Effects of dietary protein supply, weaning age and experimental enterotoxigenic *Escherichia coli* infection on newly weaned pigs: health

I. J. Wellock¹, P. D. Fortomaris², J. G. M. Houdijk¹ and I. Kyriazakis¹,³

¹Animal Nutrition and Health Department, Scottish Agricultural College, West Mains Road, Edinburgh, EH9 3JG, UK; ²Department of Animal Production, School of Veterinary Medicine, Aristotle University of Thessaloniki, 54124 Thessaloniki, Greece; ³Faculty of Veterinary Medicine, University of Thessaly, PO Box 43100 Karditsa, Greece

(Received 31 January 2008; Accepted 4 February 2008)

Weaning is often associated with post-weaning colibacillosis (PWC), caused by enterotoxigenic *Escherichia coli* (ETEC). The objective was to investigate the effects of manipulating dietary protein supply and increasing weaning age on enteric health and ETEC shedding of newly weaned pigs exposed to an experimental ETEC challenge. The experiment consisted of a complete $2 \times 2 \times 2$ factorial combination of weaning age (4 v. 6 weeks), dietary protein content (H, 230 g crude protein (CP)/kg v. L, 130 g CP/kg) and experimental ETEC challenge (+ v. −); all foods were free from in-feed antimicrobial growth promoters (AGP). An additional four treatments were added to allow the effect of protein source (DSMP, dried skimmed milk powder v. SOYA, soybean meal) and AGP inclusion (yes v. no) to be investigated in challenged pigs of both weaning ages. On day 3 post-weaning challenged pigs were administered per os with $10^9$ cfu ETEC O149. A subset of pigs was euthanased on days 0 and 6 post weaning to assess enteric health and small intestine morphology. Both weaning age and dietary protein content affected the consequences of ETEC challenge. ETEC excretion persisted longer in the 4-week-weaned pigs than those weaned at 6 weeks. Although not significant, the numbers of ETEC shed in the faeces post infection (days 4 to 14) were higher on the H than L diet, especially in the 4-week-weaned pigs ($P = 0.093$). Lowering CP level led to significantly firmer faeces post challenge (days 3 to 6) and decreased colonic digesta pH. Protein level had no effect on small intestine villous heights or crypt depths. There was no significant effect of protein source on ETEC excretion or enteric health. Results suggest that increasing weaning age and decreasing the level of dietary protein, especially in earlier weaned pigs, may help to maintain enteric health and minimise the effects of PWC.

Keywords: health, pigs, post-weaning colibacillosis, protein, weaning age

Introduction

Weaning is a critical stage of pig production and is associated with profound changes in the structure and function of the gastrointestinal tract (Pluske *et al.*, 1997). Furthermore, weaning introduces a number of stress factors that may negatively affect the immune function and intestinal microflora of pigs (Barnett *et al.*, 1989; Montagne *et al.*, 2004). These disturbances increase the risk of enteric disorders, particularly post-weaning colibacillosis (PWC), which is caused primarily by the proliferation of enterotoxigenic *Escherichia coli* (ETEC).

Traditionally the risk of enteric disorders in newly weaned pigs has been mitigated through the use of in-feed antimicrobial growth promoters (AGP). However, their use has been recently withdrawn within the EU, at least partly in recognition of the growing problem of antibiotic resistance of pathogens such as ETEC (Barton, 1999). Consequently, the weaned pig will be at greater risk from PWC and there is a growing need to find alternative, non-pharmaceutical, strategies to combat such infections. Manipulation of dietary protein supply, in order to decrease protein availability to ETEC in the distal small intestine, and the consequent production of harmful fermentation by-products, such as amines, is one way in which this could be achieved (Prohászka and Baron, 1980; Nyachoti *et al.*, 2006; Wellock *et al.*, 2006). Increasing weaning age may also decrease sensitivity to PWC, due to the establishment of the regular intake of solid feed, resulting in a more mature and stabilised gut morphology and physiology reducing the colonisation of ETEC (Nabuurs *et al.*, 1996; Pluske *et al.*, 1997).
The objective of the current experiment was to investigate the effects of post-weaning dietary protein supply and weaning age on ETEC shedding and enteric health of newly weaned pigs exposed to an experimental ETEC challenge. An analysis of the data regarding the effect of experimental treatment on post-weaning performance is reported separately (Wellock et al., 2008). It was hypothesised that reducing protein availability in the distal small intestine, through reducing dietary protein content and/or substitution of existing ingredients, mainly soya, with the more digestible dried skimmed milk powder (DSMP), and increasing weaning age would reduce ETEC proliferation and improve the enteric health of the newly weaned pig.

Material and methods

Animals and housing

A total of 104 pigs (Large white × Landrace) of both sexes weaned at either 4 (28.2 ± 1.20 days) or 6 (40.8 ± 1.31 days) weeks of age (mean ± s.d.) and weighing 8.0 (± 0.98) and 13.0 (±1.94) kg, respectively, were used in the experiment. Pigs were derived from 10 litters, with five litters weaned at 4 weeks and five litters weaned at 6 weeks of age. At weaning (day 0), pigs were removed from the sow and individually housed in one of six rooms, each containing eight pens per room. The pens (2 × 1 m) had a 0.2 m deep transparent plastic partitioning along their length to enable visual contact between adjacent pens. Each pen was bedded with sawdust and equipped with a feeder and a nipple drinker. The environmental temperature was maintained at 25.6 ± 1.20 and 22.6 ± 0.60°C for the 4- and 6-week-weaned pigs, respectively, for the first 3 days after weaning and then decreased by 2°C/week for the remainder of the experiment. Fresh feed and water was available ad libitum throughout the experiment.

Diet and feeding

All pigs had access to a standard creep (16.0 MJ DE/kg and 230 g crude protein (CP)/kg) feed free from AGP during the last 14 days of sucking to allow animals to have experience of solid feed before weaning. At weaning, pigs were given ad libitum access to one of four experimental diets for 2 weeks: a high protein (H, 230 g/kg CP) and a low protein (L, 130 g/kg CP) both with DSMP (H-DSMP and L-DSMP, respectively); a high protein where DSMP was replaced with soya bean meal (H-SOYA) and H-DSMP with ZnO (3100 mg/kg), Cu (170 mg/kg) and avilamycin (Maxus G200, Elanco Animal Health Ltd, UK; 40 mg/kg) (H-AGP). For further details of the experimental diets, see Wellock et al. (2008). On day 14 post-weaning, a standard grower ration was fed to all animals until 10 weeks of age.

Experimental design

The experiment was conducted over two rounds with one weaning age represented in each round. It consisted of a complete 2 × 2 × 2 factorial combination of weaning age (4 or 6 weeks), dietary protein content (H or L) and experimental ETEC challenge (infected, + or sham, −). An additional four treatments were added to allow the effect of protein source (DSMP or SOYA) and AGP inclusion (yes or no) to be investigated in challenged pigs of both weaning ages. This resulted in six treatments per weaning age. The experimental design and numbers of pigs in each of the 12 treatment groups are shown in Table 1. Pigs were assigned to the six treatment groups per weaning age, taking into account body weight (BW) and sex, as far as possible, with littermates equally divided across treatments. Treatments were assigned to pens with each of the six treatments per weaning age being represented in each of the six experimental rooms. It was ensured that all uninfected animals were housed in pens adjacent to other uninfected animals in order to minimise the risk of contamination between pens.

Four pigs per treatment on day 6 and an additional six pigs per weaning age on day 0 were euthanased to assess enteric health. Performance of the remaining animals was monitored until 10 weeks of age (see Wellock et al., 2008). The Animal Experiments Committee of the Scottish Agricultural College

<table>
<thead>
<tr>
<th>Treatment code</th>
<th>Number of animals</th>
<th>Weaning age (weeks)</th>
<th>CP (g/kg)</th>
<th>CP source</th>
<th>AGP</th>
<th>Infection (ETEC O149)</th>
</tr>
</thead>
<tbody>
<tr>
<td>4-L-DSMP−</td>
<td>8</td>
<td>4</td>
<td>130 (L)</td>
<td>DSMP</td>
<td>−</td>
<td>−</td>
</tr>
<tr>
<td>4-H-DSMP−</td>
<td>8</td>
<td>4</td>
<td>230 (H)</td>
<td>DSMP</td>
<td>−</td>
<td>−</td>
</tr>
<tr>
<td>4-L-DSMP+</td>
<td>8</td>
<td>4</td>
<td>130 (L)</td>
<td>DSMP</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>4-H-DSMP+</td>
<td>8</td>
<td>4</td>
<td>230 (H)</td>
<td>DSMP</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>4-L-SOYA+</td>
<td>8</td>
<td>4</td>
<td>230 (H)</td>
<td>SOYA</td>
<td>−</td>
<td>+</td>
</tr>
<tr>
<td>4-H-AGP+</td>
<td>8</td>
<td>4</td>
<td>230 (H)</td>
<td>DSMP</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>6-L-DSMP−</td>
<td>7</td>
<td>6</td>
<td>130 (L)</td>
<td>DSMP</td>
<td>−</td>
<td>−</td>
</tr>
<tr>
<td>6-H-DSMP−</td>
<td>7</td>
<td>6</td>
<td>230 (H)</td>
<td>DSMP</td>
<td>−</td>
<td>−</td>
</tr>
<tr>
<td>6-L-DSMP+</td>
<td>8</td>
<td>6</td>
<td>130 (L)</td>
<td>DSMP</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>6-H-DSMP+</td>
<td>7</td>
<td>6</td>
<td>230 (H)</td>
<td>DSMP</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>6-L-SOYA+</td>
<td>8</td>
<td>6</td>
<td>230 (H)</td>
<td>SOYA</td>
<td>−</td>
<td>+</td>
</tr>
<tr>
<td>6-H-AGP+</td>
<td>7</td>
<td>6</td>
<td>230 (H)</td>
<td>DSMP</td>
<td>+</td>
<td>+</td>
</tr>
</tbody>
</table>

DSMP = dried skimmed milk powder; ETEC = enterotoxigenic Escherichia coli; SOYA = soybean meal; AGP = antimicrobial growth promoters; ZnO = 3100 mg/kg, CuSO₄ = 170 mg/kg and avilamycin (Maxus G200) = 40 mg/kg.
approved the protocol used in the study (ED AE 20/2003) for consistency with UK Home Office regulations (PPL 60/3205).

**Experimental infection**

The pigs were infected with 10^9 colony-forming units (cfu) of ETEC (E. coli O149 239/03) suspended in 10 ml phosphate buffer solution (PBS) on day 3 post weaning to induce sub-clinical PWC, according to the methodology of Houdijk et al. (2005). The infective dose was given on this day in order to ensure that pigs were eating a sufficient quantity of solid feed before being exposed to ETEC. The ETEC used were derived from clinical cases of PWC (Veterinary Laboratories Agency, Surrey, UK) and had been characterised as having the required virulence factors to induce PWC (Adhesion factors K91, K88 (F4)). The infection was administered per os, using a stomach tube with an additional 10 ml PBS for rinsing. PBS was used in order to increase bacterial survival in the stomach and to aid safe transfer of the inoculum into the small intestine. Non-infected animals were given 20 ml of PBS as a sham infection.

**Sampling collection and measurements**

*Health status and faecal sampling.* Individual faecal scored (FS), cleanliness scores (CS) and health scores (HS) were recorded each morning throughout the experiment, and every 2 h for a period of 24 h post infection using a subjective four-point scale (1 to 4) (see Wellock et al., 2006 for further details). Briefly, an increasing score represents increasing fluidity of faeces, increasing faecal contamination and deteriorating health for FS, CS and HS, respectively. Trained individuals with no prior knowledge of the treatment allocation recorded all scores. Fresh faecal samples from all animals were collected directly from the rectum on days 0, 3 (immediately before infection), 4, 6, 9 and 14 to assess ETEC shedding.

*Post mortem sampling procedures and measurements.* The six pigs per weaning age taken directly from the sow (day 0) and four pigs per treatment on day 6 were euthanased by intracardiac injection of Euthatal (Merial Animal Health Ltd, Harlow, Essex, UK; 0.7 ml/kg) and exsanguinated. The abdomen was then opened from the sternum to the pubis, and the gastrointestinal tract removed. Upon removal the gastrointestinal tract was divided into five sections: stomach (ST), small intestine (SI), caecum (CM), proximal colon (PC) and distal colon (DC). Pieces of intestinal tissue, approximately 5 cm in length, were immediately taken at three different locations of the SI: a site located about 5 cm after the gastric pylorus (proximal small intestine), at the mid point between the gastric pylorus and ileocaecal junction (mid small intestine) and at a site 5 cm proximal to the ileocaecal junction (distal small intestine). These samples were tied at either end with cotton, filled by injecting formalin and conserved in formalin until analysis of villous height and crypt depth (see below). The pH of the digesta from the ileum (IL), CM, PC and DC was measured upon collection by inserting the electrode of a portable pH meter (Testo 230; Testo Ltd, Alton, UK) into the collected sample. Digesta samples from the IL and PC were assessed for the lactobacilli to coliform ratio (L:C) and ETEC concentration (IL only).

**Microbiology**

*Inoculum.* A bead of the E. coli O149 239/03 strain was taken from storage at −80°C, reconstituted onto a sheep blood agar plate (E&O Laboratories Ltd, UK) and incubated overnight at 37°C. A number of formed colonies were removed from the plate, seeded into nutrient broth (Oxoid Ltd, Cambridge, UK) and incubated at 37°C for 24 h while being shaken at 200 rpm. Bacteria were harvested by centrifugation, twice washed with PBS and re-suspended at a concentration of 1 × 10^8 cfu/ml. The number of cfu’s per ml was checked before being used for oral inoculation of the pigs through standard enumeration techniques.

*Detection and enumeration of ETEC O149.* Approximately 1 g of each sample (IL and faecal) was serially diluted to 10.^−9^ in sterile PBS, and 100 μl aliquots were plated on sheep blood agar (E&O Laboratories Ltd, Stirlingshire, UK). The number of ETEC colonies was counted after 24 h incubation (39°C) under aerobic conditions. Randomly picked colonies were identified and confirmed as the infective strain by slide agglutination with a specific antiserum (K91 and K88; Mast Group Ltd, Bootle, UK).

*Lactobacillus and coliform counts.* Approximately 1 g of each sample (IL and PC) was serially diluted to 10.^−8^ in maximum recovery diluent, and 100 μl aliquots were plated in MacConkey agar (E&O Laboratories Ltd, UK) for coliforms and in MRS agar (E&O Laboratories Ltd, UK) for lactobacilli. Coliform colonies were counted after 24 h incubation (39°C) under aerobic conditions and lactobacilli colonies were counted after 48 h incubation (39°C) under anaerobic conditions (Reid and Hillman, 1999).

**Villus height and crypt depth measurement**

For histological analysis, tissue samples of the proximal, mid and distal small intestine were cut in order to form a ring-shaped length of tissue. These sections were then dehydrated before being embedded in paraffin wax. Sections were cut (4 μm) and stained with haematoxylin and eosin. From the stained sections, villous height and crypt depths were determined manually. One slide per piglet was used for each of the sites and the average value taken for a minimum of four villi and crypts per slide. The villous/crypt ratio was calculated as the villous height divided by the crypt depth.

**Calculations and statistical analysis**

The data were analysed as three separate factorial designs; a 2 × 2 × 2 analysis of weaning age (4 v. 6 weeks), protein content (H v. L) and infection (+ v. −); a 2 × 2 analysis of weaning age (4 v. 6 weeks) and protein source (DSMP v. SOYA); and a 2 × 2 analysis of weaning age (4 v. 6 weeks) and AGP inclusion (yes v. no). In addition, a one-way
ANOVA was performed to determine the effect of weaning age on indicators of enteric health on day 0. The effects of the main factors, along with any interactions, in each of the three analyses were performed by restricted maximum likelihood (REML) to account for unbalanced data caused by missing pigs in four of the treatments (see Table 1) and the removal of one pig from the trial (see below). BW at 4 weeks of age was used as a covariate in all analyses. The individual pig was used as the experimental unit and litter as a random factor. Effects of sex and experimental room were found not to be significant and not included in the model. All statistical analyses were performed by Genstat 5 for Windows (2001) (release 4.2, service pack 2; Lawes Agricultural Trust, Rothamsted, Hertfordshire, UK). Individual FS, CS and HS were averaged over the four time periods (days 0 to 3, 3 to 6, 0 to 14 and 14 to 28) and ETEC, lactobacillus and coliform counts were log10 transformed before analysis and calculation of L : C. Because the effect of weaning age was investigated in each of the three separate statistical analyses, three P values for each of the factors were determined. As these were not always identical, the largest P value is presented in order to prevent over-interpretation of the results.

Results

Faecal, cleanliness and health scores
None of the animals suffered from clinical PWC or had to be removed from the experiment due to illness. However, one animal in the treatment group 6-H-AGP+ was not infected due to poor feed intake (<50 g/day), and was consequently removed from the experiment.

Table 2 shows the effect of weaning age, protein content and infection on the mean FS and CS throughout the trial period. There were no significant effects of infection on FS, CS or HS, although infected 4-week-weaned pigs tended to have increased FS values post infection compared with their non-infected counterparts, especially in the first 24-h post-infection period (P = 0.092). Throughout the experiment, the 6-week-weaned pigs had higher FS and CS than their 4-week-weaned counterparts. However, this was significant only in the pre-infection period (days 0 to 3). There was a significant weaning age by infection interaction for FS and CS over the period immediately post infection (days 3 to 6). This was because infection increased FS and CS of pigs weaned at 4 weeks of age but not those weaned at 6 weeks when compared with their non-infected counterparts.

There was a significant improvement in both FS and CS in the 3- to 6-day period with a decrease in the CP content of the diet. The mean ± s.e. FS and CS for this period were 1.70 ± 0.09 and 1.30 ± 0.05 for the H-DSMP diet and 1.45 ± 0.07 and 1.14 ± 0.03 for the L-DSMP diet, respectively. There was also a significant weaning age by CP level interaction in the 3- to 6-day period for both FS and CS, with pigs weaned at 6 weeks showing a larger increase in FS and CS with an increase in CP than those weaned at 4 weeks. There was no effect of protein source on FS, CS or HS. There was a significant (P < 0.05) decrease in FS in the 3- to 6-day and 0- to 14-day periods with the inclusion of AGP (day 0 to 14; H-DSMP = 1.41 v. H-AGP = 1.06; P = 0.044). The inclusion of AGP also decreased CS and HS throughout the experimental period, but this was not significant.

ETEC excretion
Neither the infection strain nor any other ETEC was detected in the animals prior to the challenge, nor in any of the uninfected groups post infection. The number of ETEC colonies recovered from the faeces throughout the

<table>
<thead>
<tr>
<th>Treatments</th>
<th>Days 0 to 3</th>
<th>Days 3 to 6</th>
<th>Days 0 to 14</th>
<th>Days 14 to 28</th>
</tr>
</thead>
<tbody>
<tr>
<td>4-L-DSMP−</td>
<td>1.00</td>
<td>1.20</td>
<td>1.02</td>
<td>1.24</td>
</tr>
<tr>
<td>4-H-DSMP−</td>
<td>1.00</td>
<td>1.17</td>
<td>1.36</td>
<td>1.29</td>
</tr>
<tr>
<td>4-L-DSMP+</td>
<td>1.00</td>
<td>1.48</td>
<td>1.35</td>
<td>1.37</td>
</tr>
<tr>
<td>4-H-DSMP+</td>
<td>1.00</td>
<td>1.70</td>
<td>1.31</td>
<td>1.28</td>
</tr>
<tr>
<td>6-L-DSMP−</td>
<td>1.48</td>
<td>1.83</td>
<td>1.46</td>
<td>1.31</td>
</tr>
<tr>
<td>6-H-DSMP−</td>
<td>1.81</td>
<td>2.36</td>
<td>1.57</td>
<td>1.28</td>
</tr>
<tr>
<td>6-L-DSMP+</td>
<td>1.46</td>
<td>1.31</td>
<td>1.35</td>
<td>1.32</td>
</tr>
<tr>
<td>6-H-DSMP+</td>
<td>1.79</td>
<td>1.65</td>
<td>1.60</td>
<td>1.14</td>
</tr>
<tr>
<td>s.e.d.†</td>
<td>0.276</td>
<td>0.299</td>
<td>0.314</td>
<td>0.262</td>
</tr>
<tr>
<td>Response</td>
<td>W**</td>
<td>CP*</td>
<td>W × CP*</td>
<td>W × I*</td>
</tr>
<tr>
<td></td>
<td>W**</td>
<td>CP**</td>
<td>W × CP*</td>
<td>W × I*</td>
</tr>
<tr>
<td></td>
<td>W × I**</td>
<td>W × CP*</td>
<td>W × I*</td>
<td></td>
</tr>
</tbody>
</table>

4 = 4 week weaned; 6 = 6 week weaned; W = Weaning age (6 v. 4 week); I = infection (− v. +); CP = crude protein content (H-DSMP v. L-DSMP); H = 230 g CP/kg; L = 130 g CP/kg; DSMP = dried skimmed milk powder; +/− represents presence of absence respectively of infection with enterotoxigenic Escherichia coli.

**P < 0.05, ***P < 0.01, ****P < 0.001.
experimental period for the eight experimentally challenged treatment groups is shown in Table 3. The number of ETEC shed was significantly \( P = 0.003 \) greater and persisted longer in the 4-week-weaned pigs than those weaned at 6 weeks (Figure 1). By day 9 (day 6 post infection) no 6-week-weaned animals were excreting detectable numbers of ETEC \( \geq 1 \times 10^3 \text{cfu/g} \), whereas the average excretion of the infected 4-week-weaned animals was 3.87 \( \log_{10} \text{cfu/g} \). Detectable numbers of ETEC were not recovered from 4-week-weaned animals on day 14 (day 11 post infection). There was no significant effect of CP level on the number of ETEC shed in the faeces post infection, although numbers of ETEC shed tended to be higher on the H-DSMP diet compared with the L-DSMP diet, especially in the 4-week-weaned pigs (day 1 to 11 post infection; \( H = 4.55 \) vs. \( L = 3.78 \log_{10} \text{cfu/g}; P = 0.093 \) (see Figure 1). There was no effect of protein source on the number of ETEC shed (day 1 to 11 post infection; \( \text{SOYA} = 3.44 \) vs. \( \text{DSMP} = 3.09 \log_{10} \text{cfu/g}; P = 0.522 \), but there was a significant weaning age by protein source interaction for the average number of ETEC shed over days 1 to 3 post infection. While there was little difference in the number of ETEC shed between the protein sources in the 4-week-weaned pigs, the 6-week-weaned pigs on the H-SOYA diet excreted a larger number of ETEC than those on the corresponding H-DSMP diet (see Table 3). There were no significant effects of AGP inclusion on ETEC shedding, although the 4-week-weaned pigs on the H-AGP diet shed approximately 50% less than pigs fed the H-DSMP diet. There was a significant weaning age by AGP inclusion interaction on day 3 post infection and over the 1- to 11-day post-infection period. This arose from the 6-week-weaned animals on the H-AGP diet, which showed an increase in the number of ETEC shed compared with those on the H-DSMP diet.

Table 3 shows the effect of weaning age (4 v. 6 weeks) and crude protein level (H v. L) on faecal enterotoxigenic \( E. \) coli shedding post infection. Standard error bars showed only for 4-H and 6-H.

**Figure 1** Effect of weaning age (4 v. 6 week) and crude protein level (H v. L) on faecal enterotoxigenic \( E. \) coli shedding post infection. Standard error bars showed only for 4-H and 6-H.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>Site/day (day post infection shown in brackets)</th>
<th>Faeces</th>
<th>Ileum</th>
</tr>
</thead>
<tbody>
<tr>
<td>Treatment</td>
<td>3 (0)</td>
<td>4 (1)</td>
<td>6 (3)</td>
</tr>
<tr>
<td>4-L-DSMP+</td>
<td>0.00</td>
<td>6.12</td>
<td>4.72</td>
</tr>
<tr>
<td>4-H-DSMP+</td>
<td>0.00</td>
<td>6.46</td>
<td>5.83</td>
</tr>
<tr>
<td>4-H-SOYA+</td>
<td>0.00</td>
<td>6.68</td>
<td>5.54</td>
</tr>
<tr>
<td>4-H-AGP+</td>
<td>0.00</td>
<td>5.61</td>
<td>3.73</td>
</tr>
<tr>
<td>6-L-DSMP+</td>
<td>0.00</td>
<td>5.27</td>
<td>2.38</td>
</tr>
<tr>
<td>6-H-DSMP+</td>
<td>0.00</td>
<td>4.16</td>
<td>0.38</td>
</tr>
<tr>
<td>6-H-SOYA+</td>
<td>0.00</td>
<td>6.70</td>
<td>1.90</td>
</tr>
<tr>
<td>6-H-AGP+</td>
<td>0.00</td>
<td>4.84</td>
<td>1.84</td>
</tr>
<tr>
<td>s.e.d. ( ^* )</td>
<td>0.000</td>
<td>0.915</td>
<td>0.766</td>
</tr>
<tr>
<td>Response</td>
<td>W**</td>
<td>W**</td>
<td>W**</td>
</tr>
<tr>
<td></td>
<td>W ( \times ) CP*</td>
<td>W ( \times ) PS*</td>
<td>W ( \times ) A**</td>
</tr>
</tbody>
</table>

4 = 4 week weaned; 6 = 6 week weaned; \( W = \) Weaning age (6 v. 4 week); \( CP = \) crude protein content (H-DSMP v. L-DSMP); \( PS = \) protein source (H-DSMP v. H-SOYA+); \( A = \) antimicrobials (H-AGP v. H-DSMP+); \( H = 230 \) g CP/kg; \( L = 130 \) g CP/kg; \( \text{DSMP = dried skimmed milk powder}; \text{SOYA = soybean meal}; \text{AGP = antimicrobial growth promoters}; \) + = infection with enterotoxigenic \( E. \) coli.

\( ^* \)s.e.d. for the \( W \times CP \) interaction.

\( ^* P < 0.05, ** P < 0.01, *** P < 0.001. \)
tract on days 0 and 6. Pigs weaned at 6 weeks had a significantly lower pH in the IL and higher pH in the ST on day 0 than pigs weaned at 4 weeks. The L : C ratio was higher in the 4 than 6-week-weaned pigs on day 0 in the PC (P = 0.025), mainly due to a decrease in the number of coliforms (P = 0.021).

There was no effect of infection on the pH of the different gastrointestinal sites, although the pH of the IL and PC contents tended to be higher in challenged pigs than their non-infected counterparts. For example, the mean ± s.e. pH of the IL digesta of the challenged and non-challenged pigs was 7.12 ± 0.16 and 6.64 ± 0.17 (P = 0.070), respectively. The pH of the IL and PC contents were higher in pigs weaned at 4 weeks compared with those weaned at 6 weeks (7.13 ± 0.63, P = 0.078) and (6.35 ± 0.56, P < 0.001), respectively, while tending to be lower in the ST (2.65 ± 3.45, P = 0.099). There was a tendency for an increase in pH with an increase in CP content but this was only significant in the PC (H-DSMP = 6.22 v. L-DSMP = 5.80, P = 0.012). There was a weaning age by infection interaction on the pH of ST contents, with infection decreasing pH of the 4-week-weaned pigs and increasing pH in the 6-week-weaned pigs.

There was no effect of weaning age or infection on the L : C ratio of samples taken from the IL and PC at slaughter on day 6, although non-infected animals had a higher ratio than infected animals. Increasing weaning age led to a significant increase in the number of lactobacilli in both the IL and PC (4 week = 7.47 v. 6 week = 8.25 log10 cfu/g, P = 0.002). There was no significant effect of CP level on the L : C ratio. A decrease in CP level led to a decrease (P = 0.021) in the number of lactobacilli in the PC, 8.44 ± 0.11 and 8.02 ± 0.12 log10 cfu/g for the H-DSMP and L-DSMP diets, respectively. There was no effect of CP level on coliform numbers.

There was no effect of protein source on gastrointestinal pH, but there was a significant increase in the pH of the IL (H-DSMP = 7.18 v. H-AGP = 7.57; P = 0.044) and PC (H-DSMP = 6.14 v. H-AGP = 6.67; P = 0.006) digesta with the inclusion of AGP. There was no significant effect of protein source or AGP on the L : C ratio, although there was significant reduction in the number of coliforms (H-DSMP = 7.19 v. H-AGP = 5.85; P < 0.001) and lactobacilli (H-DSMP = 8.03 v. H-AGP = 6.38; P < 0.001) with the inclusion of AGP.

Villous height and crypt depth
Table 5 shows the effect of weaning age, protein level and experimental infection on the villous heights and crypt depths for three sites of the small intestine on days 0 and 6. In all, 4-week-weaned pigs had significantly increased villous heights, decreased crypt depths and increased villous height to crypt depth ratio (V : C) compared with 6-week-weaned pigs on day 0, although this was not significant for villous height at the proximal site (P = 0.077) and V : C at the distal site of the small intestine (P = 0.062).

There was no effect of infection on villous height, crypt depth or the V : C ratio at any of the sites of the small intestine. There was a significant effect of weaning age on villous height and crypt depth in most of the sites, with 6-week-weaned animals having larger villous and deeper crypts than those weaned at 4 weeks (see Table 5). This led
to smaller V:C ratios in all 3 sites for the 6-week-weaned pigs, although this was not significant \( (P = 0.301) \). There was a significant weaning age by infection interaction for mean crypt depth and mean V:C ratio. Infected 4-week-weaned pigs had lower crypt depth and higher V:C ratio than non-infected counterparts while the reverse was true for the 6-week-weaned pigs.

There was no effect of protein level on small intestine morphology. Pigs fed the H-SOYA diet tended to have increased villous height and V:C ratio than those on the H-DSMP diet, although this was significant only for the villous height at the proximal site \( (\text{H-SOYA} = 449 \, \mu\text{m v. H-DSMP} = 403 \, \mu\text{m}; P < 0.001) \). The only significant effect of AGP on villous height or crypt depths was on crypt depth in the distal small intestine, with pigs fed AGP having shallower crypts than those on the H-DSMP diet \( (\text{H-DSMP} = 276 \, \mu\text{m v. H-AGP} = 235 \, \mu\text{m}; P = 0.021) \).

### Discussion

The oral challenge of pigs with pathogenic *Escherichia coli* has been widely used as a model of PWC (Sarmiento et al., 1988; Nollet et al., 1999; Madec et al., 2000; Van Dijk et al., 2002; Kiers et al., 2003; Montagne et al., 2004). The experimental model used here produced prolonged faecal ETEC excretion in all challenged pigs post infection. No pigs died and/or showed signs of clinical PWC, although feed intake and growth rate were temporarily impaired (see Wellock et al., 2008). This suggests that the aim of inducing sub-clinical PWC was successfully achieved. There was a rapid initial increase in ETEC shedding in the faeces of challenged pigs immediately post infection followed by a gradual decrease. Others have reported the same pattern over a similar time period (Nabuurs et al., 1993; Kiers et al., 2003).

Experimental infection had a greater impact and persisted longer in the 4-week-weaned pigs than those weaned at 6 weeks, with greater numbers of ETEC being shed immediately post infection and for a longer period of time. All challenged 6-week-weaned pigs in the trial stopped shedding detectable numbers of ETEC by day 6 post infection, while 50% of the challenged 4-week-weaned pigs were still shedding more than 10^5 cfu/g faeces. One reason as to why 4-week-weaned pigs may have been affected to a greater extent than the 6-week-weaned pigs may be due to the greater gut maturity and better immunological status of the older pigs, allowing them to better combat the infection (Vega-Loópez et al., 1995). Alternatively, it may be a consequence of using the same experimental dose for both weaning ages, rather than making this weight dependent. However, as there was no significant correlation between BW on day of infection and average daily gain as a percentage BW over the immediate post-infection period (Nabuurs et al., 1996; Vente-Spreeuwenberg et al., 2003), which may be associated with increased occurrence of diarrhoea (Nabuurs et al., 1993). These changes in gut morphology are primarily related to the low feed intake post-weaning (Kelly et al., 1991; Pluske et al., 1996), being lowest around day 4 before...
reversing to pre-weaning values between days 11 and 14 (Nabuurs et al., 1993). Older pigs therefore may be expected to show less of a decrease in villous height due to a greater feed intake. This was observed here with pigs weaned at 4 weeks of age demonstrating a much larger decrease in villous height and V:C ratio from days 0 to 6 than pigs weaned at 6 weeks of age. The fact that infection had little impact on villi atrophy, a common feature associated with clinical PWC (Kenworthy, 1976; Nabuurs et al., 1993; McDonald et al., 2001), supports the view that sub-clinical rather than clinical PWC was induced. Infection had a greater detrimental effect on the enteric health, as indicated by a higher pH, lower L:C ratio of the PC contents and lower villous heights at all three sites of the small intestine, of pigs weaned at 4 weeks than those weaned at 6 weeks of age.

Infection had a rapid and short-lived effect on FS and CS. Within 12 h post infection, 15% of infected pigs were displaying FS of 2.5 or higher from an initial score of 1. By day 6 post infection, however, most scores had returned to normal. Similar results were seen by Madec et al. (2000) who reported transient diarrhoea in 50% of inoculated pigs soon after the challenge, accompanied by signs of depression and low weight gain. No diarrhoea was observed here in the non-challenged 4-week-weaned pigs, although this was not the case for those weaned at 6 weeks, who had increased FS compared with the 4-week-weaned pigs, especially in the first 3 days of the experiment. This may have been related to their achieved feed intake immediately post weaning. While it has been suggested that a high creep intake before weaning favours immune tolerance (Hampson, 1987), others have reported an increased incidence of diarrhoea post weaning with an increased consumption of dry food prior to weaning (Barnett et al., 1989).

It was hypothesised that decreasing the amount of protein in the diet would decrease the amount of substrate available in the distal small intestine and hence decrease the rate of ETEC proliferation. This was previously observed in pigs offered the H-DSMP and H-SOYA diets. Similar results have been reported by Montagne et al. (2004) who found no significant effect on the proliferation of ETEC when animal protein was replaced with plant protein in a rice-based diet. Several studies have shown a decrease in villous height in newly weaned pigs fed a diet with a high level of soya instead of DSMP (Li et al., 1991; Makinde et al., 1996), while others have failed to do so (McCracken et al., 1999; Jiang et al., 2000). The fact that pigs fed the DSMP diets ate significantly more than those fed the SOYA diets (see Wellock et al., 2008), and that DSMP constituted only 25% of total protein in the DSMP diets, may explain why there was no significant effect of protein source on ETEC shedding or measures of enteric health.

The inclusion of AGP in the diet decreased the numbers of ETEC shed in the faeces in the 4-week-weaned pigs by approximately 50% but not in the 6-week-weaned pigs. Only 25% pigs of the challenged 4-week-weaned pigs on the AGP diet were excreting any ETEC on day 6 post infection compared with 75% of those on the equivalent non-AGP diet (H-DSMP). The addition of AGP also led to a decrease in FS, suggesting a decreased severity of PWC (McDonald et al., 2001) and risk of spread from one piglet to another when in a group situation. AGP inclusion had no effect on enteric health in terms of the gastrointestinal pH or L:C ratio, but as reported elsewhere (Broom et al., 2003; Wellock et al., 2006) there was a significant decrease in total lactobacilli and coliform numbers.

Conclusion

The results suggest that, in the absence of in-feed AGP, increasing weaning age and decreasing dietary protein supply may help to maintain enteric health and minimise the risk and of spread of PWC by reducing ETEC proliferation and faecal ETEC shedding. Decreasing the dietary CP content may be especially important in earlier weaned animals, and particularly in environments that are challenging to the health of pigs.

Acknowledgements

This research was financially supported by ABN Ltd, Frank Wright Ltd, Home-Grown Cereals Authority, Meat and Livestock
Wellock, Fortomaris, Houdijk and Kyriazakis

Commission/British Pig Executive, Primary Diets Ltd and Provimi Ltd with match funding from Defra, through the Sustainable Livestock Production LINK programme. The authors would like to acknowledge Paul Toplis of Primary Diets Ltd, UK, for the formulation and manufacture of the experimental diets, Biomathematics and Statistics Scotland for help with the statistical analysis, Terry McHale, Dave Anderson, Lesley Deans, Kostantios Zaralis, Richard Slade and Siobhan Carroll for technical assistance and Kevin Hillman for the L : C analysis.

References


