
<td>44·9</td>
</tr>
<tr>
<td>Ash</td>
<td>48·7</td>
</tr>
<tr>
<td>Soluble fibre</td>
<td>19·6</td>
</tr>
<tr>
<td>Insoluble fibre</td>
<td>95·7</td>
</tr>
<tr>
<td>Total dietary fibre</td>
<td>115</td>
</tr>
<tr>
<td>Determined energy (MJ/kg DM)</td>
<td>18·1</td>
</tr>
</tbody>
</table>

* The chemical composition of the ingredients to formulate the diet was obtained from the US Department of Agriculture National Nutrient Database (https://ndb.nal.usda.gov/).
† Vitamin and mineral premixes were obtained from Vittec Nutrition Ltd and supplied (per kg of diet as-fed): Mn, 45 mg; Zn, 80 mg; Cu, 25 mg; Co, 0·5 mg; Se, 0·3 mg; Fe, 100 mg; iodine, 1·0 mg; choline, 100 mg; all-trans retinyl acetate, 3·0 mg; cholecalciferol, 0·05 mg; α-tocopherol, 50 mg; menadione, 2·0 mg; thiamin, 1·0 mg; riboflavin, 3·0 mg; nicotinic acid, 15 mg; pantothenic acid, 20 mg; pyridoxine, 2·0 mg; cyanocobalamin, 0·01 mg; foliac acid, 0·5 mg; biotin, 0·1 mg.

#### Experimental design

**In vivo assay.** Pigs received the human-type diet for 14 d (100 g DM/kg metabolic body weight (body weight\(^{[75]}\) per d), with a gradual adaptation from a commercial to the human-type diet during the first three experimental days. Two equal meals were given per day (08.00 and 16.00 hours) with *ad libitum* access to water. The daily ration was adjusted after 1 week. Pigs were monitored at least twice a day, and cages and toys were cleaned after each meal. On day 15, pigs were fed half their daily ration in a single meal and, at 5 h postprandially, they were anaesthetised and euthanised\(^{[10]}\). The small intestine and caecum were ligated with clamps and immediately removed. Digesta from the last one-third of the small intestine (ileum) and caecum for each individual pig for in vitro fermentation were collected in plastic bags containing CO\(_2\) prior to being stored in insulated containers at \(37^\circ\)C. Aliquots of the digesta were also collected in Eppendorf tubes and stored at \(-80^\circ\)C for microbial analysis.

**In vitro fermentation assays.** In vitro ileal\(^{[6]}\) and caecal\(^{[14]}\) fermentation assays optimised for different parameters such as incubation time, amount of digesta and pH were used in this study. Therefore, parameters differed between in vitro ileal and caecal fermentation assays. Digesta were pooled to prepare the ileal and caecal inocula\(^{[20]}\).

**In vitro ileal fermentation assay.** The inoculum was prepared under a constant flow of CO\(_2\) to maintain anaerobic conditions. Equal amounts of fresh ileal digesta from each pig were weighed. The pooled ileal digesta sample (220 g) was then mixed with 1 litre of a sterilised anaerobic 0·1 M phosphate buffer solution
(plus 1 mg L-cysteine, pH 6.5 at 37°C), homogenised and filtered through four sterilised layers of cheesecloth. The inoculum was stirred constantly. A quantity of 5 ml of the inoculum was then added to a McCartney bottle containing 5 ml phosphate buffer either alone (blank) or containing 0.1 g substrate. Six bottles per blank (initial and final) or per substrate were used. Each bottle was flushed with CO₂, capped and incubated at 37°C for 2 h (13) with the exception of the initial blank that was not incubated. Initial blank bottles and the bottles after fermentation were cooled down on ice. Half of the bottles (blanks and substrates) were placed in an autoclave (121°C for 20 min) before determination of OM to allow calculation of OM fermentability. The other half of the bottles was thoroughly mixed, and an aliquot (1 ml) was collected in an Eppendorf tube. The Eppendorf tubes were centrifuged (14 000 × g for 15 min at 4°C), and the supernatant (0.5 ml) was collected and stored at –20°C for subsequent determination of the organic acids.

**In vitro caecal fermentation assay.** In pigs, terminal ileal digesta are released into the caecum where fermentation with the caecal microbiota occurs. Thus, in this study, caecal digesta instead of faeces were used to prepare the caecal inoculum. The *in vitro* caecal fermentation assay was the same as that for the *in vitro* ileal fermentation assay, with the exception that the inoculum was prepared from a pooled 330 g sample of caecal digesta and the fermentation lasted for 24 h (13). Two substrates (FOS and inulin) were also fermented *in vitro* with the caecal inoculum for 2 h.

**Chemical and microbial analyses**

The diet and substrates were analysed in duplicate for DM, ash and OM (DM minus ash) (13). The diet was analysed for starch (Kit AA/AMG; Megazyme), crude protein (N × 6.25; using an elemental analyser LECO), ether extract (using a Soxhlet apparatus and petroleum ether extraction), gross energy (LECO AC-350 Automatic Calorimeter), soluble and insoluble dietary fibres (22), DM, ash and OM contents were also determined on the material remaining after the *in vitro* fermentation.

Organic acids (formic, acetic, propionic, butyric, isobutyric, valeric, isovaleric and lactic acids) were determined in *in vitro* fermentation samples using GC (23). The ileal and caecal digesta were analysed for total bacteria using quantitative PCR and bacterial community composition (Illumina) and number of glycoside hydrolases (*in silico*) as follows:

\[
\text{Number of bacteria}_{\text{genus}} (16S \text{ rRNA gene copy number of bacteria}_{\text{genus}}/g \text{ DM}) = 16S \text{ rRNA gene copy number of total bacteria/g DM} \\
\times \text{ relative abundance coefficient}_{\text{genus}}
\]

\[
\text{Number of glycoside hydrolase}_{\text{family}_i} = \sum (\text{number of bacteria}_{\text{genus}_j} \times \text{glycoside hydrolase}_{\text{family}_i})
\]

where glycoside hydrolase_{family}_i is the number of sequences in each glycoside hydrolase family for the genus j.

**Statistical analysis**

Based on the known means and variances for the *in vitro* ileal fermentation of pectin and cellulose (30 and 3.7 (SD 1.13 %), two *in vitro* replicates were required to detect a statistical difference between treatments with a power > 80 % at a two-tail 5 % significance. In this study, three technical replicates were used.

The statistical analyses were performed using the Mixed Model procedure of SAS (SAS/STAT version 9.4). A two-way ANOVA model was used to determine the effects of inoculum (ileal v. caecal), substrate (AG, cellulose, FOS, inulin, mucin, pectin and starch) and their interaction on OM fermentability, organic acid production and energy from organic acids. When the interaction effect was not statistically significant, it was removed from the final model.

The model diagnostics for each response variable were tested using the ODS Graphics procedure and the Repeated statement.
of SAS. The Repeated statement in the Mixed Model procedure allowed testing for the homogeneity of variance by fitting models with the Restricted Maximum Likelihood method and comparing them using the log-likelihood ratio test. The selected model for all response variables had similar Studentised residuals (i.e. equal variances) across treatments. When the F-value of the model was significant \((P \leq 0.05)\), the means were compared using Least Significant Difference. Correlations were determined between OM fermentability and total energy from organic acids \((n = 21)\) for each GIT location using the Correlation procedure of SAS.

**Results**

All pigs remained healthy throughout the study and once adapted to their diet consumed all of their daily ration.

The microbial population of the ileal digesta at the genus level mainly comprised *Streptococcus* (67%), while the caecal digesta mainly comprised *Streptococcus* (15%), *Ruminococcaceae* (15%), *Ruminococcus* (15%), *Clostridiales* (11%) and *Prevotella* (9%) (online Supplementary Fig. S1).

There was an interaction \((P < 0.001; \text{Fig. 1})\) between the inoculum and substrate for in vitro OM fermentability. **During in vitro ileal fermentation**, pectin had the highest fermentability (26%), \(P \leq 0.05\) followed by AG, FOS and resistant starch, while cellulose, inulin and mucin had the lowest fermentability. In contrast, during in vitro caecal fermentation, FOS, cellulose, inulin, mucin, pectin and resistant starch had the highest \((P \leq 0.05)\) fermentation, while AG and cellulose had the lowest fermentation. The fermentation of FOS, inulin and mucin was higher (17, 26 and 24% units, respectively; \(P \leq 0.05\)) during in vitro caecal fermentation than during in vitro ileal fermentation, whereas the fermentation of AG, cellulose, pectin and starch was similar \((P > 0.05)\) for in vitro ileal and caecal fermentations.

To investigate the effect of incubation time during in vitro caecal fermentation, FOS and inulin were fermented in vitro with the caecal inoculum for both 2 and 24 h. The extent of in vitro caecal OM fermentability during 2 h was 16 and 14% for FOS and inulin, respectively.

There was an interaction \((P \leq 0.05; \text{Table 2})\) between inoculum and substrate for the in vitro production of acetic, propionic, butyric, isovaleric and lactic acids. For acetic acid, FOS, inulin and mucin in general had the highest \((P \leq 0.05)\) in vitro ileal and caecal productions, though mucin had one of the lowest in vitro caecal acetic acid productions \((P > 0.05)\). For propionic, butyric and isovaleric acids, there were no differences \((P > 0.05)\) between substrates during in vitro ileal fermentation. However, during in vitro caecal fermentation, there were large differences \((P \leq 0.05)\) between substrates. For instance, the in vitro caecal production of propionic acid was 3.3-fold higher for FOS when compared with cellulose. For lactic acid, resistant starch had the highest \((P \leq 0.05)\) in vitro ileal and caecal productions, while FOS (ileal fermentation) and pectin (caecal fermentation) had the lowest. In general, the in vitro caecal production of acetic, propionic and butyric acids was higher \((P \leq 0.05)\) than the in vitro ileal counterpart. In contrast, the in vitro caecal production of isovaleric and lactic acids was in general either similar \((P > 0.05)\) or lower \((P \leq 0.05)\) than the in vitro ileal counterpart.

There was no interaction \((P > 0.05)\) between inoculum and substrate for the in vitro production of formic, isobutyric and valeric acids (Table 2). The in vitro fermentation of AG rendered a very high \((P < 0.001)\) production of formic acid (>50-fold compared with other substrates), followed by pectin and FOS. The production of formic acid was 5% higher for in vitro ileal fermentation compared with in vitro caecal fermentation \((P < 0.001)\). In contrast, the production of isobutyric and valeric acids was 10.3- and 8.4-fold higher for in vitro caecal fermentation in comparison with in vitro ileal fermentation \((P \leq 0.05)\), with no difference between substrates for their in vitro productions \((P > 0.05)\).

The calculated gross energy of the organic acids produced during in vitro fermentation reflected the interaction between the inoculum and the substrate for OM fermentability \((P < 0.001; \text{Fig. 2})\). During in vitro ileal fermentation (2 h), AG, inulin, mucin and resistant starch had higher \((P \leq 0.05)\) energy productions (energy content of organic acid production) than FOS and pectin, while during in vitro caecal fermentation (24 h), resistant starch had the highest \((P \leq 0.05)\) gross energy production followed by FOS and inulin, while cellulose and mucin had the lowest. For all substrates, apart from cellulose, more energy \((P \leq 0.05)\) from organic acids was obtained during in vitro caecal fermentation than in vitro ileal fermentation. The gross energy production for in vitro ileal fermentation represented between 11% (FOS and starch) and 100% (cellulose) of the energy produced during in vitro caecal fermentation.

**Discussion**

The study aimed to characterise the in vitro ileal fermentability of several purified substrates in the growing pig, selected as an animal model for ileal fermentation in the adult human. The quantitative significance of in vitro ileal fermentation for the pig was gauged by comparison with pig in vitro caecal fermentation.
### Table 2. Organic acid production during in vitro ileal and caecal fermentation of different purified substrates. Ileal and caecal digesta were collected from pigs fed a high-fibre human-type diet (Mean values with their standard errors)

<table>
<thead>
<tr>
<th>Inoculum</th>
<th>Formic acid (mmol/kg substrate)</th>
<th>Acetic acid (mmol/kg substrate)</th>
<th>Propionic acid (mmol/kg substrate)</th>
<th>Butyric acid (mmol/kg substrate)</th>
<th>Isovaleric acid (mmol/kg substrate)</th>
<th>Valeric acid (mmol/kg substrate)</th>
<th>Lactic acid (mmol/kg substrate)</th>
<th>Total organic acids (mmol/kg substrate)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
<td>Mean SEM</td>
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<td></td>
<td>S</td>
<td>I</td>
<td>S</td>
<td>I</td>
<td>S</td>
<td>I</td>
<td>S</td>
<td>I</td>
</tr>
<tr>
<td><strong>Substrate</strong></td>
<td><strong>AG</strong></td>
<td><strong>Cellulose</strong></td>
<td><strong>FOS</strong></td>
<td><strong>Inulin</strong></td>
<td><strong>Mucin</strong></td>
<td><strong>Pectin</strong></td>
<td><strong>RS</strong></td>
<td><strong>Substrate</strong></td>
</tr>
<tr>
<td>Ileal</td>
<td>2.99 ± 0.51</td>
<td>7.34 ± 0.95</td>
<td>3.53 ± 0.60</td>
<td>1.59 ± 0.86</td>
<td>7.98 ± 0.71</td>
<td>3.79 ± 1.17</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Caecal</td>
<td>3.27 ± 0.56</td>
<td>4.60 ± 0.95</td>
<td>0.79 ± 0.60</td>
<td>0.00 ± 0.82</td>
<td>5.23 ± 0.71</td>
<td>1.04 ± 1.17</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Acetic acid (mmol/kg substrate)</td>
<td>2.98 ± 0.11</td>
<td>4.65 ± 0.06</td>
<td>4.07 ± 0.11</td>
<td>4.91 ± 0.29</td>
<td>3.34 ± 0.15</td>
<td>3.37 ± 0.12</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Ileal</td>
<td>6.05 ± 0.00</td>
<td>6.92 ± 0.09</td>
<td>5.84 ± 0.12</td>
<td>5.67 ± 0.05</td>
<td>6.57 ± 0.05</td>
<td>6.57 ± 0.05</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Caecal</td>
<td>6.77 ± 0.01</td>
<td>6.77 ± 0.09</td>
<td>6.53 ± 0.12</td>
<td>6.51 ± 0.09</td>
<td>6.51 ± 0.09</td>
<td>6.51 ± 0.09</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Propionic acid (mmol/kg substrate)</td>
<td>0.21 ± 0.14</td>
<td>0.17 ± 0.14</td>
<td>0.11 ± 0.14</td>
<td>1.44 ± 1.39</td>
<td>0.00 ± 0.03</td>
<td>0.02 ± 0.04</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Ileal</td>
<td>0.00 ± 0.06</td>
<td>1.76 ± 0.12</td>
<td>155 ± 16</td>
<td>171 ± 11</td>
<td>164 ± 6</td>
<td>0.336 ± 0.02</td>
<td>–</td>
<td>0.336 ± 0.02</td>
</tr>
<tr>
<td>Caecal</td>
<td>0.54 ± 0.13</td>
<td>0.51 ± 0.13</td>
<td>0.50 ± 0.13</td>
<td>0.73 ± 0.27</td>
<td>0.64 ± 0.14</td>
<td>0.49 ± 0.12</td>
<td>0.018</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Butyric acid (mmol/kg substrate)</td>
<td>0.55 ± 0.08</td>
<td>0.66 ± 0.14</td>
<td>0.56 ± 0.14</td>
<td>0.19 ± 0.22</td>
<td>0.39 ± 0.14</td>
<td>0.69 ± 0.03</td>
<td>0.163</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Ileal</td>
<td>3.68 ± 0.23</td>
<td>2.96 ± 0.10</td>
<td>2.96 ± 0.09</td>
<td>2.58 ± 0.10</td>
<td>3.81 ± 0.10</td>
<td>4.02 ± 0.03</td>
<td>0.001</td>
<td>0.001</td>
</tr>
<tr>
<td>Caecal</td>
<td>3.65 ± 0.09</td>
<td>3.65 ± 0.08</td>
<td>3.65 ± 0.08</td>
<td>3.65 ± 0.08</td>
<td>3.65 ± 0.08</td>
<td>3.65 ± 0.08</td>
<td>0.163</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Isovaleric acid (mmol/kg substrate)</td>
<td>0.27 ± 0.16</td>
<td>1.04 ± 0.80</td>
<td>1.67 ± 1.18</td>
<td>1.67 ± 1.18</td>
<td>0.00 ± 0.02</td>
<td>0.45 ± 0.39</td>
<td>0.027</td>
<td>0.025</td>
</tr>
<tr>
<td>Ileal</td>
<td>2.48 ± 0.19</td>
<td>13.7 ± 0.83</td>
<td>6.93 ± 3.21</td>
<td>3.15 ± 1.24</td>
<td>2.44 ± 1.50</td>
<td>2.55 ± 0.82</td>
<td>0.027</td>
<td>0.025</td>
</tr>
<tr>
<td>Caecal</td>
<td>0.00 ± 0.05</td>
<td>0.05 ± 0.05</td>
<td>0.00 ± 0.05</td>
<td>0.00 ± 0.05</td>
<td>0.00 ± 0.05</td>
<td>0.00 ± 0.05</td>
<td>0.001</td>
<td>0.001</td>
</tr>
<tr>
<td>Valeric acid (mmol/kg substrate)</td>
<td>10.76 ± 12.0</td>
<td>13.4 ± 31.0</td>
<td>146 ± 12.4</td>
<td>157 ± 4.3</td>
<td>99 ± 7.5</td>
<td>172 ± 4.8</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Lactic acid (mmol/kg substrate)</td>
<td>56.3 ± 12.4</td>
<td>7.59 ± 9.9</td>
<td>28.2 ± 9.1</td>
<td>7.2 ± 7.2</td>
<td>105 ± 18.1</td>
<td>874 ± 106</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Ileal</td>
<td>492 ± 18.7</td>
<td>159 ± 18.7</td>
<td>156 ± 18.7</td>
<td>136 ± 18.7</td>
<td>207 ± 18.7</td>
<td>18.7 ± 18.7</td>
<td>–</td>
<td>18.7 ± 18.7</td>
</tr>
<tr>
<td>Caecal</td>
<td>997 ± 62.6</td>
<td>626 ± 62.6</td>
<td>1314 ± 62.6</td>
<td>626 ± 62.6</td>
<td>626 ± 62.6</td>
<td>626 ± 62.6</td>
<td>–</td>
<td>626 ± 62.6</td>
</tr>
</tbody>
</table>

AG, arabinoxylan; FOS, fructo-oligosaccharide; RS, resistant starch.
* Means for the caecal inoculum data differ from the ileal counterpart (P < 0.05).
† Different repeated statement for each organic acid were required to have similar Studentised residuals as described in the Statistical analysis section (e.g. for total organic acids, the best repeated statement had a common sse for each inoculum, while for lactic acid had a standard error for each individual substrate by inoculum combination). Means within each inoculum (ileal or caecal) without a common letter differ (P < 0.05).
‡ For formic, isobutyric and valeric acid production, there was no significant (P > 0.05) interaction between substrate and inoculum. Therefore, the interaction term was removed from the model.
§ A natural logarithm transformation of the raw data was required to achieve the model assumptions of normality and homoscedasticity. The values in parentheses represent the mean for each response variable after back transformation.
|| Negative values were found. They are explained by the artifact of correcting a small number for the material found in the blanks. In these instances, values are assumed to be zero (i.e. the organic acid was not produced).
For five out of seven of the substrates investigated, *in vitro* ileal OM fermentability exceeded 5%, demonstrating the potential importance of ileal fermentation. As more total OM enters the ileum compared with the caecum, the amount of fermented OM could be similar or even higher in the ileum compared with the caecum as reported elsewhere for pigs fed a human-type diet (66% more fermented OM in the ileum compared with the caecum) (7). Although this may be the case for some substrates, the generally higher production of organic acids in the caecal fermentation observed here demonstrates the importance of hindgut fermentation.

Although optimised *in vitro* ileal and caecal fermentation assays were used in this study (5,14), these methodologies do have limitations. For instance, they use fixed parameters (e.g. fermentation time) that in the animal may vary for different diets and substrates. The use of a pooled inoculum does not provide for animal to animal variation but does allow characterising the fermentability of different substrates. The use of purified substrates in the fermentation also assumes that the substrates are not fermented to some degree in anterior GIT regions. For example, in the present work, we used intact pectin for the *in vitro* caecal fermentation, but 26% of pectin was fermented *in vitro* in the ileum. Anterior fermentation may modify the structure of the substrates, thus affecting their fermentability. Although the present results should be regarded as indicative only, the methodologies do assist in developing an understanding of the fermentation of substrates in parts of the GIT. Based on our results, the ileum appears to be a site of substantial bacterial fermentation in the growing pig.

The large difference in OM fermentability seen among substrates during *in vitro* ileal fermentation may be explained by several factors such as: (i) a low bacterial phylogenetic diversity in the ileum and therefore a low diversity and number of enzymes able to cleave substrate bonds. For instance, and based on the ileal and caecal bacterial populations (online Supplementary Table S1), the predicted number of glycoside hydrolase families present in the ileum and caecum was sixty-five and seventy-three, respectively, for the pig fed a human-type diet (online Supplementary Table S2). It is important however to highlight that different glycoside hydrolase families could have the same activities (e.g. α-galactosidase, EC 3.2.1.7, could be expressed in families 18, 19, 23, 24, 25 and 73), although it is unknown if they may have the same capacity to hydrolyse glycoside bonds; (ii) structural differences in the substrates. For instance, the main difference between FOS and inulin is their chain length (on average five and twenty-five degrees of polymerisation, respectively) (27). This difference may have affected their *in vitro* ileal OM fermentability. Previous studies have shown that short fructans disappeared earlier than long fructans when fermented with either individual *Bifidobacterium* strains or a faecal human inoculum (9,27); (iii) monomeric composition of the substrate and specificity of the microbial glycoside hydrolases. For instance, pectin that had a greater *in vitro* ileal fermentation than the other substrates (e.g. inulin) has a greater diversity of monomeric units (galacturonic acids (65%), xylogalacturonan, apiogalacturonan, rhamnogalacturonan I and II, arabinose, galactose, rhamnose, xylose and fucose (20). *v.* inulin (2,1) bonds of fructose with a terminal α(1→2) linked D-glucose. Based on the monomeric unit composition, the predicted number of glycoside hydrolases available to break down pectin is 4.7-fold higher than for inulin; (iv) microbial preferences towards substrates, which could be explained by differences in other microbial enzymes (e.g. glycosyltransferases) and carbohydrate-binding modules (29). For instance, after 24 h of *in vitro* fermentation with faecal inocula, specific bacteria were attached only to insoluble wheat bran (e.g. *Roseburia faecis*), high amylose starch (e.g. *Ruminococcus bromii*) and a commercial porcine gastric mucin (e.g. *Bifidobacterium bifidum*). In contrast, other species attached to all substrates (e.g. *Escherichia rectale* and *Bifidobacterium longum*). (30)

In this study, fermentation was also investigated by determining the production of organic acids. Acetic, formic and lactic acids were found in quantifiable amounts for *in vitro* ileal fermentation. However, *in vitro* ileal productions of the other organic acids were low. Considering that organic acids produced in the ileum are absorbed (metabolised) in the ileum (27), it could be expected that they have similar beneficial effects both locally and systemically (e.g. intestinal tissue proliferation, GIT motility) to those absorbed in the large intestine (31,32). The relatively high production of acetic, formic and lactic acids may be explained by the high relative abundance of the *Streptococcus* genus (67% of the total 16S rRNA gene copies) in the ileal digesta (31,32). For instance, streptococci can rapidly ferment available carbohydrates to lactate. This is achieved through efficient uptake and conversion of sugars using the relatively simple and energy efficient phosphotransferase systems (22). Ileal digesta did not host members of the Bacteroidetes phylum (online Supplementary Fig. S2). Bacteria in the Bacteroidetes phylum possess a range of carbohydrate enzymes (e.g. α- and β-glucosidases) that allow them to utilise substrates at a competitive advantage over other members of the gut microbial consortium, resulting in the production of acetate, succinate and especially propionate (33). The lack of butyric acid production during *in vitro* ileal fermentation can also be attributed to the absence of bacterial families including Lachnospiraceae and Ruminococcaceae, which are key butyrate producers (34).
During in vitro ileal fermentation, the production of formic, acetic and lactic acids differed between substrates. For instance, FOS had the highest production of acetic acid, but the lowest production of lactic acid, while AG had the lowest production of acetic acid, but the highest production of formic acid. These differences could be explained by the same factors described above (e.g. chemical compositions of the fibre sources). Pectin, which had the highest in vitro OM fermentability, did not have the highest production of the analysed organic acids. Other intermediate microbial metabolites (e.g. pyruvic acid) not analysed in this study could have been produced during pectin fermentation in greater amounts before being converted to endpoint organic acids later in the fermentation (e.g. pyruvic acid can be metabolised to lactic acid).56

In this study, pigs fed a human-type diet were used as an animal model for the adult human due to their similarities in foregut digestion. The ileal fermentation observed here may also be important for the adult human. However, more work is needed to validate the pig as a model for ileal fermentation and to elucidate the physiological significance of ileal fermentation in humans. The results reported here give encouragement for this work to be undertaken.

The differences in OM fermentability between the substrates observed for in vitro ileal fermentation were reduced during in vitro caecal fermentation. This could be ascribed to some of the factors discussed above such as a greater amount and diversity of caecal microbial enzymes. Some caecal bacteria may have different pathways to produce enzymes, which are able to degrade both dietary and non-dietary materials,57,58 which explains why mucin was also fermented during in vitro caecal fermentation. It may also be related to the longer time of fermentation. Important differences in the in vitro caecal production of formic, acetic, propionic, butyric, isovaleric and lactic acids were observed. In general, FOS had the greatest production of acetic, propionic, butyric and valeric acids, while cellulose had the lowest. The low production of these organic acids for cellulose may be related to its low in vitro caecal fermentability due to the absence of cellulose-digesting enzymes in many of the GIT metagenome members, while the high levels of FOS and AG may be explained by several factors such as its high in vitro caecal fermentability and a lower number of pathways to produce organic acids as reviewed elsewhere.59

To demonstrate the quantitative significance of ileal fermentation in the pig itself, in vitro ileal fermentation was compared with in vitro caecal fermentation. The considerably greater in vitro caecal OM fermentability of FOS, inulin and mucin compared with their in vitro ileal counterparts in the pig could be attributed to both the longer fermentation time and greater amounts of glycoside hydrolases. For instance, in this study, the extent of caecal OM fermentability during 2 h of in vitro caecal fermentation for FOS (16%) and inulin (14%) was lower than that for their 24 h counterparts (33% and 32%, respectively). This is supported by the rate of SCFA production reported elsewhere for FOS and inulin.60 For FOS, the in vitro ileal OM fermentability was comparable to the in vitro caecal OM fermentability for 2 h. In contrast for inulin, the in vitro ileal OM fermentability was still lower. This lower fermentability could be related to a lower number of inulin degrading enzymes in the ileal digesta compared with caecal digesta of the pigs fed the human-type diet. In this study, the predicted number of endo-inulinase (EC 3.2.1.7) and exo-inulinase (EC 3.2.1.80) enzymes was 2-fold higher in the caecal digesta compared with ileal digesta. Similarly, the predicted number of inulin lyases (DFA-I-forming (EC 4.2.2.17) and (DFA-III-forming (EC 4.2.2.18)) in caecal digesta was 9-fold greater. For the remaining substrates (AG, cellulose, pectin and starch), there was no difference in their in vitro ileal and caecal OM fermentability. Despite similarities in OM fermentability between the ileum and caecum for AG, pectin and resistant starch, the total organic acid production and energy derived from the organic acids were higher during in vitro caecal fermentation for all substrates. A correlation (r = 0.52; P ≤ 0.05; n = 21) was found between OM fermentability and organic acid energy production for in vitro caecal fermentation, but not for in vitro ileal production. The lack of a correlation for in vitro ileal fermentation could be attributed to the different microbial composition. Different microbial populations may have different energy requirements and different efficiencies of fermentation.60–62 Based on energy and OM disappearance in the hindgut of pigs fed diets containing different levels of fibre, and on energy lost as methane and heat, assuming that they are lost in the hindgut,63 8.7 kJ of energy is expected to be produced per gram of fermented ileal digesta. This value is similar to 8-3 kJ/g dietary fibre fermented suggested by the European Commission.64 In this study, the energy derived from organic acids ranged between 0.6 and 11.7 kJ/g substrate fermented (in vitro) in the ileum (except for cellulose), and between 2.5 and 12.8 kJ/g substrate fermented (in vitro) in the caecum. This suggests that important amounts of energy from the microbial fermentation of fibre substrates can be produced in the ileum. Further, in some instances, organic acids other than the ones determined in this study may have contributed to the energy arising from fermentation.

In conclusion, the extent of ileal fermentation of both dietary and non-dietary substrates appears to be quantitatively significant in the growing pig adapted to a human-type diet, and such ileal fermentation may also be important in the adult human. The extent of ileal fermentation, however, depends upon the substrate. This could be attributed to factors such as structural differences of the substrates and available glycoside microbial hydrolases. The energy from fibre fermentation in the ileum is estimated to be between 0.6 and 11.7 kJ/g substrate fermented. Some substrates appear to be fermented to the same extent as in the caecum.

Acknowledgements

The authors acknowledge the help of Edward de Haas and Megan Cognée during the study.

This study was funded by the Centre of Research Excellence Fund from the Tertiary Education Commission and the New Zealand Ministry of Education.

C. A. M. and P. J. M. were responsible for planning the study. P. B. was responsible for the microbial analysis. C. A. M. was responsible for conducting the experiments, did the statistical
analysis and prepared the first draft of the manuscript which was revised by P. J. M. and P. B. All authors read and approved the final manuscript. The authors declare that there are no conflicts of interest.

Supplementary material

For supplementary materials referred to in this article, please visit https://doi.org/10.1017/S0007114520000542

References


35. Sakamoto M & Benno Y (2006) Reclassification of Bacteroides distasonis, Bacteroides goldsteinii and Bacteroides merdae as In vivo ileal v. caecal fermentations


