

## Effect of maternal supplementation with seaweed extracts on growth performance and aspects of gastrointestinal health of newly weaned piglets after challenge with enterotoxigenic *Escherichia coli* K88

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(Submitted 2 May 2014 – Final revision received 3 September 2014 – Accepted 7 September 2014 – First published online 27 October 2014)

### Abstract

In the present study, a 2 × 2 factorial arrangement was conducted to investigate the effect of maternal supplementation with seaweed extracts (–SWE *v.* +SWE, *n* 20) from day 83 of gestation until weaning (day 28) on post-weaning (PW) growth performance, faecal score, faecal enterotoxigenic *Escherichia coli* (ETEC) toxin quantification, intestinal histology and cytokine mRNA of unchallenged and ETEC-challenged pigs. Pigs were ETEC challenged on day 9 PW. There was a maternal treatment × challenge (SWE × ETEC) interaction effect on growth performance and faecal score (*P* < 0.05). Pigs from SWE-supplemented sows and ETEC-challenged (SE) had higher average daily gain (ADG) during 0–13 d PW and reduced faecal score during 0–72 h post-challenge than those from basal-fed sows and ETEC-challenged (BE) (*P* < 0.05). However, there was no difference between unchallenged pigs from the SWE-supplemented sows (SC) and basal-fed sows (BC) (*P* > 0.10). Pigs from the SWE-supplemented sows had reduced heat-labile enterotoxin gene copy numbers than those from the basal-fed sows (*P* < 0.05). Maternal SWE supplementation increased the villus height in the ileum of pigs (*P* < 0.05). There was a SWE × ETEC interaction effect (*P* < 0.05) on *IL-6* mRNA and a SWE × gastrointestinal (GI) region interaction effect (*P* < 0.05) on transforming growth factor-β1 (*TGF-β1*) and *TNF-α* mRNA. *IL-6* mRNA was down-regulated in SC pigs than BC pigs (*P* < 0.05). However, there was no difference in *IL-6* mRNA between SE and BE pigs. The mRNA of *TGF-β1* and *TNF-α* was down-regulated in the colon of pigs from the SWE-supplemented sows compared with those from the basal-fed sows (*P* < 0.05). However, there was no difference in *TGF-β1* and *TNF-α* mRNA in the ileum between the pigs from the SWE-supplemented sows and basal-fed sows. In conclusion, maternal SWE supplementation improves ADG and the aspects of GI health of weaned pigs following an ETEC challenge.

**Key words:** Enterotoxigenic *Escherichia coli* K88 challenge: Inflammatory cytokines: Pig performance: Seaweed extracts

Enterotoxigenic *Escherichia coli* (ETEC) K88 is a major cause of diarrhoea in neonates and recently in weaned pigs<sup>(1,2)</sup>. It results in reduced growth rate, increased morbidity and mortality, and great economic loss in pig production worldwide<sup>(3,4)</sup>. Post-weaning (PW) diarrhoea due to ETEC K88 generally occurs in pigs 3–10 d PW<sup>(5,6)</sup>. ETEC K88 colonises in the small intestine and releases specific enterotoxins that impair intestinal barrier function, which indirectly induces fluid losses<sup>(4–6)</sup>. Several factors, such as stress of weaning, lack of milk and dietary changes, contribute to the severity of the disease. Furthermore, weaning is often associated with undesirable morphological and physiological changes

in the gastrointestinal (GI) tract that is related to a reduced voluntary feed intake and, subsequently, susceptibility to intestinal health imbalance<sup>(7,8)</sup>. The ban on the inclusion of antibiotic growth promoters in weaning pig diets in Europe (European Community Regulation no. 1831/2003) has been associated with an increase in diarrhoea, weight loss and mortality due to *E. coli* in pigs PW<sup>(9)</sup>. Traditional measures to ameliorate or reduce PW disorders have centred on dietary manipulations PW<sup>(10–13)</sup>. However, recent research has indicated that maternal dietary supplementation with seaweed extracts (SWE) containing laminarin and fucoidan has beneficial effects on weaning-associated intestinal dysfunction

**Abbreviations:** ADFI, average daily feed intake; ADG, average daily gain; BC, pigs weaned from basal-fed sows and unchallenged; BE, pigs weaned from basal-fed sows and enterotoxigenic *Escherichia coli* K88-challenged on days 9 and 10 post-weaning; BW, body weight; cDNA, complementary DNA; CD, crypt depth; EAST1, enteroaggregative *Escherichia coli* heat-stable enterotoxin 1; ETEC, enterotoxigenic *Escherichia coli*; G:F, gain:feed ratio; GCN, gene copy number; GI, gastrointestinal; *IFN-γ*, interferon-γ; *LT*, heat-labile enterotoxin; qPCR, quantitative real-time PCR; PW, post-weaning; SAS, Statistical Analysis System; SC, pigs weaned from seaweed extract-supplemented sows and unchallenged; SE, pigs weaned from seaweed extract-supplemented sows and enterotoxigenic *Escherichia coli* K88-challenged on days 9 and 10 PW; *Stb*, heat-stable enterotoxin b; SWE, seaweed extracts; *TGF-β1*, transforming growth factor-β1; VH, villus height.

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and growth depression immediately after weaning<sup>(14)</sup>. Furthermore, pigs weaned from SWE-supplemented sows had a higher average daily gain (ADG) between days 0 and 21 PW compared with those weaned from untreated sows<sup>(14)</sup>.

Experiments on the effect of maternal dietary supplementation with SWE on growth performance and indicators of GI health of weaned piglets after challenge with ETEC K88 PW are limited. The objective of the present study was to investigate the effect of maternal supplementation with SWE from day 83 of gestation until weaning (day 28) on pig growth performance, faecal score, ETEC K88 toxin quantification, intestinal architecture and gene expression profiles of cytokines of both unchallenged and ETEC K88-challenged weaned piglets. The hypothesis of the present study is that maternal supplementation with SWE would enhance the growth performance and improve the aspects of GI health and immune status of weaned piglets, making them more immune competent to deal with the PW adversities, represented in the present experiment by an ETEC K88 challenge.

### Materials and methods

All procedures described in the present experiment were conducted under experimental license from the Irish Department of Health in accordance with the Cruelty to Animals Act 1876 and the European Communities (Amendments of the Cruelty to Animals Act, 1876) Regulations.

#### *Experimental design and animal diets – gestation and lactation periods*

A total of twenty cross-bred pregnant gilts (Large White × Landrace genetic lines, Hermitage) were assigned to one of two dietary treatments (*n* 10 sows per treatment): basal gestation/lactation diet (T1) and basal gestation/lactation diet supplemented with 10.0 g SWE/d (T2) from day 83 of gestation until weaning (day 28). The quantity of SWE (BioAtlantis Limited) used in the present study was based on previous work by Leonard *et al.*<sup>(14)</sup>. The SWE supplement (10.0 g/d) contained laminarin (1.0 g), fucoidan (0.8 g) and ash (8.2 g), and was extracted from a *Laminaria* spp. according to the procedure described by Lynch *et al.*<sup>(15)</sup>. The gestation diet was formulated to contain 140 g/kg of crude protein, 13.0 MJ/kg of digestible energy and 5.5 g/kg of total lysine. The lactation diet was formulated to contain 190 g/kg of crude protein, 14.5 MJ/kg of digestible energy and 10.0 g/kg of total lysine. All amino acids requirements were met relative to lysine<sup>(16)</sup>. The ingredients and chemical analysis of the experimental diets are presented in Table 1.

#### *Animals and management*

Before day 83 of gestation, the gilts were housed in groups of ten. From day 83 until day 106 of gestation, they were housed individually in crates (2.0 × 0.6 m) in the gestation house. In the farrowing house, the gilts were housed individually in farrowing pens (2.2 × 2.4 m). The gestation house and farrowing room temperature was maintained at 20°C throughout the

experiment. The experimental supplement (SWE) was top-dressed on the gestation and lactation diets each morning (09.00 hours) to ensure consumption. The dams received specific amounts of feed in the following quantities: 2.5 kg/d of gestation diet from day 83 until day 106 of gestation. They were fed 2 kg/d of lactation diet from day 107 of gestation until the day of farrowing, and then the feed supply was increased by 1.0 kg/d until day 3 post-farrowing and by 0.5 kg/d until day 6 post-farrowing. Afterwards, the sows were allowed semi-*ad libitum* consumption of the lactation diet, which was adjusted for each sow depending on daily intake. The sows were fed in three equal meals provided at 09.00, 13.00 and 17.00 hours. The sows had *ad libitum* access to drinking-water throughout the experimental period.

#### *Farrowing and piglet management*

All farrowings were supervised. At parturition, each piglet (meat-line boars × Large White × Landrace genetic lines) was individually weighed and tagged. Between 6 and 12 h after the birth of the last piglet, litter size was adjusted by cross-fostering piglets within dietary treatments to ensure that sows nursed a similar number of piglets (*n* 12 piglets/sow), and this was maintained throughout the suckling period. The individual piglet body weight (BW) was recorded at birth, on day 7 and at weaning, and ADG calculated from these data. Piglets received an intramuscular injection of Fe-dextran (Ferdex 100; Medion Farma Jaya) on day 7 after birth. No creep feed was offered to the piglets throughout the lactation period.

#### *Post-weaning period – experimental design and animal management*

At weaning, a total of forty male pigs (*n* 2 pigs/litter) with an average BW of 8.4 (SD 0.14) kg were selected. The selection was based on the average BW of the litter. The pigs were individually penned in a fully slatted pen (1.7 m × 1.2 m). The pigs were fed *ad libitum* with a standard diet for this age of pig until day 13 PW (Table 1). The diet was formulated to contain 210 g/kg of crude protein, 15.0 MJ/kg of digestible energy and 14.5 g/kg of ileal digestible lysine. Water was available from nipple drinkers at all times. Feed was available up to weighing of the pigs, and then weighed back for the purpose of calculating feed intake. Representative feed sample was taken on day 0 (day of weaning) for chemical analysis. The ingredient composition and chemical analysis of the PW diet are presented in Table 1.

House temperature was maintained at 30°C during the first 7 d and then reduced to 28°C until day 13 PW. On days 9 and 10 PW, twenty pigs (ten pigs weaned from the basal-fed sows and ten pigs weaned from the SWE-supplemented sows) were housed in unsanitised rooms and orally challenged with ETEC K88 to induce PW diarrhoea and the other twenty pigs were housed in sanitised rooms and received only PBS (Oxoid), resulting in four groups: pigs weaned from the basal-fed sows and unchallenged (control pigs, *n* 10; BC); pigs weaned from the basal-fed sows and ETEC k88-challenged on days 9 and 10 PW (*n* 10; BE); pigs



**Table 1.** Ingredients and chemical analysis of the experimental diets (g/kg, unless otherwise indicated)

	Gestation diet*	Lactation diet*	Post-weaning diet
<b>Ingredients</b>			
Wheat	343.8	318.8	380.0
Barley	300.0	300.0	233.5
Soyabean meal	67.0	182.0	170.0
Full-fat soyabean			120.0
Dried maize distillers' grains	60.0	100.0	
Soya hulls	70.0		
Beet pulp	100.0		
Whey powder			50.0
Soya oil	30.0	70.0	10.0
Vitamins and minerals†‡	3.0	2.5	3.0
Salt	3.0	5.0	3.0
Dicalcium phosphate	11.2	12.0	12.5
Limestone	12.0	12.0	11.0
Lys-HCl		2.0	4.0
D-L-Met		1.0	1.5
L-Thr		1.0	1.5
<b>Chemical analysis</b>			
DM	870.9	873.2	866.1
Crude protein (N × 6.25)	140.0	190.0	210.6
Gross energy (MJ/kg)	16.9	17.1	17.1
Ash	55.2	57.1	48.4
Neutral-detergent fibre	–	–	115.1
DE (MJ/kg)§	13.5	14.5	14.5
Lys§	5.5	10.0	14.5
Met and Cys§	3.3	6.0	8.4
Thr§	3.85	7.0	9.1
Trp§	1.0	1.8	2.5
Ca§	8.7	9.3	9.5
P§	5.0	5.2	6.1

DE, digestible energy.

\*T1, basal diet; T2, basal diet supplemented with 10.0 g seaweed extract/d.

 † Gilt/sow diet provided (per kg diet): 250 mg choline chloride; 140 mg Fe; 120 mg Zn as ZnO; 67 mg  $\alpha$ -tocopherol; 47 mg Mn as MnO; 25 mg Cu as CuSO<sub>4</sub>; 12 mg nicotinic acid; 10 mg pantothenic acid; 4 mg phytylmenaquinone; 2 mg riboflavin; 2 mg thiamin; 1.8 mg retinol; 0.6 mg iodine as calcium iodate on a calcium sulphate/calcium carbonate carrier; 0.3 mg Se as sodium selenite; 0.025 mg cholecalciferol; 0.01 mg cyanocobalamin; 0.015 mg pyridoxine.

 ‡ Weaner diet provided (per kg diet): 100 mg Zn as ZnO; 40 mg  $\alpha$ -tocopherol; 25 mg Mn as MnO; 25 mg Cu as CuSO<sub>4</sub>; 0.3 mg retinol; 0.2 mg iodine as calcium iodate on a calcium sulphate/calcium carbonate carrier; 0.3 mg Se as sodium selenite; 0.05 mg cholecalciferol.

 § Calculated from amino acid and DE values of ingredients<sup>(46)</sup>.

weaned from the SWE-supplemented sows and unchallenged ( $n$  10; SC); pigs weaned from the SWE-supplemented sows and ETEC k88-challenged on days 9 and 10 PW ( $n$  10; SE). Each treatment was housed in separated rooms, to avoid cross-contamination between the treatments. The unsanitised rooms were uncleaned following a previous batch of weaned pigs, while the sanitised rooms were cleaned and disinfected following the previous batch of weaned pigs. The pigs were weighed at weaning (day 0), day 9 (before the ETEC K88 challenge) and at the end of the experiment (day 13). The pigs were humanely killed on day 13 by lethal injection with pentobarbital sodium (Euthatal Solution, 200 mg/ml; Merial Animal Health Limited) at a rate of 0.71 ml/kg BW to the cranial vena cava, and the entire GI tract was immediately removed.

#### *Enterotoxigenic Escherichia coli K88 challenge – bacterial culture, oral challenge, health status and sample collection*

The ETEC K88 strain (Abbotstown, O149:K91:K88), obtained from Animal Health and Veterinary Laboratories Agency

(AHVLA, Addlestone, Surrey, UK), was grown in Luria-Bertani broth. The pathogen was derived from clinical cases of PW colibacillosis and characterised as having the required virulence factors to induce PW colibacillosis. The expanded culture of ETEC K88, approximately  $1 \times 10^8$  colony-forming units/ml, was further prepared for oral dosing as described by Zhang *et al.*<sup>(17)</sup>. The suspension was calculated on the basis of optical density established by serial dilution before bacterial cell count. Pigs were not tested for susceptibility to these adhesion factors, but were from a herd where persistent PW ETEC K88 shedding was recorded. At 08.00 hours on days 9 and 10 PW, the pigs ( $n$  20) were orally given a dose of 10 ml PBS containing approximately  $1 \times 10^8$  colony-forming units of ETEC K88 using a syringe attached to a polyethylene tube held in the back of the oral cavity. The unchallenged control pigs received 10 ml of sterilised PBS only. The bacterial solution and PBS was slowly dribbled into the pig's throat so that the swallowing reflex was triggered and passage of the inoculant into the lungs was minimised.

The pigs were checked for diarrhoea to evaluate their status before and after challenging with ETEC K88.

Severity of diarrhoea was characterised by using the following faecal consistency score system described by Pierce *et al.*<sup>(18)</sup>: (1) hard firm faeces; (2) slightly soft faeces; (3) soft, partially formed faeces; (4) loose, semi-liquid faeces (mild diarrhoea); (5) watery, mucous-like faeces (severe diarrhoea). Faecal scores were taken before the ETEC challenge (day 9, 0h) and every 12h after the challenge until day 13 (72h). Faecal consistency score was performed by two trained personnel with no prior knowledge of the sow's dietary treatment.

Fresh faecal samples were collected directly from the rectum of each pig at 0, 24 and 48h after ETEC K88 challenge and stored at  $-20^{\circ}\text{C}$  for quantification of ETEC K88 toxin genes (heat-labile enterotoxin – *LT*, heat-stable enterotoxin b – *STb* and enteroaggregative *E. coli* heat-stable enterotoxin 1 – *EAST1*). Diarrhoea caused by ETEC K88 is attributable to the action of one or more enterotoxins produced by the bacteria that have colonised the small intestine, such as *LT*, *STb* and *EAST1*<sup>(4,19)</sup>.

### Chemical analysis

The feed samples (gestation, lactation and PW diets) were analysed for N, DM, ash, gross energy and neutral-detergent fibre). The feed samples were milled through a hammer mill provided with a 1mm screen (Christy and Norris Hammer Mill; Christy Turner Limited). The nutrient contents (N, DM, ash, gross energy and neutral-detergent fibre) in the diets were determined as described by Heim *et al.*<sup>(13)</sup>. The total laminarin content of the SWE was determined using a Megazyme kit (Megazyme International Ireland Limited). Fucoïdan levels were determined using the method described by Usov *et al.*<sup>(20)</sup>.

### Collection of tissue samples

Immediately after slaughter, the entire digestive tract was removed by blunt dissection, and sections of the duodenum (10cm from the stomach), the jejunum (60cm from the stomach) and the ileum (15cm from the caecum) were excised and fixed in 10% phosphate-buffered formalin for villus height (VH) and crypt depth (CD) measurements. Ileum and colon tissues (second loop of the proximal colon) were excised, emptied by dissecting them along the mesentery and rinsed using sterile PBS (Oxoid). Tissue sections of  $1\text{cm}^2$ , which was stripped of the overlying smooth muscle, were cut from each tissue and stored in 15ml RNAlater<sup>®</sup> solution (Applied Biosystems) overnight at  $4^{\circ}\text{C}$ . RNAlater<sup>®</sup> was then removed before storing the samples at  $-80^{\circ}\text{C}$ .

### Microbiology – DNA extraction and quantitative PCR from faecal samples

Microbial genomic DNA was extracted from faecal samples using a QIAamp DNA stool kit (Qiagen) in accordance with the manufacturer's instructions. Quantity and quality of DNA was assessed using a Nanodrop ND1000 spectrophotometer (Thermo Scientific). Standard curves were generated as described by O'Shea *et al.*<sup>(21)</sup>. Briefly, genomic DNA from all

faecal samples was pooled and amplified through routine PCR using ETEC K88 toxin primers (*LT*, *STb* and *EAST1*). The ETEC K88 toxin primers are presented in Table 2. All primers were designed using Primer Express<sup>™</sup> software (Applied Biosystems) and synthesised by MWG Biotech. Serial dilutions of these amplicons served to generate standard curves using quantitative real-time PCR (qPCR – ABI 7500 Real-Time PCR System; Applied Biosystems Limited), allowing estimations of absolute quantification based on gene copy number (GCN)<sup>(22)</sup>. qPCR were performed in a final reaction volume of  $20\mu\text{l}$  containing  $3\mu\text{l}$  of template DNA,  $1\mu\text{l}$  of forward ( $100\text{pM}$ ) and  $1\mu\text{l}$  of reverse primers ( $100\text{pM}$ ),  $10\mu\text{l}$  SYBR Green PCR Master Mix (Applied Biosystems) and  $5\mu\text{l}$  nuclease-free water. The thermal cycling conditions involved an initial denaturation step at  $95^{\circ}\text{C}$  for 10min followed by forty cycles of  $95^{\circ}\text{C}$  for 15s and  $65^{\circ}\text{C}$  for 1min. Dissociation analyses of the qPCR product were performed to confirm the specificity of the resulting qPCR products. All samples were prepared in duplicate. The mean cycle threshold ( $C_t$ ) values of duplicates of each sample were used for calculations.

### Duodenal, jejunal and ileal morphology

The preserved segments (duodenum, jejunum and ileum) were prepared using standard paraffin-embedding techniques. The samples were sectioned at  $5\mu\text{m}$  thickness and stained with haematoxylin and eosin<sup>(23)</sup>. Measurements of fifteen well-orientated and intact villi and crypts were taken for each segment. The VH and the CD were measured as described by Heim *et al.*<sup>(13)</sup>. The results are expressed as mean VH or CD in  $\mu\text{m}$ . The VH:CD ratio was calculated.

### RNA extraction, complementary DNA synthesis and quantitative real-time PCR

RNA was extracted from approximately 50mg of tissue samples using the GenElute Mammalian Total RNA Miniprep Kit (Sigma-Aldrich Corporation), according to the manufacturer's instructions. Total RNA was quantified using  $1.5\mu\text{l}$  of total RNA on a NanoDrop Spectrophotometer ND1000 (Thermo Fisher Scientific, Inc.), and samples with a 260:280 ratio  $\geq 2.0$  were considered suitable for complementary DNA (cDNA) synthesis. Total RNA integrity (i.e. quality and quantity) was assessed by analysing  $1\mu\text{l}$  of total RNA using the Agilent 2100 Bioanalyser version A.02.12 (Agilent Technologies, Inc.) using RNA Nano LabChips<sup>®</sup> (Caliper Technologies Corporation). cDNA synthesis was performed using  $1\mu\text{g}$  of total RNA and oligo(dT)<sub>20</sub> primers in a final reaction volume of  $20\mu\text{l}$  using Superscript<sup>™</sup> III First-Strand synthesis kit (Invitrogen Life Technologies) following the manufacturer's instructions. The final reaction volume of  $20\mu\text{l}$  was then adjusted to  $250\mu\text{l}$  using nuclease-free water. All primers for the selected cytokine genes *IL-1*, *IL-2*, *IL-6*, *IL-8*, *IL-10*, *IL-12A* (*p35*), *IL-17*, *TNF- $\alpha$* , interferon- $\gamma$  (*IFN- $\gamma$* ), transforming growth factor- $\beta$ 1 (*TGF- $\beta$ 1*), forkhead box P3 (*FOXP3*) were designed using Primer Express<sup>™</sup> (PE Applied Biosystems) and synthesised by MWG Biotech. Primer sequences are



**Table 2.** Porcine oligonucleotide primers used for real-time PCR

Genes	Accession no.	Primer (5' → 3')	Product length (bp)
<i>LT*</i>	.	F: GCTACAAATGCCTATGCATCTACACA R: CATGCTCCAGCAGTACCATCTCTAAC	275
<i>STb†</i>	.	F: ACGGCGTTACTATCCTGTCTATGTGC R: TTGGTCTCGGTCAGATATGTGATTCT	125
<i>EAST1†</i>	.	F: TGCCATCAACACAGTATATCC R: GCGAGTGACGGCTTTGT	109
<i>IL-1</i>	NM_214029.1	F: CAGCCAACGGGAAGATTCTG R: AATGGCTTCCAGGTCGTCAT	77
<i>IL-2</i>	NM_213861.1	F: TAACCTCAACTCCTGCCACAATG R: GTGCACCGTTTGCCATGA	87
<i>IL-6</i>	NM_214399.1	F: GACAAAGCCACCACCCTAA R: CTCGTTCTGTGACTGCAGCTTATC	69
<i>IL-8</i>	NM_213867.1	F: TGCACTTACTCTTGCCAGAAGT R: CAAACTGGCTGTTGCCTTCTT	82
<i>IL-10</i>	NM_214041.1	F: GCCTTCGGCCCAAGTAA R: AGAGACCCGGTCAGCAACAA	71
<i>IL-12A (p35)</i>	NM_213993.1	F: CGTGCCCTCGGGCAATTATA R: CGCAGGTGAGGTCGCTAGTT	68
<i>IL-17</i>	NM_001005729.1	F: CAAGCGGTGGCGTTTTGCCT R: GTCTCCGTCGGGGATGGGCT	57
<i>IFN-γ</i>	NM_213948.1	F: TCTAACCTAAGAAAGCGGAAGAGAA R: TTGCAGGCAGGATGACAATTA	81
<i>TGF-β1</i>	NM_214015.1	F: AGGGCTACCATGCCAATTTCT R: CGGGTTGTGCTGGTTGTACA	101
<i>TNF-α</i>	NM_214022.1	F: TGGCCCTTGAGCATCA R: CGGGCTTATCTGAGGTTTGAGA	68
<i>FOXP3</i>	NM_001128438.1	F: GTGGTGCACTCTCTGGAACAAC R: AGGTGGGCTGCATAGCA	68
<i>PPIA</i>	NM_214353.1	F: CGGGTCCTGGCATCTTGT R: TGGCAGTGCAAATGAAAACTG	75
<i>BM</i>	NM_213978.1	F: CGGAAAGCCAAATTACCTGAAC R: TCTCCCGGTTTTTCAGCAAAT	83

*LT*, heat-labile enterotoxin; F, forward; R, reverse; *STb*, heat-stable enterotoxin b; *EAST1*, enteroaggregative *Escherichia coli* heat-stable enterotoxin 1; *IFN-γ*, interferon-γ; *TGF-β1*, transforming growth factor-β; *FOXP3*, forkhead box P3; *PPIA*, peptidylprolyl isomerase A; *BM*, β2-microglobulin.

\* Zhang *et al.*<sup>(47)</sup>.

† Metzler-Zebeli *et al.*<sup>(48)</sup>.

presented in Table 2. A panel of pro- and anti-inflammatory cytokines chosen was selected based on previous work by Walsh *et al.*<sup>(12)</sup> and Leonard *et al.*<sup>(24)</sup>. The efficiencies of all primer sets were established using a semi-log curve of quantity *v.* control, of two-fold serial dilutions of cDNA as reported previously by Smith *et al.*<sup>(25)</sup>. The following two porcine reference genes were used as described previously by Ryan *et al.*<sup>(26)</sup>: β2-microglobulin (*BM*) and peptidylprolyl isomerase A (*PPIA*). qPCR was then carried out on cDNA using the ABI PRISM 7500 Fast Sequence Detection System for ninety-six-well plates (Applied Biosystems). All samples were prepared in duplicate using the SYBR Green Fast PCR Master Mix (Applied Biosystems), cDNA as the template and specific primers for the genes selected. For each reaction, 5 μl cDNA, 1.2 μl forward and reverse primer mix (300 nM), 3.8 μl nuclease-free water and 10 μl Fast SYBR Green PCR Master Mix (PE Applied Biosystems) were added and made up to a final volume of 20 μl. The two-step PCR programme was as follows: 95°C for 10 min for one cycle, followed by 95°C for 15 s and 60°C for 1 min for forty cycles.

The mean *C<sub>t</sub>* values of duplicates of each sample were used for calculations. Normalised relative quantities were obtained using qbase PLUS software (Biogazelle).

### Statistical analysis

All data were analysed using the general linear model procedure of Statistical Analysis System (SAS; SAS Institute, Inc.)<sup>(27)</sup>. All data were initially checked for normality using the univariate procedure in SAS<sup>(27)</sup>. For pig performance (BW, ADG, average daily feed intake (ADFI) and gain:feed ratio (G:F)), faecal score, ETEC K88 toxin gene expression and cytokine gene expression, the data were analysed by repeated-measures analysis using the PROC MIXED procedure of SAS<sup>(28)</sup>. The model included the effects of sow dietary treatment, ETEC K88 challenge and time, and the associated two-way and three-way interactions. Gene copy estimates of ETEC K88 toxins were log-transformed before statistical analysis. The statistical model used for cytokine gene expression included both GI sites (ileum *v.* colon), sow dietary treatment and ETEC K88 challenge and the associated two-way and three-way interactions. Small intestinal morphology data were analysed as a 2 × 2 factorial arrangement, with the pig as the experimental unit. The statistical model used included the effects of sow dietary treatment and ETEC K88 challenge and the associated interaction. Contrast statements were used to compare (1) basal-fed sows *v.* SWE-supplemented sows (lactation effect), (2) non-challenge *v.* ETEC K88

**Table 3.** Effect of sow dietary treatment on weaned pig performance following an enterotoxigenic *Escherichia coli* (ETEC) K88 challenge\* (Least-squares mean values with their standard errors)

Sow SWE Pig ETEC	BC		BE		SC		SE		SEM		P	
	No	No	No	Yes	Yes	No	Yes	Yes	Yes	SWE	ETEC	SWE × ETEC
	Days 0-9	Days 9-13	Time	SWE × ETEC × time								
n	10	10	10	10	10	10	10	10	10	0.28		
BW day 0 (kg)	8.4	8.3	8.6	8.4	8.4	8.4	8.4	8.4	8.4	0.28		
Growth performance												
BW (kg)	9.4	10.6	9.8	9.2	9.8	10.7	9.8	10.6	0.28	0.018	0.053	0.257
ADG (kg/d)	0.11	0.30	0.13	0.12	0.15	0.22	0.15	0.20	0.03	0.019	0.018	0.343
ADFI (kg/d)	0.18	0.37	0.21	0.24	0.17	0.32	0.17	0.29	0.01	<0.001	0.014	0.049
G:F (kg/kg)	0.60	0.82	0.64	0.56	0.88	0.68	0.88	0.70	0.15	0.180	0.054	0.349

BC, pigs weaned from the basal-fed sows and unchallenged; BE, pigs weaned from the basal-fed sows and ETEC K88-challenged on days 9 and 10 post-weaning (PW); SC, pigs weaned from the seaweed extracts (SWE)-supplemented sows and unchallenged; SE, pigs weaned from the SWE-supplemented sows and ETEC K88-challenged on days 9 and 10 PW; BW, body weight; ADG, average daily gain; ADFI, average daily feed intake; G:F, gain:feed ratio.  
\* ETEC K88 oral challenge: days 9 and 10 PW.

challenge PW effect and (3) the interaction between the lactation effect and the PW effect. The sow data for the lactation period were analysed as a complete randomised block design, with the sow as the experimental unit. The pdiff function of SAS was used to separate means.

All data presented in the tables are expressed as least-square means with their standard errors of the mean. The probability level that denotes significance is  $P < 0.05$ .

**Results**

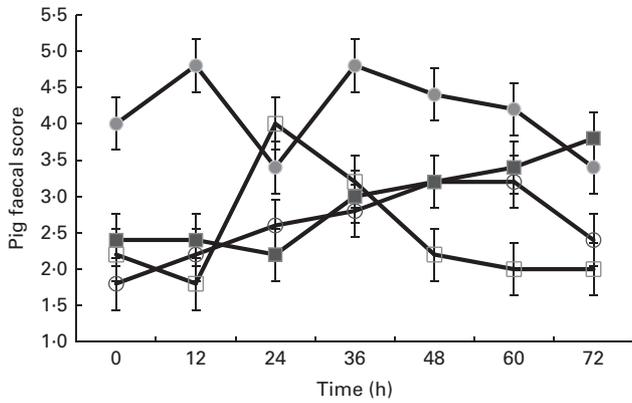
*Lactation period*

Gilts offered the SWE extracts had a longer gestation period than basal-fed gilts (114.5 v. 113.5 d, SEM = 0.27,  $P < 0.05$ ). Litter size (14.4 (SD 0.72) piglets) and number of live-born piglets (12.9 (SD 0.62)) were not influenced by sow dietary treatments ( $P > 0.10$ ). Sow dietary treatment had no effect on piglet BW at birth (1.76 (SD 0.66) kg), on day 7 after birth (2.70 (SD 0.09) kg) and at weaning (8.33 (SD 0.22) kg), and had no effect on ADG from birth to day 7 of lactation (0.189 (SD 0.010) kg/d) and from birth to weaning (0.240 (SD 0.069) kg/d) ( $P > 0.10$ ). Piglets suckling SWE-supplemented sows had higher ADG compared with those suckling basal-fed sows (0.249 v. 0.230 kg/d; SEM = 0.064;  $P < 0.05$ ) from day 8 of lactation to weaning.

*Post-weaning period*

**Growth performance.** The effect of sow dietary treatment and ETEC K88 challenge on pig growth performance PW is presented in Table 3. No difference was recorded in pig BW at weaning (day 0) among the dietary treatments ( $P > 0.10$ ). There was a sow dietary treatment × ETEC K88 challenge (SWE × ETEC) interaction effect on pig BW (days 9 and 13), ADG (days 0-9 and 9-13) and G:F (days 0-9) ( $P < 0.05$ ). The BE pigs had a decreased BW, ADG and G:F compared with the SE pigs ( $P < 0.05$ ). However, there was no difference in BW, ADG and G:F between the BC and SC pigs ( $P > 0.10$ ). There was a SWE × ETEC × time interaction effect on pig ADFI ( $P < 0.05$ ). The BE pigs had a decreased ADFI compared with BC pigs during 0-9 and 9-13 d ( $P < 0.05$ ). However, there was no difference in ADFI between the SC and SE pigs during 9-13 d ( $P > 0.10$ ), and the SE pigs had decreased ADFI compared with the SC pigs during 0-9 d ( $P < 0.05$ ).

**Faecal score.** The effect of sow dietary treatment and ETEC K88 challenge on pig faecal score from 0 h (immediately before the ETEC K88 challenge) to 72 h post-challenge is presented in Fig. 1. There was a SWE × ETEC × time interaction effect on pig faecal score from 0 to 72 h post-challenge ( $P < 0.05$ ). The BE pigs had higher faecal score at 0, 12, 36, 48, 60 and 72 h post-challenge compared with the SE pigs ( $P < 0.05$ ). However, there was no difference in faecal score at 0, 12 and 36 h post-challenge between the BC and SC pigs ( $P > 0.10$ ); at 24 h, the BC pigs had lower faecal score compared with the SC pigs, and at 48, 60 and 72 h post-challenge, the BC pigs had higher faecal score compared with the SC pigs ( $P < 0.05$ ).



**Fig. 1.** Differences in pig faecal score over time at 0 (immediately before the enterotoxigenic *Escherichia coli* (ETEC) K88 challenge), 12, 24, 36, 48, 60 and 72 h post-challenge. Values are means, with their standard errors represented by vertical bars. BC, pigs weaned from basal-fed sows and unchallenged (-SWE -ETEC K88; ○); BE, pigs weaned from basal-fed sows and ETEC K88-challenged on days 9 and 10 post-weaning (PW) (-SWE +ETEC K88; ●); SC, pigs weaned from SWE-supplemented sows and unchallenged (+SWE -ETEC K88; □); SE, pigs weaned from SWE-supplemented sows and ETEC K88-challenged on days 9 and 10 PW (+SWE +ETEC K88; ■). There was an effect detected for the SWE × ETEC × time interaction ( $P < 0.001$ ).

**Gene expression of enterotoxigenic *Escherichia coli* K88 toxins in the faeces.** The effect of sow dietary treatment and ETEC K88 challenge on *EAST1*, *LT* and *STb* log GCN/g faeces at 0, 24 and 48 h post-challenge is presented in Table 4. There was a time effect on *LT*, *EAST1* and *STb* log GCN/g faeces ( $P < 0.05$ ). At 24 h post-challenge, *LT* and *STb* log GCN/g faeces were decreased compared with the faecal collection time point at 0 and 48 h post-challenge ( $P < 0.05$ ). In contrast, *EAST1* log GCN/g faeces was increased at 24 h compared with the 0 and 48 h time points ( $P < 0.05$ ). The ETEC K88 challenge increased *EAST1* and *STb* log GCN/g faeces ( $P < 0.05$ ). Pigs weaned from the SWE-supplemented sows had reduced *LT* log GCN/g faeces compared with those weaned from the basal-fed sows ( $P < 0.05$ ).

**Small-intestinal morphology.** The effect of sow dietary treatment and ETEC K88 challenge on VH, CD and the VH:CD ratio in the duodenum, jejunum and ileum of pigs on day 13 PW is shown in Table 5. Pigs challenged with ETEC K88 tended to have a reduced VH:CD ratio in the duodenum ( $P = 0.064$ ) and deeper CD in the jejunum ( $P < 0.05$ ) compared with unchallenged pigs. Pigs weaned from the SWE-supplemented sows had longer VH and higher VH:CD

ratio in the ileum compared with those weaned from the basal-fed sows ( $P < 0.05$ ).

**Cytokine gene expression.** The effect of sow dietary treatment and ETEC K88 challenge on the expression of pro- and anti-inflammatory cytokines in the ileum and in the colon tissues is presented in Table 6. There was a SWE × ETEC interaction effect on the gene expression of the cytokine *IL-6* ( $P < 0.05$ ). The gene expression of *IL-6* was down-regulated in the ileum and colon tissues of SC pigs compared with BC pigs ( $P < 0.05$ ). However, there was no difference in the gene expression of *IL-6* in the ileum and colon tissues between the SE and BE pigs ( $P > 0.10$ ). There was an ETEC K88 × GI region interaction effect on the gene expression of the cytokines *IL-6*, *IL-10*, *IL-12A (p35)* and *TGF-β1* ( $P < 0.05$ ). The gene expression of *IL-10* and *IL-12A (p35)* was up-regulated in the ileum of ETEC K88-challenged pigs compared with their expression in the ileum of unchallenged pigs ( $P < 0.05$ ). However, there was no difference in the gene expression of *IL-10* and *IL-12A (p35)* in the colon between the ETEC K88-challenged and unchallenged pigs ( $P > 0.10$ ). The gene expression of *IL-6* and *TGF-β1* was down-regulated in the colon of ETEC K88-challenged pigs compared with their expression in the colon of unchallenged pigs ( $P < 0.05$ ). However, there was no difference in the gene expression of *IL-6* and *TGF-β1* in the ileum between the ETEC K88-challenged and unchallenged pigs ( $P > 0.10$ ). There was a SWE × GI region interaction effect on the gene expression of the cytokines *IL-8*, *TGF-β1* and *TNF-α* ( $P < 0.05$ ). The gene expression of *IL-8* was up-regulated in the ileum of pigs weaned from the SWE-supplemented sows compared with its expression in the ileum of pigs weaned from the basal-fed sows ( $P < 0.05$ ). However, there was no difference in the gene expression of *IL-8* in the colon between the pigs weaned from the SWE-supplemented sows and those weaned from the basal-fed sows ( $P > 0.10$ ). The gene expression of *TGF-β1* and *TNF-α* was down-regulated in the colon of pigs weaned from the SWE-supplemented sows compared with their expression in the colon of pigs weaned from the basal-fed sows ( $P < 0.05$ ). However, there was no difference in the gene expression of *TGF-β1* and *TNF-α* in the ileum between the pigs weaned from the SWE-supplemented sows and those weaned from the basal-fed sows ( $P > 0.10$ ). The gene expression of *IFN-γ* tended to be up-regulated in the ileum and colon tissues of pigs weaned from the SWE-supplemented sows compared with those weaned from the basal-fed sows ( $P = 0.09$ ).

**Table 4.** Effect of sow dietary treatment on enterotoxigenic *Escherichia coli* (ETEC) K88 toxin log gene copy number/g faeces following an ETEC K88 challenge\*

(Least-squares mean values with their standard errors,  $n$  10 pigs/treatment group)

Items	SWE		ETEC		Time (h)			SEM	$P$ †		
	Yes	No	Yes	No	0	24	48		SWE	ETEC	Time
<i>LT</i>	4.78	5.72	5.52	4.98	6.31	2.39	7.04	0.31	0.022	0.190	< 0.001
<i>STb</i>	6.48	6.50	6.81	6.18	6.86	5.89	6.72	0.19	0.944	0.016	0.005
<i>EAST1</i>	6.44	6.47	6.66	6.24	6.13	6.86	6.36	0.12	0.854	0.009	< 0.001

SWE, seaweed extracts; *LT*, heat-labile enterotoxin; *STb*, heat-stable enterotoxin b; *EAST1*, enteroaggregative *E. coli* heat-stable enterotoxin 1.

\* ETEC K88 oral challenge: days 9 and 10 post-weaning.

† There was no effect detected for the SWE × ETEC interaction ( $P > 0.10$ ) or the SWE × ETEC × time interaction ( $P > 0.10$ ).

**Table 5.** Effect of sow dietary treatment and enterotoxigenic *Escherichia coli* (ETEC) K88 challenge\* on small-intestinal morphology of pigs on day 13 post-weaning

(Least-squares mean values with their standard errors, *n* 10 pigs/treatment group)

Items	SWE		ETEC		SEM	<i>P</i> (effect)†	
	No	Yes	No	Yes		SWE	ETEC
<b>Duodenum</b>							
VH (μm)	150	155	164	141	11.2	0.730	0.149
CD (μm)	129	133	128	134	5.8	0.705	0.529
VH:CD	1.2	1.2	1.3	1.1	0.1	0.985	0.064
<b>Jejunum</b>							
VH (μm)	112	116	108	120	9.3	0.784	0.378
CD (μm)	111	116	107	120	4.5	0.377	0.054
VH:CD	1.0	1.0	1.0	1.0	0.1	0.858	0.828
<b>Ileum</b>							
VH (μm)	80	115	102	93	9.0	0.008	0.474
CD (μm)	109	106	103	113	6.5	0.752	0.288
VH:CD	0.8	1.2	1.1	0.9	0.1	0.025	0.326

SWE, seaweed extracts; VH, villus height; CD, crypt depth.

\*ETEC K88 oral challenge: days 9 and 10 post-weaning.

†There was no effect detected for the SWE × K88 interaction ( $P > 0.10$ ).

The gene expression of *IL-10* was up-regulated in the ileum and colon tissues of pigs challenged with ETEC K88 compared with unchallenged pigs ( $P < 0.05$ ). No effect was observed on the gene expression of the cytokines *FOXP3*, *IL-1*, *IL-17* and *IL-2* in the ileum and colon tissues ( $P > 0.10$ ).

## Discussion

The objective of the present experiment was to evaluate the effects of maternal dietary supplementation with SWE from day 83 of gestation until weaning on weaned piglet growth performance, faecal score, faecal ETEC K88 toxin quantification, intestinal histology and gene expression profiles of pro- and anti-inflammatory cytokines following an ETEC K88 challenge on days 9 and 10 PW. It was hypothesised that maternal supplementation with SWE would improve the growth performance and indicators of GI health of weaned pigs after an ETEC K88 challenge PW. The positive response observed in pigs weaned from SWE-supplemented sows on growth performance, faecal score, intestinal morphology and immunological response PW compared with those weaned from basal-fed sows supports the hypothesis.

### Lactation period

The SWE supplement (10.0 g/d) contained laminarin (1.0 g), fucoidan (0.8 g) and ash (8.2 g). It is speculated that this level of ash in the SWE supplement would have no impact on sow performance as it made up only a very small proportion of total ash intake. The present data indicate that SWE supplementation from day 83 of gestation increased the gestation length of the gilt. Their gestation length was prolonged by 1 d compared with the basal-fed gilts. In the swine species, the gestation length is an average of 114 d, with 85% of farrowing concentrating between 114 and 116 d of gestation<sup>(29)</sup>. However, this increase in gestation length

did not influence piglet BW at birth. There was also no effect of maternal dietary treatment on piglet BW on day 7 of age and at weaning. Chau *et al.*<sup>(30)</sup> observed no effect of sow dietary supplementation with yeast-derived β-glucans in the growth rate of piglets. Similarly Leonard *et al.*<sup>(24)</sup> also reported no effect of sow supplementation with a similar SWE from day 107 of gestation to weaning (26 d) on piglet growth during the lactation period.

### Post-weaning

The SE pigs had higher BW, ADG, ADFI and G:F compared with the BE pigs. However, there was no difference in BW, ADG, ADFI and G:F between the BC and SC pigs. The SE pigs had a lower faecal score at 0, 12, 36, 48, 60 and 72 h post-challenge compared with the BE pigs. However, there was no difference in faecal score at 0, 12 and 36 h post-challenge between the BC and SC pigs. At 48, 60 and 72 h post-challenge, the SC pigs had a lower faecal score compared with the BC pigs. The lack of a negative effect of ETEC K88 challenge on the growth performance and faecal score of pigs weaned from the SWE-supplemented sows could be due to a number of reasons. First, it may be attributed to the immunomodulatory effects of SWE in enhancing cellular and humoral immune function, as well as suppressing the *E. coli* population<sup>(24)</sup>. Leonard *et al.*<sup>(24)</sup> demonstrated that sow supplementation with SWE reduced colonic *E. coli* numbers in piglets at weaning. A suppressed colonic *E. coli* population may ultimately alleviate the incidence and severity of PW diarrhoea<sup>(31)</sup>. The present data demonstrated that maternal supplementation with SWE down-regulated the gene expression of a panel of pro-inflammatory cytokines (*IL-6*, *TGF-β* and *TNF-α*) in the ileum and colon tissues. It has been shown that less activation of the immune system during the weaning period could result in an improvement in growth performance<sup>(32)</sup>. Second, it may be attributable to an increase in *E. coli*-phagocytising effect (not analysed in the present experiment). According to Leonard *et al.*<sup>(33)</sup>, piglets suckling SWE-supplemented sows had a greater percentage of *E. coli* phagocytising leucocytes and a reduced percentage of *E. coli*-phagocytising lymphocytes at weaning, indicating an enhancement of immune function against presenting pathogens. Third, it may be attributable to the positive changes in the intestinal architecture (increased VH and VH:CD ratio in the ileum). A reduction in VH and the VH:CD ratio has been associated with poor performance and increased incidences of scouring in pigs challenged with ETEC<sup>(34)</sup>. Villi are critical components of the digestive tract and their geometry provides an indicator of the absorptive capacity of the small intestine<sup>(7)</sup>. Turnover of the intestinal epithelium reflects a dynamic equilibrium between the production of enterocytes in the crypts and their subsequent desquamation from the villus<sup>(7)</sup>. The VH:CD ratio is a useful criterion for assessing intestinal health and function<sup>(7,35)</sup>. The results obtained from the present study showed that pigs weaned from the SWE-supplemented sows had longer villi and a higher VH:CD ratio in the ileum. Furthermore, sow dietary supplementation with SWE clearly prevented the damage to intestinal morphology, otherwise

**Table 6.** Effect of sow dietary treatment on the transcriptional response of pig genes related to immune response following an enterotoxigenic *Escherichia coli* (ETEC) K88 challenge on days 9 and 10 PW (Least-squares mean values with their standard errors, *n* 10 piglets per treatment group)

Items	SWE			ETEC			GI region			P (effect)*					
	No	Yes	SEM	No	Yes	SEM	Colon	Ileum	SEM	SWE	ETEC	GI	SWE × ETEC	SWE × GI	ETEC × GI
	FOXP3	1.06	1.07	0.06	1.09	1.04	0.07	1.07	1.05	0.06	0.898	0.560	0.823	0.271	0.266
IFN-γ	1.05	1.40	0.14	1.15	1.30	0.14	1.17	1.29	0.14	0.088	0.458	0.559	0.530	0.123	0.823
IL-1	1.01	1.15	0.06	1.02	1.13	0.07	1.10	1.06	0.06	0.143	0.249	0.659	0.177	0.165	0.488
IL-2	1.11	1.10	0.09	1.15	1.07	0.09	1.10	1.11	0.09	0.954	0.521	0.957	0.312	0.212	0.755
IL-6††	1.10	1.00	0.05	1.08	1.01	0.06	1.03	1.06	0.05	0.231	0.373	0.683	0.039	0.224	0.031
IL-8§	1.20	1.28	0.15	1.29	1.19	0.15	1.14	1.34	0.15	0.719	0.667	0.346	0.934	0.008	0.823
IL-10‡	1.12	1.03	0.08	0.95	1.20	0.08	1.04	1.12	0.08	0.484	0.046	0.496	0.496	0.911	0.036
IL-12A (p35)‡	0.99	1.09	0.05	0.97	1.10	0.05	1.03	1.05	0.05	0.234	0.097	0.792	0.856	0.743	0.030
IL-17	1.13	1.10	0.09	1.10	1.12	0.09	1.05	1.18	0.09	0.818	0.852	0.306	0.168	0.184	0.937
TGF-β1‡§	1.07	0.98	0.04	1.04	1.00	0.04	1.03	1.02	0.04	0.102	0.452	0.870	0.115	0.050	0.004
TNF-α§	1.09	1.00	0.05	1.02	1.07	0.05	1.05	1.04	0.05	0.232	0.448	0.894	0.636	0.050	0.513

SWE, seaweed extracts; ETEC, enterotoxigenic *Escherichia coli* K88; GI, gastrointestinal; FOXP3, forkhead box P3; IFN-γ, interferon-γ; TGF-β1, transforming growth factor-β1.

\* There was no effect detected for the SWE × ETEC × GI interaction ( $P > 0.01$ ).

† There was an effect detected for the SWE × ETEC interaction ( $P < 0.05$ ).

‡ There was an effect detected for the ETEC × GI interaction ( $P < 0.05$ ).

§ There was an effect detected for the SWE × GI interaction ( $P < 0.05$ ).

caused by ETEC K88 infection alone, which is in accordance with the improved performance and lower incidence of diarrhoea in pigs weaned from those sows.

Diarrhoea caused by ETEC K88 is attributable to the action of one or more enterotoxins produced by the bacteria that have colonised the small intestine, such as *EAST1*, either as the only toxin gene or with the classic toxins associated with this pathotype, *LT* and *STb*, to impair the intestinal barrier function, which indirectly induces fluid losses<sup>(4,36,37)</sup>. In the present experiment, the ETEC K88 challenge increased the gene expression of *EAST1* and *STb*. Interestingly, in the present experiment, pigs weaned from the SWE-supplemented sows had reduced *LT* log GCN/g faeces compared with those weaned from the basal-fed sows. According to Fairbrother *et al.*<sup>(4)</sup>, ETEC K88 is a major cause of diarrhoea in pigs during days 3–10 PW. The pathogen colonises in the small intestine, releasing specific enterotoxins that induce fluid losses<sup>(4–6)</sup>. An *in vivo* study on gnotobiotic piglets revealed that *LT* is the major virulence factor of K88 ETEC, because it enhances bacterial colonisation of the intestine<sup>(38)</sup>. It may also be postulated from the present study that the reduction of ETEC K88 toxin *LT* in pigs weaned from the SWE-supplemented sows may be associated with the reduced faecal score of pigs weaned from those sows. However, a potential effect of ETEC was already present before the ETEC challenge begun, as evidenced by the GCN of ETEC toxins. Despite the presence of high GCN of ETEC toxins before the challenge, faecal scores were low, except for the BE pigs. Therefore, caution should be exercised when using the presence of toxin genes in the faeces as an indicator of ill health and diarrhoea in weaned pigs.

Maternal supplementation with SWE from day 83 of gestation and throughout the lactation period enhanced the immune response of weaned pigs reflected by a down-regulation of the gene expression of the pro-inflammatory cytokines *IL-6*, *TGF-β* and *TNF-α*. If maternal supplementation with SWE suppresses the secretion of pro-inflammatory cytokines, then less activation of the immune system during the weaning period would be achieved, which could result in an improvement in growth performance<sup>(32)</sup>. The gene expression of *IL-10* and *IL-12A* (*p35*) (pro-inflammatory cytokines) was up-regulated in the ileum of ETEC K88-challenged pigs compared with their expression in the ileum of unchallenged pigs. According to Girard *et al.*<sup>(39)</sup>, the first step in the pathogenesis of ETEC is adhesion to the ileal epithelium with the subsequent production of enterotoxins inducing intestinal acute diarrhoea and inflammation. Then, *E. coli* can be probably engulfed by antigen-presenting cells, e.g. macrophages, and evokes different courses of inflammation<sup>(40)</sup>. An interesting observation is that the gene expression of cytokines classically involved in ETEC challenge (i.e. *IL-1*, *IL-8* and *TNF-α*) was low or null in the present study. Perhaps some early sign may have already been passed in the small intestine on day 4 post-challenge. During ETEC infection, it is imperative to generate an adequate inflammatory response against the pathogen, accompanied by efficient regulation, in order to achieve protection without damaging the host tissues<sup>(41)</sup>. In the present study, the gene expression

of *IL-8* was up-regulated in the ileum of pigs weaned from the SWE-supplemented sows compared with its expression in the ileum of pigs weaned from the basal-fed sows. The gene expression of *IFN- $\gamma$*  tended to be up-regulated in the ileum and colon tissues of pigs weaned from the SWE-supplemented sows compared with those weaned from the basal-fed sows. The tissue collection was carried out on day 13 PW, the period during which PW diarrhoea due to ETEC K88 generally occurs in pigs<sup>(5,6)</sup>. Leonard *et al.*<sup>(24)</sup> reported that maternal supplementation with SWE enhanced the expression of a pro-inflammatory cytokine (*TNF- $\alpha$* ) in the ileum after an *ex vivo* lipopolysaccharide challenge at weaning. Rice *et al.*<sup>(42)</sup> also reported an increase of a pro-inflammatory cytokine (*IL-6*), in the serum, after oral administration of  $\beta$ -glucan. According to Xiao *et al.*<sup>(43)</sup>, monocytes/macrophages are the major effectors of innate immunity, functioning as the first barrier of natural defence.  $\beta$ -Glucans can stimulate macrophages, neutrophils and natural killer cells, and could promote T-cell-specific responses by the induction of cytokines such as *IFN- $\gamma$* , *IL-8* and *IL-12* from macrophages, neutrophils and natural killer cells<sup>(43–45)</sup>. Rice *et al.*<sup>(42)</sup> demonstrated that laminarin delivered orally can be internalised by intestinal epithelial cells and gut-associated lymphoid cells in the murine model. However, the possibility of mammary uptake of laminarin or fucoidan or both and subsequent introduction to the GI tract of suckling piglets was not examined in the present experiment. Furthermore, in the present experiment, the up-regulation of pro-inflammatory cytokines (*IL-8* and *IFN- $\gamma$* ) by maternal supplementation with SWE did not affect growth performance.

### Conclusion

The data reported in the present study indicate that maternal supplementation with SWE from day 83 of gestation to weaning prevented the negative effect on growth performance and diarrhoea induced by the ETEC K88 challenge. It does so by improving the function of intestinal health, offsetting or preventing impaired intestinal morphology, reducing faecal ETEC K88 toxin (*LT*) and down-regulating a panel of inflammatory cytokines. Collectively, these results indicated an important immunomodulatory role of maternal SWE supplementation on piglet immune function at weaning that may help alleviate the negative impact of a disease challenge at this period. These results provide new insights into the protective activity of maternal SWE supplementation, making it a good alternative for improving the health status of pigs with ETEC-associated diarrhoea.

### Acknowledgements

The present study was supported financially by the Irish Research Council for Science, Engineering and Technology (IRCSET) and BioAtlantis Limited. The IRCSET and BioAtlantis Limited had no role in the design or analysis of the study or in the writing of this article.

The authors' contributions are as follows: G. H. wrote the manuscript, collected the samples and carried out the

laboratory analysis; J. V. O'. D. was the principal investigator responsible for the design of the experiment, supervision of the data collection and statistical analysis and correction of the manuscript; T. S. designed the study and corrected the manuscript; C. J. O'. S. and D. N. D. contributed to the data collection and laboratory collection. All authors approved the final version of the manuscript.

None of the authors had a financial or personal conflict of interest in relation to the present study.

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