

Interaction of soyasaponins with plant ingredients in diets for Atlantic salmon, *Salmo salar* L.

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(Submitted 21 March 2011 – Final revision received 18 July 2011 – Accepted 2 August 2011 – First published online 14 September 2011)

Abstract

The effects of combining soyasaponins with plant ingredients on intestinal function and fish health were investigated in an 80 d study with Atlantic salmon (270 g) distributed thirty each into twenty-four tanks with seawater. Soyasaponins were supplemented (2 g/kg) to diets with maize gluten (MG), pea protein concentrate (PPC) and sunflower (SFM), rapeseed (RSM) or horsebean meals. A diet with soyabean meal (SBM) and another with wheat gluten and soyasaponins served as reference diets. Marked soyasaponin effects were observed when combined with PPC. This combination induced inflammation in the distal intestine (DI) similar to SBM, reduced feed intake, apparent digestibility of lipid, most amino acids and ash, decreased bile salt levels in intestinal chyme and decreased leucine aminopeptidase (LAP) activity but increased trypsin activity in the DI. No enteritis was observed in other diet groups, but small consistent negative soyasaponin effects were seen on lipid and fatty acid digestibility, faecal DM and LAP activity of the DI. Soyasaponin combination with RSM reduced digestibility of all nutrients including minerals. The mineral effect was also seen for SFM, whereas with MG and SFM a positive soyasaponin effect on feed intake was observed. Caution should be exercised to avoid ingredient combinations giving high saponin levels, a condition that appears to be a key factor in diet-induced enteritis together with certain plant ingredients.

Key words: Soyasaponins: Plant protein ingredients: Antinutritional factors: Fish feed: Gastrointestinal tract

Alternative dietary protein sources to supplement and replace limited marine ingredients in fish feeds are important for the future of the fish farming industry. Plant ingredients such as soyabeans hold promise with their good amino acid profiles that can easily be improved for fish requirements by supplementation with deficient amino acids. However, soyabean meal (SBM) inclusion has been demonstrated to induce enteritis and reduce performance in salmonids^(1–3) and carp⁽⁴⁾. The factors responsible for the disorders have not been conclusively identified, but soyasaponins, and possibly other bioactive antinutritional factors (ANF) in SBM, are implicated in the aetiology^(5–7).

Saponins are heat-stable glycosides present in soyabean and other legumes such as pea and lupin^(8,9). Saponins, with their

membrane-active nature and affinity to cholesterol and bile salts^(10,11), possess a number of potential biological effects compatible with the negative effects observed in fish fed diets containing SBM. There have been conflicting findings from studies on dietary effects of saponins to teleost fish. In one study, a saponin-rich extract from SBM and *Quillaja* saponins, both at a 0·3% dietary inclusion rate of saponin, greatly reduced feed intake and growth in Chinook salmon and depressed growth in rainbow trout⁽¹²⁾. Furthermore, the *Quillaja* saponin diets (0·15 and 0·3% dietary inclusion) both induced substantial damage to the intestinal mucosa for both Chinook salmon and rainbow trout. On the other hand, Francis *et al.*⁽¹³⁾ concluded that saponins from *Quillaja*

Abbreviations: AD, apparent digestibility; ANF, antinutritional factors; BBM, brush-border membrane; CP, crude protein; DI, distal intestine; EAA, essential amino acids; FDM, faecal DM; FER, feed efficiency ratio; GIT, gastrointestinal tract; HB, horsebean; HBM, horsebean meal; LAP, leucine aminopeptidase; MG, maize gluten; MI, mid-intestine; NEAA, non-essential amino acids; OSI, organo-somatic index; PI, pyloric intestine; PP, pea protein; PPC, pea protein concentrate; RS, rapeseed; RSM, rapeseed meal; SBM, soyabean meal; SFM, sunflower meal; ST, stomach; TGC, Thermal-unit growth coefficient; WG, wheat gluten.

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at similar dietary inclusion levels as previously indicated did not affect feed intake, but rather induced better feed conversion efficiency and growth in common carp (*Cyprinus carpio* L.) and Nile tilapia (*Oreochromis niloticus* L.). More recently, support has been strengthened for the involvement of saponins in the distal intestinal enteritis induced by SBM in Atlantic salmon; similar morphological changes were induced in Atlantic salmon when soyasaponins were added to a lupin kernel basal diet at inclusion levels of 0.17 and 0.26%^(6,7).

Findings from feeding salmonids diets with SBM have set a precedent in the approach to the evaluation of new plant ingredients for use as alternative protein sources in fish feeds. The present study was designed to investigate the impact of soyasaponins on fish performance and physiology and to find possible interactions between soyasaponins and current plant protein source, ingredients that, to some extent, have been investigated earlier as ingredients in diets for Atlantic salmon^(14–16).

Materials and methods

Diets

Effects of adding soyasaponin to maize gluten (MG; *Zea mays* L.), pea protein concentrate (PPC; *Lathyrus aphaca*), sunflower meal (SFM, *Helianthus annuus*), rapeseed meal (RSM, *Brassica napus*) or horsebean meal (HBM, *Vicia faba* var. *equina*) as dietary plant protein source were investigated. The five protein sources were included at levels as high as practically possible in commercial diet formulations, taking the varying fibre content into consideration, with standardisation regarding protein replacement and protein:energy ratio. As the fibre and protein content of the selected sources differed greatly, two levels of dietary protein replacement were used; MG and PPC were included at a level corresponding to 33% of total protein, while SFM, RSM and HBM were at 21%. The diets were formulated to contain an equal crude protein (CP):energy ratio of 20 g/MJ. The strategy was to let dietary energy levels vary to avoid using fillers in the diets that may influence results. Each of these five protein sources was investigated without and with soyasaponin supplementation (0.2%). Hereafter, the term saponin will refer to soyasaponin regarding the supplementation in the present study. The supplementation level approximated a level provided by a 20–30% SBM dietary inclusion. In addition, two reference diets were made, one with wheat gluten (WG, *Triticum* spp.) at an inclusion corresponding to 33% of total protein and supplemented with saponins, and a second with SBM (*Glycine max*) included at 21% of total protein, a level known to cause clear enteritis but usually without severe depression of feed intake.

In all of the diets, protein from the various plant sources partially replaced marine fish protein derived from a combination of Nordic LT (Norsildmel AS, Bergen, Norway) and South American Superprime fishmeals. All diets were supplemented with standard vitamin and micromineral premixes and contained 100 mg/kg yttrium oxide as an inert marker

for the calculation of nutrient apparent digestibilities (AD). The diets were produced by extrusion at the BioMar Feed Technology Centre (Brande, Denmark) with a pellet size of 5 mm in batches of 50 kg. Detailed diet formulations are shown in Table 1 and chemical composition, as analysed, is shown in Tables 2 and 3.

Experimental animals and conditions

The present experiment was conducted in compliance with laws regulating experimentation with live animals in Norway as overseen by the Norwegian Animal Research Authority (Forsøksdyrutvalget). The feeding trial was conducted at the land-based Nofima Marin research station at Sunndalsøra, Norway. Atlantic salmon (*Salmo salar* L.) post smolts of the Sunndalsøra breed with mean weight of 270 g ± 10% were allocated, thirty each, into twenty-four, 1 m³ fibreglass tanks with 500 litres of saltwater flowing at a rate of 20 litres/min. Water temperature varied between 9 and 13°C. The oxygen content and salinity of the outlet water were monitored. Salinity ranged between 31.4 and 33.3 with an average of 32.4 g/l. A 24 h lighting regimen was employed during the experimental period.

The fish were fed to satiation using automatic disc feeders refilled every 3 d with weighed amounts; waste feed was collected daily, separated from faeces, weighed and stored at –20°C. Every 3 d, the waste feed level and the percentage recovery of DM from each diet were used to determine the approximate feed intake for each tank. The feed intake for each tank was in turn used to adjust the feeding level every 3 d to provide at least 20% excess feed per d. Before starting the experiment, all fish per tank were weighed in bulk.

Sampling

At the start of the feeding period, twenty fish from the same group as the fish in the experiment were sampled, euthanised by a lethal dose of tricaine methane-sulfonate (MS222; Argent Chemical Laboratories, Inc., Redmond, WA, USA), individually weighed and frozen for whole body composition analysis.

The feeding trial ran for 80 d with an intermediate sampling (day 30) and a terminal sampling (day 80). At both samplings, tank order and fish sampling were conducted randomly. All sampled fish were euthanised by overdosing with tricaine methane-sulfonate (MS222) and weight and length recorded.

On day 30, the fish were weighed in bulk and three fish from each tank were weighed individually and sampled for histology. The fish were dissected and the gastrointestinal tract (GIT) removed and cleaned of associated adipose tissue. Liver, and mid (MI) and distal intestines (DI) were weighed to calculate organo-somatic indices (OSI). Histology samples were taken from the pyloric, mid and distal segments of the intestine and the liver, fixed in phosphate-buffered formalin (4% formaldehyde) for 24 h and then transferred to ethanol (70%).

At termination of the experiment (day 80), twelve fish from each tank were randomly selected and euthanised as

Table 1. Formulation of the experimental diets*

Ingredients	Diets (%)											
	MG-0	MG-S	PP-0	PP-S	SF-0	SF-S	RS-0	RS-S	HB-0	HB-S	WG-S	SBM
Nordic LT-meal†	23.1	23.1	21.5	21.5	26.2	26.2	24.8	24.8	24.5	24.5	23.9	26.5
Superprime fish meal‡	23.1	23.1	21.5	21.5	26.2	26.2	24.8	24.8	24.5	24.5	23.9	26.5
WG§	–	–	–	–	–	–	–	–	–	–	21.0	–
MG	26.0	26.0	–	–	–	–	–	–	–	–	–	–
PPC¶	–	–	31.0	31.0	–	–	–	–	–	–	–	–
HP soya**	–	–	–	–	–	–	–	–	–	–	–	20.0
HP sunflower††	–	–	–	–	23.0	23.0	–	–	–	–	–	–
RSM‡‡	–	–	–	–	–	–	27.0	27.0	–	–	–	–
HB§§	–	–	–	–	–	–	–	–	34.6	34.6	–	–
Saponins	–	0.2	–	0.2	–	0.2	–	0.2	–	0.2	0.2	–
Tapioka¶¶	6.0	6.0	6.0	6.0	6.0	6.0	6.0	6.0	0.0	0.0	6.0	6.0
Fish oil	11.8	11.8	10.7	10.7	10.4	10.4	9.9	9.9	9.6	9.6	12.9	11.3
RS oil	11.8	11.8	10.7	10.7	10.4	10.4	9.9	9.9	9.6	9.6	12.9	11.3
Vitamin–mineral-mix***	0.38	0.38	0.38	0.38	0.38	0.38	0.38	0.38	0.38	0.38	0.38	0.38
Lys	0.21	0.21	–	–	–	–	–	–	–	–	0.13	–
DL-Met	–	–	0.37	0.37	–	–	–	–	–	–	–	–
Carophyll pink	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04	0.04
Monocalcium phosphate	0.30	0.30	0.51	0.51	–	–	–	–	–	–	0.30	–

MG, maize gluten, -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal; PPC, PP concentrate; RSM, RS meal.

* The plant ingredients were included at levels about as high as possible taking into account bulkiness and expected effects on feed intake and fish performance decided based on earlier experiences. The low-fibre plant ingredients – MG, PPC and WG – were included at levels corresponding to 33% of total crude protein, whereas the high-fibre plant ingredients – SF meal, RSM, HB meal and SBM – were included at levels corresponding to 21% of total protein.

† Nordic LT 94 fishmeal supplied by Norsildmel AS, Bergen, Norway.

‡ Superprime fishmeal supplied by Köster Marine Proteins GmbH, Hamburg, Germany.

§ WG supplied by Roquette (Beinheim, France).

|| MG supplied by Cargil Nordic (SAS van Gent, Holland).

¶ PPC made from yellow peas by air classification; supplied by AgriMarin Nutrition (Stavanger, Norway).

** SBM supplied by Scanmills AS (Kolding, Denmark).

†† Heat-treated and hexane-extracted SF meal; supplied by DLA Agro (Galten, Denmark).

‡‡ Heat-treated RSM supplied by Emmelev Mølle (Otterup, Denmark).

§§ Whole horsebeans supplied by Overgaard Gods (Havndal, Denmark).

||| The 95% soyasaponin extract from soyabeans (*Glycine max*) supplied by Organic Technologies (Ohio, USA).

¶¶ Tapioca supplied by KMC, Brande, Denmark.

*** Supplied to ensure that the diets cover requirements for vitamins and minerals.

described earlier. From six fish, blood was collected in heparinised vacutainers for plasma preparation. From these same fish, the GIT was removed. Intestinal contents from the stomach (ST), cranial and caudal halves of the pyloric intestine (PI1 and PI2), MI and the cranial and caudal halves of the DI (DI1 and DI2) were collected into pre-weighed tubes. Tubes were subsequently frozen in liquid N₂ and stored at –80°C for bile salt and trypsin analyses. The organs were put back in place; the fish were re-weighed, individually packed into plastic bags and frozen at –20°C for whole body analyses. From the other six fish, the GIT was removed and freed of associated adipose tissue before histology samples were collected from the cardiac (ST1) and pyloric ST (ST2), PI, MI, DI, liver, spleen, head kidney, trunk (urinary) kidney, gills and muscle, and fixed in neutral buffered formalin for 24 h and then transferred into 70% ethanol. The liver was weighed before the histological samples were collected. The remaining MI and DI tissues were collected into pre-weighed containers, frozen in liquid N₂ and stored at –80°C for brush-border membrane (BBM) enzyme activity analysis as described previously. Faeces were stripped from the fish remaining in the tanks after sampling for AD measurements. Faeces were stored frozen at –20°C until freeze-drying and analysis. Samples of each diet were collected at start and end of the experiment and stored at –40°C for proximate analysis.

Analyses

Chemical analyses. Diet and faecal samples were analysed for DM (after heating at 105°C for 16–18 h), ash (combusted at 550°C to constant weight), nitrogen (CP) (by the semi-micro-Kjeldahl method, Kjeltec-Auto System, Tecator, Höganäs, Sweden), fat (diethyl ether extraction in a Fosstec analyser (Tecator, Höganäs, Sweden) after HCl-hydrolysis), starch (measured as glucose after hydrolysis by α-amylase (Novo Nordisk A/S, Bagsvaerd, Denmark) and amylo-glucosidase (Bohringer Mannheim GmbH, Mannheim, Germany)), followed by glucose determination by the 'Glut-DH method' (Merck, Darmstadt, Germany)), gross energy (using the Parr 1271 Bomb calorimeter, Parr, Moline, IL, USA) and yttrium (by inductivity coupled plasma mass-spectroscopy as described by Refstie *et al.*⁽¹⁷⁾). The amino acids in the diet were analysed using a Biochrom 30 amino acid analyser (Cambridge, UK) following the EC Commission Directive 98/64/EC (1999), after hydrolysis in 6 M-HCl for 23 h at 110°C. Tryptophan and tyrosine were analysed after basic hydrolysis.

Plasma variables. Plasma was analysed for NEFA, cholesterol, total TAG and glucose following standard procedures at the Central Laboratory of the Norwegian School of Veterinary Science (NVH), Oslo, Norway.

Trypsin activity and bile salt analyses in intestinal content. Faecal trypsin and bile salt analyses were performed on

pooled freeze-dried gastrointestinal contents from ST, PI1, PI2, MI, DI1 and DI2.

Trypsin activity was determined colorimetrically, according to Kakade *et al.*⁽¹⁸⁾, using the substrate benzoyl-arginine-*P*-nitroanilide (Sigma no. B-4875; Sigma Chemical Company, St Louis, MO, USA) and a curve derived from a standardised bovine trypsin solution.

Bile salt concentration was determined using the enzyme cycling amplification/Thio-NAD method (Inverness Medical, Cheshire, UK) in the ADVIA[®] 1650 Chemistry System (Siemens Healthcare Diagnostics, Inc., Deerfield, IL, USA) at the Central Laboratory of NVH.

Brush-border membrane enzyme activity analyses. Activity of the BBM enzyme leucine aminopeptidase (LAP; EC 3.4.11.1) was measured in intestinal tissue homogenates. The homogenates were prepared from tissues thawed in ice-cold Tris-mannitol buffer (1:20 w/v) containing the serine proteinase inhibitor 4-(2-aminoethyl) benzenesulfonyl fluoride HCl (Pefabloc[®] SC; Pentapharm Limited, Basel, Switzerland). Activity of LAP was determined colorimetrically using L-leucine- β -naphthylamide as the substrate as described by Krogdahl *et al.*⁽¹⁹⁾.

Protein concentration of the homogenates was estimated using the BioRad[®] Protein Assay (BioRad Laboratories, Munich, Germany). Tissue protein concentration was used in the determination of LAP specific activity.

Intestinal histology. Histology samples were processed according to standard histological techniques and stained with haematoxylin and eosin. The sections were randomised to ensure blinded examination and evaluated using a light microscope. A visual analogue (continuous) scale type scoring system as described by Penn *et al.*⁽²⁰⁾ was used to evaluate the intestinal histology. The following tissue characteristics were evaluated: (1) length and fusion (bridging) of the mucosal folds; (2) width and cellularity (leucocyte infiltration, connective tissue hyperplasia) of the lamina propria and submucosa; (3) degree of supranuclear absorptive vacuolisation and nucleus position of enterocytes; (4) frequency of intraepithelial lymphocytes and goblet cells. These are characteristics reported altered in SBM-induced enteritis in Atlantic salmon⁽²⁾.

Calculations

CP was calculated as $N \times 6.25$. Thermal-unit growth coefficient (TGC) was calculated as: $TGC = (FBW^{1/3} - IBW^{1/3}) \times (\Sigma D^{\circ})^{-1}$, where IBW and FBW are the initial and final body weights (tank means) and ΣD° is the thermal sum (feeding days \times average temperature in $^{\circ}C$). Feed intake was estimated by subtracting uneaten (waste) feed from fed feed on a DM basis. The uneaten feed was corrected for DM losses during feeding and collection using estimates of recovery of uneaten feed as described by Helland *et al.*⁽²¹⁾. Feed efficiency ratio (FER) was calculated as: $G \times F$, where G is the weight gain and F the consumption of DM from the feed. Instantaneous (daily) feed intake (% of body weight) was estimated as: $FE = 100 \times F_d \times (W_{d-1} + (F_{d-1} \times FER_p))$, where F_d represents feed intake at day, W_{d-1} and F_{d-1} are weight and feed intake the previous day and FER_p is

Table 2. Proximate composition and mineral content (n=2) of the experimental diets as analysed

Nutrient	Diets												SD*	
	MG-0	MG-S	PP-0	PP-S	SF-0	SF-S	RS-0	RS-S	HB-0	HB-S	WG-S	SBM		
DM (g/kg)	947	949	945	949	935	937	940	935	932	946	938	954	954	2.2
CP (g/kg DM)	504	503	475	492	462	451	449	438	450	474	565	522	522	5.6
Lipid (g/kg DM)	303	304	276	272	287	285	277	282	256	252	308	283	283	1.8
Starch (g/kg DM)	91	93	73	76	45	44	49	52	121	122	74	59	59	0.7
Gross energy (MJ/kg DM)	26	26	25	25	25	25	25	25	24	24	26	25	25	0.4
Ash (g/kg DM)	78	76	90	93	97	96	97	92	91	92	84	103	103	1.4
P (mg/kg DM)	12396	12796	14240	15047	14735	15162	14683	14027	13411	13576	13306	14436	14436	800
Ca (mg/kg DM)	15900	16262	16107	16993	17705	18475	19002	18034	17618	17445	18368	19033	19033	1500
Na (mg/kg DM)	1405	1453	1996	2011	2901	3071	2842	2699	1933	1871	1598	2209	2209	400
Mg (mg/kg DM)	6437	7145	5883	6706	7985	7186	7035	6648	7876	7935	8399	8385	8385	550
Zn (mg/kg DM)	198	197	216	221	214	217	217	207	217	218	213	219	219	13
Fe (mg/kg DM)	116	123	125	131	126	129	129	134	177	138	132	144	144	10
Mn (mg/kg DM)	53	51	60	60	57	57	65	66	56	57	60	57	57	4.3
Cu (mg/kg DM)	9	10	12	12	14	14	9	9	12	12	8	10	10	0.6

MG, maize gluten; -0, non-inclusion; -S, inclusion; PP-, rapeseed; RS-, sunflower; SF-, pea protein; PP-, rapeseed; HB-, horsebean; WG, wheat gluten; SBM, soyabean meal; CP, crude protein. * For the minerals, the sd indicate the upper limit of acceptable standard deviation for the procedure. For the other variables, the sd indicate pooled sd as calculated from the individual observations.

FER during the present experimental period. OSI were calculated as percentages of the weight of the organ in relation to body weight. AD was estimated by the indirect method using Y_2O_3 as an inert marker⁽²²⁾ and calculated as: $AD_N = 100 - (100 \times (M_{feed}/M_{faeces}) \times (N_{faeces}/N_{feed}))$, where M_{feed} and M_{faeces} are percentage concentration of the inert marker (Y_2O_3) in feed and faeces, respectively, and N_{feed} and N_{faeces} represent percentage concentration of a nutrient in feed and faeces, respectively. Nutrient retention – retentions of CP, individual amino acids and energy – was calculated as: $100 \times ((FBW \times C_1) - (IBW \times C_0)) \times (F \times C_{diet})^{-1}$, where C_{diet} is the content in the diets, and C_0 and C_1 are the initial and final contents in the fish.

Statistical analyses

The results were analysed using SAS Enterprise Guide 4.1 statistical software (SAS Institute, Inc., Cary, NC, USA). Tank means were used as the statistical unit in the analyses because individual fish responses were not considered independent within a tank. Saponin inclusion and basal diet were evaluated as class variables in a two-way ANOVA with interaction. Significant interaction between the main effects was observed for many variables and a one-way ANOVA was then used as an aid for the interpretation of data. When the interaction was significant, effects of saponin supplementation were evaluated based on differences in effect within the various basal diets, whereas effects of basal diet were evaluated based on the differences observed between fish fed the unsupplemented diets, i.e. MG-0, pea protein (PP-0), sunflower (SF-0), rapeseed (RS-0) and horsebean (HB-0). When there was no significant interaction, the basal diets were compared according to their main effects and mentioned as MG, PPC, SFM, RSM and HBM. The one-way ANOVA also allowed inclusion in the comparison of the SBM and the

WG-S diets that were excluded from the two-way ANOVA. The level of significance was set at $P < 0.05$, and P -values between 0.05 and 0.1 were considered as indications of effects mentioned as trends. The Duncan’s multiple range test was employed as the mean separating technique.

Results

The results for the two observation periods were generally similar but the effects were less clear after the first period, day 0–30. Results that follow are presented with main emphasis on the observations for the period, day 31–80. In the following presentation of results, under each subheading, the effects of saponin supplementation are presented first; thereafter, considerations regarding the basal diets are given. The basal diets are presented and discussed using abbreviations; MG represents the basal diet with MG and is used when there was no significant interaction between saponin supplementation and basal diet. When the interaction was significant, the results for the unsupplemented diet is given and presented as MG-0. Similar abbreviations are used for the other basal diets. When the protein source is mentioned as such, the full term is used.

Reference diets

The fish fed the SBM reference diets showed all the expected signs of enteritis usually observed in the DI of Atlantic salmon and confirmed that the experiment had the right conditions to reveal if saponins are involved in the development of this condition. The signs were: reduced bile salt concentration, low activity of BBM LAP, high chyme trypsin activity, reduced OSI, reduced mucosal fold height, increased width and cell infiltration of lamina propria and submucosa, etc.^(1,19). The results for the other reference diet, the WG-S, did not add

Table 3. Amino acid composition (*n* 2) in the experimental diets

	Diets (% of protein)												Pooled sd
	MG-0	MG-S	PP-0	PP-S	SF-0	SF-S	RS-0	RS-S	HB-0	HB-S	WG-S	SBM	
Essential amino acids													
Arg	5.4	5.5	7.7	7.8	7.3	7.3	6.8	6.8	7.6	7.6	5.7	7.2	0.08
His	2.7	2.7	3.0	3.0	3.1	3.1	3.1	3.1	3.1	3.1	2.8	3.1	0.04
Ile	4.3	4.3	4.5	4.4	4.6	4.5	4.6	4.5	4.6	4.5	4.3	4.7	0.06
Leu	10.9	10.9	8.0	8.0	7.9	7.9	8.0	8.0	8.1	8.1	7.8	8.1	0.16
Lys	6.2	6.3	8.1	8.3	7.8	7.8	8.0	8.0	8.2	8.2	6.3	8.1	0.09
Met	2.7	2.8	3.3	3.2	3.1	3.1	3.0	3.0	2.7	2.7	2.6	2.9	0.05
Phe	5.1	5.3	4.8	5.0	4.7	4.9	4.6	4.8	4.6	4.8	5.0	4.9	0.10
Thr	3.9	3.9	4.2	4.2	4.4	4.4	4.5	4.5	4.3	4.2	3.7	4.3	0.06
Trp	1.0	1.0	1.2	1.2	1.3	1.3	1.4	1.3	1.3	1.2	1.2	1.3	0.03
Val	5.1	5.1	5.3	5.2	5.5	5.4	5.6	5.5	5.4	5.3	4.9	5.4	0.08
Non-essential amino acids													
Ala	6.6	6.6	5.5	5.4	5.8	5.8	5.8	5.8	5.7	5.7	4.8	5.7	0.10
Asp + Asn	8.3	8.3	10.4	10.5	9.9	9.9	9.6	9.6	10.3	10.2	7.5	10.4	0.12
Cys	1.2	1.2	1.0	1.0	1.0	1.0	1.2	1.2	1.0	1.0	1.3	1.0	0.02
Glu + Gln	17.3	17.1	15.5	15.4	15.7	15.6	15.3	15.3	15.2	15.5	22.8	15.5	0.26
Gly	4.6	4.6	5.1	5.2	5.9	5.9	5.8	5.8	5.5	5.5	4.9	5.5	0.07
Pro	6.2	6.1	4.4	4.2	4.6	4.5	5.0	4.9	4.6	4.4	6.8	4.4	0.17
Ser	4.4	4.4	4.4	4.5	4.3	4.3	4.3	4.3	4.4	4.4	4.3	4.3	0.08
Tyr	4.0	4.0	3.6	3.5	3.3	3.2	3.3	3.3	3.4	3.4	3.3	3.4	0.08

MG, maize gluten, -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.

important information to the results obtained with the basal diets. The results for the SBM and WG-S are therefore not further presented here, except in a few cases.

Feed intake. Supplementation of the basal diets with saponins affected DM feed intake, but the effect was dependent on the basal diet (Table 4). The saponin–basal diet interaction was significant in the second feeding period, day 31–80. Inclusion of saponins significantly decreased feed intake for the PPC basal diet, but for the other basal diets feed intake seemed to be constant or to slightly increase, especially in the case of SFM basal diet for which the increase was significant.

Fish fed different basal diets differed in feed intake. In the first period, feed intake was highest in fish fed SF-0 and PP-0, and lowest in fish fed HB-0. In the second feeding period, there was a general increase in feed intake for all basal diets, with the HB-0 showing the highest increase of about 100%. The highest feed intakes were observed for fish fed PP-0 and HB-0. The fish fed RS-0 had the lowest intake.

Body weight and thermal-unit growth coefficient. As for feed intake, there was a significant interaction between saponin supplementation and basal diet for body weight and TGC (see Table 4). Fish fed the PP-S diet showed significantly lower values than fish fed the PP-0 diet both at day 30 and at the final sampling. For the other basal diets, saponin supplementation rather increased growth, but not significantly.

There were significant differences between the basal diets regarding growth parameters observed on day 30. Fish fed diet PP-0 had the highest values, while fish fed RS-0 had the lowest. Also at termination, fish fed PP-0 had the highest weight and TGC, and fish fed RS-0 remained the smallest. The ranking of the basal diets for TGC was as follows: PP-0^a > HB-0^{a,b} > MG-0^{a,b,c,d} > SF-0^{c,d,e} > RS-0^e (diets with different superscripts differed significantly). The HB-0 and SF-0 had switched place compared to the first period, and were now significantly different.

Feed efficiency ratio. Saponin supplementation did not significantly affect the FER (see Table 4) for any of the basal diets. Differences in FER were caused by differences between the basal diets. During the first 30 d of the trial, FER was highest for fish fed PPC followed by MG and HBM with no significant difference between the three. The RSM had the lowest FER, significantly lower than the other basal diets. In the second feeding period, the picture was somewhat different with the MG showing the highest FER followed by PPC, HBM, SFM and RSM. The latter two differed significantly from the MG.

Apparent nutrient digestibility

Macronutrients. Saponin inclusion was associated with a small but significant reduction in AD of lipids (Table 5). The effect was most pronounced for the PPC, but the overall saponin–basal diet interaction was not significant. Saponin supplementation also affected protein AD, but the effect depended on the basal diet. For the PPC diet, the effect was negative. A negative trend was also apparent for RSM. For MG and HBM, the trend was positive, but only marginally. Saponin inclusion did not affect starch AD significantly. For energy AD, the saponin effect was clearly negative for

the PPC and RSM, whereas for the other basal diets, the trend was positive. Regarding faecal DM (FDM), significant negative effects of saponin supplementation were seen, depending on the basal diet. The effect was greatest for PPC, for which saponin supplementation reduced FDM by 6% units. For SFM, RSM and HBM, the reduction was about 1% unit, whereas for MG, the reduction was negligible.

Significant differences were observed regarding AD of all the macronutrients and energy due to differences between the basal diets. Fish fed MG-0 had the highest protein and energy AD; fish fed RS-0, the lowest. For lipid AD, the highest value was observed in fish fed PPC diets, being significantly higher than all the other basal diets, whereas the lowest value was observed in fish fed SFM diets. The largest variation in nutrient AD was observed for starch. The highest value was observed with MG and the lowest with HBM. FDM was about 14% for most basal diets, except the HBM (observed for the HB-0), which showed significantly higher FDM than the other supplemented diets.

Amino acids

Essential amino acids. Generally, AD coefficients for all of the essential amino acids (EAA) were high, predominantly above 90% (see Table 6). Saponin inclusion effects on individual EAA differed and depended on the basal diet. For MG, saponin inclusion was consistently associated with a significantly higher AD for all EAA except Leu. Even for Leu, the trend was maintained as the MG-S diet had a numerically higher AD than the MG-0 diet. Saponin inclusion for the rest of the basal diets had either a negative effect on AD of the EAA or no effect. For PPC, the AD of all EAA was markedly lower for fish fed the saponin-supplemented diet. The pattern was similar for the RSM, except that differences in the AD of His, Lys and Phe were not significant. There were no clear saponin supplementation effects for SFM and HBM for any of the EAA.

The MG-0 and HB-0 generally showed high AD values across all the EAA, and AD of Leu, Trp and Phe were significantly higher for MG-0 than for HB-0. Also, the PP-0 diet showed quite high amino acid AD. In contrast, fish fed the RS-0 consistently had the lowest AD values for all the EAA, while fish fed the SF-0 were intermediate. Among the EAA, Trp showed the lowest values.

Non-essential amino acids and taurine. Generally, AD of the non-essential amino acids (NEAA) were not as high as for the EAA (see Table 7). Saponin effects were similar to those observed for the EAA AD values. Saponin inclusion in the MG basal diet was associated with significantly higher digestibilities of Ala, Asp-Asn and Gly, and with numerically higher values for the remaining NEAA. Saponin inclusion in the PPC showed significantly lower AD in the saponin-supplemented diet for all the NEAA. The greatest difference was seen for Cys for which AD dropped 12% units due to the saponin inclusion. Saponin inclusion was also associated with significantly lower AD for RSM diets for most NEAA except Cys and Asp-Asn.

The AD of all NEAA were generally highest for fish fed the MG-0 diet and lowest for fish fed the RS-0 diet. The lowest AD and the greatest differences were found for Cys.

Table 4. Feed intake, growth and feed utilisation efficiency of Atlantic salmon during the feeding period

	BW (g)			DM intake (% mean BW/d)		TGC × 1000		FER (DM basis, g/g)	
	Day 0	Day 30	Day 80	Day 0–30	Day 31–80	Day 0–30	Day 31–80	Day 0–30	Day 31–80
Two-way ANOVA model									
<i>P</i> (model)	0.18	0.16	0.01	0.02	0.002	0.10	0.0004	0.05	0.01
Pooled SEM	1.4	9.5	25.8	0.04	0.03	0.2	0.1	0.8	0.04
<i>P</i> values effect tests in the two-way ANOVA model									
Saponin	0.99	0.87	0.73	0.36	0.19	0.89	0.25	0.55	0.64
Basal diet	0.08	0.12	0.01	0.01	0.002	0.06	0.0001	0.01	0.001
Interaction	0.34	0.18	0.03	0.09	0.01	0.16	0.01	0.37	0.27
Marginal means for the two-way ANOVA model									
Saponin supplementation									
No saponin	262	319	537	0.52	0.78	1.09	2.63	1.18	1.34
With saponin	262	318	543	0.54	0.80	1.07	2.72	1.14	1.35
Basal diet									
MG	262	318	547 ^a	0.50 ^{b,c}	0.77 ^{b,c}	1.09	2.79 ^{a,b}	1.24 ^a	1.45 ^a
PPC	260	330	578 ^a	0.58 ^{a,b}	0.81 ^b	1.34	2.87 ^{a,b}	1.28 ^a	1.39 ^{a,b}
SFM	263	325	549 ^a	0.60 ^a	0.81 ^b	1.20	2.64 ^b	1.14 ^a	1.31 ^{b,c}
RSM	262	303	461 ^b	0.52 ^{a,b,c}	0.71 ^c	0.81	2.06 ^c	0.89 ^b	1.22 ^c
HBM	265	314	563 ^a	0.44 ^c	0.87 ^a	0.95	3.01 ^a	1.23 ^a	1.37 ^{a,b}
Means of the diets for the one-way ANOVA model									
MG-0	261	316 ^c	529 ^{b,c,d}	0.49 ^{c,d,e}	0.76 ^{b,c,d}	1.07 ^{b,c}	2.67 ^{a,b,c,d}	1.25 ^a	1.42 ^{a,b,c}
MG-S	264	321 ^{a,b,c}	566 ^{a,b,c}	0.52 ^{c,d,e}	0.78 ^{b,c,d}	1.11 ^{b,c}	2.92 ^{a,b,c}	1.22 ^a	1.48 ^{a,b}
PP-0	260	347 ^{a,b}	640 ^a	0.65 ^{a,b}	0.86 ^{a,b}	1.63 ^a	3.19 ^a	1.42 ^a	1.41 ^{a,b,c,d}
PP-S	260	313 ^c	517 ^{b,c,d}	0.52 ^{c,d,e}	0.75 ^{c,d}	1.04 ^{b,c}	2.56 ^{b,c,d}	1.15 ^{a,b}	1.38 ^{b,c,d}
SF-0	264	320 ^{a,b,c}	518 ^{b,c,d}	0.55 ^{b,c,d}	0.72 ^d	1.09 ^{b,c}	2.40 ^{c,d,e}	1.13 ^{a,b}	1.34 ^{c,d}
SF-S	262	330 ^{a,b,c}	580 ^{a,b}	0.65 ^{a,b}	0.89 ^a	1.31 ^{a,b}	2.89 ^{a,b,c}	1.15 ^{a,b}	1.27 ^{d,e}
RS-0	263	302 ^c	448 ^d	0.48 ^{c,d,e}	0.71 ^d	0.76 ^c	1.95 ^e	0.91 ^b	1.16 ^e
RS-S	261	305 ^c	475 ^{c,d}	0.56 ^{b,c}	0.71 ^d	0.86 ^{b,c}	2.18 ^{d,e}	0.87 ^b	1.28 ^{d,e}
HB-0	264	309 ^c	551 ^{a,b,c}	0.42 ^e	0.85 ^{a,b,c}	0.87 ^{b,c}	2.95 ^{a,b}	1.17 ^{a,b}	1.38 ^{b,c,d}
HB-S	265	319 ^{b,c}	576 ^{a,b}	0.46 ^{c,d,e}	0.89 ^a	1.04 ^{b,c}	3.07 ^{a,b}	1.30 ^a	1.36 ^{b,c,d}
WG-S	264	319 ^{b,c}	568 ^{a,b,c}	0.44 ^{d,e}	0.75 ^d	1.05 ^{b,c}	2.91 ^{a,b,c}	1.38 ^a	1.52 ^a
SBM	264	350 ^a	605 ^{a,b}	0.72 ^a	0.79 ^{a,b,c,d}	1.61 ^a	2.82 ^{a,b,c}	1.26 ^a	1.37 ^{b,c,d}

BW, body weight; TGC, thermal growth coefficient; FER, feed efficiency ratio; MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.
^{a,b,c,d,e} Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).

The AD of taurine, a sulfonic acid found in high concentration in both pancreatic juice and bile, is worth mentioning. There was no significant effect of saponin supplementation, but the PPC and RSM diets showed significantly lower taurine AD than the other basal diets.

Fatty acids. The AD of fatty acids are presented in Table 8. Saponin supplementation decreased AD of most fatty acids, significantly for 16:1*n*-7, 18:0, 18:1*n*-9, 20:2*n*-6, 22:1*n*-11 and sum MUFA. The same trend was seen for all other fatty acids except 18:1*n*-11 and 18:3*n*-3. No significant interactions between saponins and basal diet were observed.

The AD of most of the fatty acids were influenced by the basal diet. The trends for the SFA such as 16:0 and 18:0 were similar. The HBM and PPC showed the highest values, RSM and MG the lowest, significantly lower than the former two. The SFM showed intermediate values and did not differ from the other basal diets. The AD of MUFA was generally high, with PPC showing the highest in this category. The SFM generally showed the lowest AD, and the difference between the two was significant for most of the MUFA. In between, MG, RSM and HBM mostly ranked 2, 3 and 4 without being significantly different, but for some of the MUFA significant differences compared to PPC and SFM were observed. The AD of the *n*-6 and *n*-3 generally showed the same ranking of the basal diets as the MUFA.

Minerals. The AD values for minerals are presented in Table 9. They represent the result of intake of minerals from seawater and diet in addition to mineral absorption and secretion by the intestine. Sea water may supply the fish with more Na, Ca and Mg than the diet and consequently AD for these minerals may become negative, which was the case in particular for Mg. Also, AD for Mn showed several negative values, and for AD of the sum of all minerals, i.e. ash, many values were negative.

Saponin inclusion negatively affected the AD of Na and Mg for PPC, RSM and SFM. Ash AD showed the same picture and a similar trend was apparent for AD of Ca. For MG and HBM, there was no clear effect. Saponin supplementation increased or tended to increase AD of Cu, Fe, Mn, Zn and P for MG and SFM, and decreased AD of these minerals for RSM. For PPC and HBM, the results varied between the minerals. The only significant effects for these two basal diets were a positive effect of saponins for PPC on AD of Cu and a negative effect for HBM on Fe.

Significant differences were observed between the basal diets for all minerals except Ca. For Na, fish fed the HB-0 and MG-0 diets had the highest AD, significantly different from fish fed PP-0 and RS-0 diets which had the lowest. Fish fed the SF-0 diet showed intermediate values, not significantly different from other fish. Fish fed HB-0 also showed the highest AD of Mg, but the value was significantly different only

Table 5. Apparent digestibility of macronutrients and energy, and faecal DM (FDM) for Atlantic salmon fed the experimental diets (%)

	Crude protein	Lipid	Starch	Energy	FDM
Two-way ANOVA model					
<i>P</i> (model)	<0.0001	0.0487	<0.0001	<0.0001	<0.0001
Pooled SEM	0.5	0.4	1.6	0.5	0.4
<i>P</i> values effect tests in the two-way ANOVA model					
Saponin	0.46	0.05	0.58	0.15	<0.0001
Basal diet	<0.0001	0.04	<0.0001	<0.0001	<0.0001
Interaction	0.05	0.20	0.10	0.02	<0.0001
Marginal means for the two-way ANOVA model					
Saponin supplementation					
No saponin	86.1	96.1 ^a	81.5	84.3	14.9 ^a
With saponin	85.9	95.6 ^b	80.9	83.8	13.1 ^b
Basal diet					
MG	89.2 ^a	95.7 ^b	82.7 ^b	89.5 ^a	13.9 ^b
PPC	86.3 ^b	96.6 ^a	84.2 ^b	85.1 ^b	11.1 ^c
SFM	85.9 ^b	95.3 ^b	81.8 ^b	82.4 ^c	14.1 ^b
RSM	82.0 ^c	95.6 ^b	89.7 ^a	79.8 ^d	13.9 ^b
HBM	86.6 ^b	96.2 ^{ab}	67.7 ^c	83.5 ^c	16.9 ^a
Means of the diets for the one-way ANOVA model					
MG-0	88.6 ^{b,c}	95.5 ^b	80.3 ^{d,e}	89.0 ^b	13.9 ^{b,c,d}
MG-S	89.9 ^b	95.9 ^b	85.2 ^{b,c,d}	90.1 ^b	13.8 ^{c,d}
PP-0	87.4 ^{c,d}	97.3 ^a	83.8 ^{c,d}	86.2 ^c	14.1 ^{b,c,d}
PP-S	85.1 ^e	95.9 ^b	84.6 ^{b,c,d}	84.1 ^d	8.1 ^f
SF-0	85.9 ^{d,e}	95.6 ^b	82.9 ^d	82.2 ^{e,f}	14.6 ^{b,c}
SF-S	85.9 ^{d,e}	95.0 ^b	80.8 ^{d,e}	82.6 ^{d,e}	13.5 ^{c,d}
RS-0	82.6 ^f	96.1 ^b	90.4 ^a	80.8 ^f	14.4 ^{b,c,d}
RS-S	81.5 ^f	95.2 ^b	89.0 ^{a,b}	78.7 ^g	13.5 ^d
HB-0	86.3 ^{d,e}	96.2 ^b	70.3 ^f	83.4 ^{d,e}	17.3 ^a
HB-S	87.0 ^d	96.1 ^b	65.2 ^g	83.6 ^{d,e}	16.5 ^a
WG-S	92.4 ^a	95.6 ^b	77.3 ^e	91.7 ^a	15.0 ^b
SBM	86.9 ^d	95.3 ^b	88.4 ^{a,b,c}	86.3 ^c	9.5 ^e

MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.
^{a,b,c,d,e,f,g} Mean values within a column with unlike superscript letters were significantly different (*P*<0.05).

Table 6. Apparent digestibility of essential amino acids for Atlantic salmon fed the experimental diets (%)

	Lys	Met	Arg	Leu	Trp	His	Ile	Phe	Thr	Val
Two-way ANOVA model										
<i>P</i> (model)	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	0.001	<0.0001	<0.0001	<0.0001
Pooled SEM	0.4	0.3	1.0	0.4	0.4	0.6	0.3	0.5	0.3	0.3
<i>P</i> values effect tests in the two-way ANOVA model										
Saponin	0.01	0.01	0.01	0.004	0.04	0.26	0.29	0.03	0.14	0.01
Basal diet	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	0.0002	<0.0001	<0.0001	<0.0001
Interaction	0.001	0.002	0.0002	0.0004	0.004	0.01	0.01	0.002	0.002	0.001
Marginal means for the two-way ANOVA model										
Saponin supplementation										
No saponin	92.5 ^a	90.8 ^a	92.8 ^a	92.0 ^a	86.3 ^a	90.3	90.4	90.0	88.2 ^a	90.5 ^a
With saponin	92.3 ^b	90.6 ^b	92.6 ^b	91.5 ^b	84.9 ^b	89.9	89.7	90.0	87.3 ^b	89.9 ^b
Basal diet										
MG	93.5 ^a	91.9 ^a	93.5 ^a	95.0 ^a	86.5 ^a	91.7 ^a	92.2 ^a	92.7 ^a	90.4 ^a	92.6 ^a
PPC	91.9 ^d	90.4 ^c	92.0 ^d	89.9 ^c	79.0 ^c	88.4 ^b	87.5 ^b	88.9 ^b	84.4 ^d	87.5 ^c
SFM	92.4 ^c	90.5 ^b	92.7 ^c	91.5 ^b	87.2 ^b	89.7 ^c	90.6 ^b	89.8 ^b	87.8 ^c	90.7 ^b
RSM	89.8 ^e	89.4 ^c	91.5 ^c	89.3 ^d	84.6 ^d	88.5 ^d	87.4 ^c	87.4 ^b	84.4 ^e	87.5 ^d
HBM	93.4 ^b	90.5 ^b	93.2 ^b	92.2 ^a	87.7 ^b	90.8 ^a	91.0 ^b	90.3 ^a	89.6 ^b	91.2 ^b
Means of the diets for the one-way ANOVA model										
MG-0	93.0 ^{c,d}	91.3 ^{c,d}	93.1 ^{c,d}	94.7 ^b	85.6 ^{d,e}	91.1 ^c	91.6 ^c	92.1 ^c	89.8 ^c	92.0 ^c
MG-S	94.0 ^b	92.5 ^b	93.9 ^b	95.4 ^{a,b}	87.3 ^{b,c}	92.4 ^b	92.7 ^b	93.3 ^b	91.0 ^b	93.2 ^b
PP-0	93.7 ^{b,c}	91.7 ^{b,c}	93.4 ^c	91.9 ^{c,d}	85.1 ^{d,e}	91.0 ^c	90.5 ^d	90.4 ^d	88.7 ^{c,d}	90.3 ^{d,e}
PP-S	91.9 ^e	90.4 ^e	92.0 ^f	89.9 ^e	79.0 ^g	88.4 ^g	87.5 ^{e,f}	88.9 ^e	84.4 ^{f,g}	87.5 ^{f,g}
SF-0	92.4 ^{d,e}	90.4 ^{d,e}	92.5 ^{e,f}	91.5 ^{c,d}	87.3 ^{b,c}	89.6 ^{e,f}	90.5 ^d	89.6 ^{d,e}	87.9 ^{d,e}	90.7 ^{d,e}
SF-S	92.5 ^{d,e}	90.6 ^{d,e}	92.8 ^{d,e}	91.5 ^{c,d}	87.2 ^{b,c}	89.8 ^{d,e,f}	90.6 ^{c,d}	90.1 ^d	87.8 ^{d,e}	90.7 ^{d,e}
RS-0	90.2 ^f	90.1 ^e	92.0 ^f	89.9 ^e	86.0 ^{c,d}	88.9 ^{f,g}	88.2 ^e	87.8 ^f	85.2 ^f	88.0 ^f
RS-S	89.5 ^f	88.7 ^f	91.0 ^g	88.7 ^f	83.1 ^f	88.1 ^g	86.6 ^f	87.0 ^f	83.7 ^g	86.9 ^g
HB-0	93.4 ^{b,c}	90.4 ^e	93.2 ^{c,d}	92.3 ^c	87.5 ^{b,c}	90.8 ^{c,d}	91.0 ^{c,d}	90.1 ^d	89.5 ^c	91.3 ^{c,d}
HB-S	93.4 ^{b,c}	90.6 ^{d,e}	93.2 ^{c,d}	92.2 ^c	87.8 ^b	90.9 ^c	90.9 ^{c,d}	90.5 ^d	89.6 ^c	91.1 ^{c,d,e}
WG-S	95.4 ^a	93.9 ^a	95.3 ^a	95.6 ^a	92.0 ^a	94.4 ^a	94.9 ^a	94.9 ^a	93.0 ^a	94.8 ^a
SBM	93.0 ^{c,d}	90.8 ^{d,e}	92.9 ^{c,d,e}	91.3 ^d	84.3 ^{e,f}	90.2 ^{c,d,e}	90.3 ^d	90.2 ^d	87.2 ^e	90.0 ^e

MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.

^{a,b,c,d,e,f,g} Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).

when compared to fish fed MG-0 which had the lowest value. Fish fed HB-0 also had the highest AD for Fe, Mn and Zn, for which SF-0 had the lowest values. For Cu and P, the PP-0 diet had the highest values, whereas SF-0 again ranked the lowest. For Cu, Mn and P, there were no significant differences between the basal diets. For Fe, the HB-0 had a significantly higher AD than the diet with second highest AD, which was RS-0. For Zn, HB-0 was also significantly higher than the two lowest, PP-0 and SF-0.

Retention of nutrients

Retention of crude protein and energy. Table 10 presents retention values for CP and energy expressed as percentages of both ingested and digested material. Retention as a percentage of digested material showed greater variance and distinguished the diets less clearly than when expressed as a percentage of ingested material. The retention values were generally high and highest for fish fed the MG-S and WG-S diets.

Saponins did not affect nitrogen and energy retention values significantly, and no significant saponin–basal diet interactions were observed.

Retention of ingested CP differed significantly between the basal diets and was lower for RSM than the other basal diets. A similar trend was discernible for retention of ingested energy and digested CP. For retention of digested energy, the

trend was less clear, but the RSM had the lowest values also for digested energy.

Retention of essential amino acids and cysteine. Retention of amino acids showed similar results whether expressed per ingested or digested amino acids. Details for the retention of digested amino acids are presented in Table 11. No values were obtained for retention of Trp. Saponin inclusion showed no significant effect on the retention of any of the amino acids. Moreover, no saponin–basal diet interactions were significant for any of the amino acids.

There were significant differences between basal diets for retention of both ingested (data not shown) and digested methionine, cysteine, arginine, leucine and tyrosine as well as ingested lysine. The highest values were mostly seen for MG and HBM, whereas PPC and RSM had the lowest values. Apparent retention of digested cysteine was the highest among all amino acids, close to 90% for diets with HBM.

Bile salts and trypsin in intestinal content. Results of analysis of bile salts concentration and trypsin activity in the intestinal content along the intestine are presented in Table 12. Saponins reduced bile salt concentration in fish fed PPC by about 60% in the PI1, PI2 and MI sections. For MG and HBM, the effect seemed to be the opposite in PI1 and PI2. In the MI, the pattern seemed to change for MG and HBM towards a negative effect that tended to continue

Table 7. Apparent digestibility of non-essential amino acids and taurine in Atlantic salmon fed the experimental diets (%)

	Ala	Asp-Asn	Cys	Glu-Gln	Gly	Pro	Ser	Tyr	Taurine
Two-way ANOVA model									
<i>P</i> (model)	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	0.05
Pooled SEM	0.3	0.4	0.3	0.3	0.3	0.2	0.4	0.5	4.0
<i>P</i> values effect tests in the two-way ANOVA model									
Saponin	0.01	0.01	0.80	0.09	0.12	0.05	0.0004	0.001	0.70
Basal diet	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	<0.0001	0.01
Interaction	0.002	0.003	0.01	0.001	0.003	0.001	0.0001	0.0001	0.56
Marginal means for the two-way ANOVA model									
Saponin supplementation									
No saponin	90.9 ^a	83.2 ^a	73.8	92.5	82.6	86.7 ^a	88.0 ^a	89.5 ^a	48.0
With saponin	90.4 ^b	82.8 ^b	71.6	92.0	82.1	85.6 ^b	87.2 ^b	88.6 ^b	47.0
Basal diet									
MG	93.6 ^a	85.7 ^a	84.9 ^a	94.8 ^a	84.9 ^a	92.5 ^a	91.3 ^a	92.8 ^b	50.5 ^a
PPC	88.6 ^d	81.9 ^c	57.2 ^b	89.9 ^c	79.9 ^{b,c}	81.6 ^b	84.4 ^d	86.4 ^d	38.5 ^b
SFM	90.2 ^c	81.6 ^b	70.3 ^b	92.0 ^c	81.4 ^c	85.6 ^b	87.3 ^c	89.2 ^{a,b}	56.9 ^a
RSM	88.8 ^e	79.0 ^d	69.6 ^c	90.6 ^d	80.4 ^d	81.9 ^c	84.0 ^e	85.7 ^c	41.2 ^b
HBM	91.2 ^b	85.0 ^b	75.5 ^b	92.7 ^b	83.9 ^{ab}	87.0 ^a	89.1 ^b	89.4 ^a	48.6 ^{a,b}
Means of the diets for the one-way ANOVA model									
MG-0	93.1 ^b	84.7 ^c	83.6 ^b	94.4 ^b	83.9 ^c	92.0 ^b	90.8 ^b	92.5 ^b	47.3 ^{b,c,d,e}
MG-S	94.0 ^a	86.7 ^b	86.2 ^b	95.2 ^b	85.8 ^b	92.9 ^b	91.9 ^b	93.2 ^{a,b}	53.6 ^{a,b,c,d}
PP-0	90.5 ^{c,d}	85.3 ^{b,c}	69.1 ^d	92.5 ^{c,d}	83.0 ^{c,d}	85.8 ^d	88.1 ^{c,d}	89.9 ^c	42.1 ^{c,d,e}
PP-S	88.6 ^f	81.9 ^d	57.2 ^e	89.9 ^f	79.9 ^{g,h}	81.6 ^f	84.4 ^e	86.4 ^d	38.5 ^e
SF-0	90.1 ^{d,e}	81.5 ^d	70.4 ^d	92.0 ^{c,d}	81.1 ^{f,g}	85.7 ^d	87.2 ^d	89.3 ^c	56.1 ^{a,b,c}
SF-S	90.2 ^{d,e}	81.8 ^d	70.2 ^d	92.1 ^{c,d}	81.6 ^{d,e,f}	85.6 ^d	87.4 ^d	89.1 ^c	57.7 ^{a,b}
RS-0	89.5 ^e	79.6 ^e	70.7 ^d	91.0 ^e	81.4 ^{e,f}	83.3 ^e	84.8 ^e	86.6 ^d	42.9 ^{c,d,e}
RS-S	88.1 ^f	78.3 ^e	68.4 ^d	90.1 ^f	79.3 ^h	80.5 ^d	83.2 ^f	84.9 ^e	39.5 ^{d,e}
HB-0	91.1 ^c	84.9 ^c	75.1 ^c	92.7 ^c	83.7 ^c	86.8 ^c	88.9 ^c	89.2 ^c	51.6 ^{a,b,c,d,e}
HB-S	91.2 ^c	85.1 ^{b,c}	76.0 ^c	92.7 ^c	84.2 ^c	87.2 ^c	89.2 ^c	89.5 ^c	45.7 ^{b,c,d,e}
WG-S	93.8 ^{a,b}	88.8 ^a	91.1 ^a	97.4 ^a	89.2 ^a	95.0 ^a	93.8 ^a	94.3 ^a	64.6 ^a
SBM	90.2 ^{d,e}	83.9 ^c	70.1 ^d	91.7 ^{d,e}	82.8 ^{c,d,e}	85.3 ^d	87.0 ^d	88.6 ^c	50.7 ^{a,b,c,d,e}

MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.

^{a,b,c,d,e,f,g,h} Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).

Table 8. Apparent digestibility of fatty acids for Atlantic salmon fed the experimental diets (%)

	SFA	MUFA	n-3 PUFA	n-6 PUFA	16:0	16:1n-7	18:0	18:1n-7	18:1n-9	18:1n-11	18:2n-6	18:3n-3	20:1n-9	20:2n-6	22:1n-11	20:5n-3	22:6n-3
Two-way ANOVA model																	
<i>P</i> (model)	0.04	0.004	0.02	0.0002	0.05	0.001	0.02	0.02	0.002	0.0002	<0.0001	0.07	0.08	0.05	0.04	0.0002	0.06
Pooled SEM	1.2	0.2	0.2	0.2	1.2	0.1	1.6	0.3	0.3	0.1	0.1	0.2	0.3	0.5	0.4	0.1	0.3
<i>P</i> values effect tests in the two-way ANOVA model																	
Saponin	0.08	0.02	0.09	0.07	0.10	0.05	0.05	0.08	0.03	0.30	0.06	0.93	0.06	0.04	0.02	0.06	0.03
Basal diet	0.02	0.001	0.01	<0.0001	0.03	0.0002	0.02	0.01	0.0004	<0.0001	<0.0001	0.02	0.15	0.08	0.06	<0.0001	0.06
Interaction	0.22	0.18	0.30	0.51	0.22	0.21	0.13	0.47	0.16	0.94	0.41	0.31	0.11	0.11	0.11	0.41	0.27
Marginal means for the two-way ANOVA model																	
Saponin supplementation																	
No saponin	87.4	98.4 ^a	98.7	98.6	87.1	98.6 ^a	83.2 ^a	97.7	98.7 ^a	99.3	98.7	99.2	97.4	94.0 ^a	97.1 ^a	99.1	97.6 ^a
With saponin	85.9	98.1 ^b	98.4	98.4	85.8	98.4 ^b	80.9 ^b	97.3	98.5 ^b	99.3	98.6	99.2	97.0	93.2 ^b	96.5 ^b	99.0	97.1 ^b
Basal diet																	
MG	84.8 ^c	98.4 ^{a,b}	98.8 ^a	98.9 ^a	84.8 ^c	98.8 ^{a,b}	79.6 ^c	97.8 ^{a,b}	98.9 ^{a,b}	99.5 ^a	99.1 ^a	99.5	97.4	93.0	97.1	99.3 ^a	97.6
PPC	88.3 ^{a,b}	98.7 ^a	98.9 ^a	99.2 ^a	88.2 ^{a,b}	99.0 ^a	84.1 ^{a,b}	98.2 ^a	99.1 ^a	99.6 ^a	99.3 ^a	99.6	97.6	94.1	97.4	99.4 ^a	97.8
SFM	85.9 ^{b,c}	97.7 ^c	98.2 ^b	97.7 ^c	85.6 ^{b,c}	98.0 ^d	81.3 ^{b,c}	97.2 ^{b,c}	98.0 ^d	98.9 ^b	97.7 ^c	98.9	96.8	93.0	96.2	98.6 ^b	96.9
RSM	85.4 ^c	98.2 ^b	98.6 ^{a,b}	98.5 ^b	85.2 ^c	98.6 ^{b,c}	79.9 ^c	96.7 ^c	98.6 ^{b,c}	99.5 ^a	98.6 ^b	99.1	97.1	93.3	96.7	99.1 ^a	97.5
HBM	88.8 ^a	98.1 ^b	98.2 ^b	98.4 ^b	88.4 ^a	98.3 ^c	85.6 ^a	97.6 ^{a,b}	98.4 ^c	99.1 ^b	98.5 ^b	98.9	96.9	94.4	96.6	98.7 ^b	97.0
Means of the diets for the one-way ANOVA model																	
MG-0	84.0 ^c	98.4 ^b	98.7 ^{a,b,c}	98.9 ^{b,c}	84.0 ^d	98.7 ^{b,c}	78.6 ^d	97.7 ^b	98.8 ^{b,c,d}	99.5 ^a	99.1 ^{b,c}	99.4	97.3 ^{a,b}	92.9 ^{b,c,d}	97.0 ^{b,c}	99.3 ^{a,b}	97.6 ^{a,b,c}
MG-S	85.5 ^{b,c}	98.5 ^b	98.9 ^{a,b}	99.0 ^{a,b}	85.5 ^{b,c,d}	98.8 ^b	80.6 ^{c,d}	97.9 ^{a,b}	98.9 ^b	99.5 ^a	99.2 ^{a,b}	99.5	97.5 ^{a,b}	93.1 ^{b,c,d}	97.3 ^{a,b}	99.3 ^{a,b}	97.6 ^{a,b,c}
PP-0	90.3 ^a	99.1 ^a	99.3 ^a	99.4 ^a	90.0 ^a	99.2 ^a	87.4 ^a	98.7 ^a	99.4 ^a	99.6 ^a	99.5 ^a	99.7	98.3 ^a	95.6 ^a	98.2 ^a	99.5 ^a	98.4 ^a
PP-S	86.3 ^{b,c}	98.3 ^b	98.6 ^{b,c}	99.0 ^{a,b}	86.4 ^{a,b,c,d}	98.7 ^{b,c}	80.8 ^{b,c,d}	97.7 ^b	98.7 ^{b,c,d}	99.5 ^a	99.1 ^{a,b,c}	99.4	96.9 ^b	92.7 ^{c,d}	96.6 ^{b,c,d}	99.2 ^{a,b}	97.1 ^{b,c}
SF-0	86.9 ^{a,b,c}	97.8 ^c	98.2 ^c	97.8 ^e	86.6 ^{a,b,c,d}	98.1 ^{de}	82.9 ^{a,b,c,d}	97.3 ^{b,c}	98.1 ^{e,f}	98.9 ^{b,c}	97.8 ^f	98.6	96.7 ^b	93.3 ^{b,c,d}	96.4 ^{b,c,d}	98.7 ^c	97.1 ^{b,c}
SF-S	84.9 ^c	97.6 ^c	98.2 ^c	97.6 ^e	84.6 ^{c,d}	98.0 ^e	79.8 ^d	97.0 ^{b,c}	97.9 ^f	98.8 ^c	97.7 ^f	99.2	96.9 ^b	92.7 ^{c,d}	96.0 ^{c,d}	98.6 ^c	96.8 ^c
RS-0	86.9 ^{a,b,c}	98.5 ^b	98.8 ^{a,b}	98.7 ^{b,c,d}	86.7 ^{a,b,c,d}	98.7 ^{b,c}	82.0 ^{b,c,d}	97.0 ^{b,c}	98.9 ^{b,c}	99.5 ^a	98.8 ^{b,c,d,e}	99.2	97.6 ^{a,b}	93.5 ^{b,c,d}	97.3 ^{a,b}	99.3 ^{a,b}	97.8 ^{a,b}
RS-S	84.0 ^c	98.0 ^{b,c}	98.3 ^{b,c}	98.3 ^d	83.7 ^d	98.4 ^{b,c,d}	77.8 ^d	96.4 ^c	98.4 ^{c,d,e}	99.4 ^a	98.4 ^e	98.9	96.6 ^b	93.1 ^{b,c,d}	96.1 ^{c,d}	99.0 ^b	97.2 ^{b,c}
HB-0	88.8 ^{a,b}	98.1 ^{b,c}	98.3 ^{b,c}	98.4 ^{c,d}	88.3 ^{a,b,c}	98.4 ^{c,d}	85.4 ^{a,b,c}	97.7 ^b	98.5 ^{b,c,d,e}	99.1 ^b	98.5 ^{d,e}	98.9	97.0 ^b	94.5 ^{a,b}	96.6 ^{b,c,d}	98.7 ^c	97.1 ^{b,c}
HB-S	88.8 ^{a,b}	98.0 ^{b,c}	98.1 ^c	98.3 ^d	88.6 ^{a,b}	98.3 ^d	85.8 ^{a,b}	97.6 ^b	98.4 ^{d,e,f}	99.0 ^{b,c}	98.4 ^e	98.9	96.9 ^b	94.2 ^{a,b,c}	96.6 ^{b,c,d}	98.6 ^c	96.9 ^c
WG-S	85.4 ^{b,c}	98.3 ^b	98.8 ^{a,b}	98.6 ^{b,c,d}	84.3 ^d	98.7 ^{b,c}	79.1 ^d	97.5 ^b	98.7 ^{b,c,d}	99.5 ^a	98.7 ^{c,d,e}	99.4	97.1 ^b	93.0 ^{b,c,d}	96.6 ^{b,c,d}	99.3 ^{a,b}	97.6 ^{a,b,c}
SBM	84.4 ^c	98.0 ^{b,c}	98.5 ^{b,c}	98.8 ^{b,c,d}	84.0 ^d	98.7 ^{b,c}	77.8 ^d	97.2 ^{b,c}	98.5 ^{b,c,d,e}	99.4 ^a	98.9 ^{b,c,d}	99.4	96.6 ^b	92.1 ^d	95.8 ^d	99.2 ^b	96.9 ^{b,c}

MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.

^{a,b,c,d,e,f} Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).

throughout the digestive tract (DI1 and DI2). However, no significant effects of saponins were observed in the two distal sections.

Chyme bile salt concentration in the proximal intestinal sections depended on basal diet, and in MI was highest for MG-0, significantly higher than SF-0 and HB-0. The other basal diets, diet PP-0 and RS-0, showed intermediate bile salt concentrations in MI. This pattern was not clearly reflected in the other intestinal sections.

Saponin supplementation elevated trypsin activity in content from the DI1 and DI2 regions in fish fed PPC. For the other basal diets, there was no significant effect in these sections. No significant saponin effects were seen in the more proximal sections for any of the basal diets. The basal diets, however, differed regarding effects on trypsin activity in content from DI1 and DI2. In DI1, fish fed SF-0 and HB-0 had lower trypsin activity than fish fed MG-0. In the very distal section, DI2, trypsin activity was highest for fish fed PP-0 and significantly higher than for fish fed MG-0, SF-0 and HB-0. Fish fed RS-0 had intermediate values.

The reference SBM diet also induced high trypsin activity in DI1, an activity that was significantly higher than values from other diets except the PP-S diet, which showed an even higher value. Also in the DI2, the highest trypsin activities were recorded for the PP-S diet. However, the SBM was almost as high and not significantly different from the value observed

for the PP-S diet. The values observed for PP-S and SBM in the DI2 were several times higher than the lowest value which was observed for the SF-S diet. For all other diets, there was a great drop in trypsin activity between DI1 and DI2.

Brush-border membrane leucine aminopeptidase activity. Activities of LAP in the DI are presented in Table 13, expressed as specific activity and as capacity (activity per kg fish). In the PI and MI sections, no significant effect of saponin supplementation was observed for either the specific activity (*P* value/pooled SEM, 0.73/44 for PI and 0.28/25 for MI) or capacity (*P* value/pooled SEM, 0.60/39 for PI and 0.09/2 for MI). In DI, saponin inclusion reduced activity expressed both ways for all basal diets, but not to the same degree. The reduction was significant for fish fed PPC and RSM; most pronounced for PPC with a reduction of 74%.

The basal diets did not differ significantly regarding effects on LAP activity in the PI and MI although there was a trend towards higher activity in the MI in fish fed HBM. In the DI, basal diet had significantly different effects on both capacity and specific activity of LAP. The lowest activity was observed in fish fed PPC, intermediate when fed MG, SFM and RSM, and highest when fed HBM.

Organo-somatic index. Saponin numerically reduced OSI of the DI for fish fed all basal diets except MG (Table 14). However, fish fed the PPC were the only ones showing

Table 9. Apparent digestibility coefficients (%) for selected dietary minerals for Atlantic salmon fed the experimental diets

	Ash	Na	Ca	Mg	Cu	Fe	Mn	Zn	P
Two-way ANOVA model									
<i>P</i> (model)	<0.0001	<0.0001	0.32	<0.0001	<0.0001	0.02	0.002	0.0004	0.04
Pooled SEM	3.0	4.2	21.2	19.5	1.7	7.0	1.9	1.7	2.3
<i>P</i> values effect tests in the two-way ANOVA model									
Saponin	0.0001	<0.0001	0.19	0.001	0.001	0.98	0.32	0.53	0.71
Basal diet	0.0003	<0.0001	0.29	<0.0001	<0.0001	0.01	0.0004	0.0002	0.002
Interaction	<0.0001	<0.0001	0.41	0.04	0.04	0.05	0.05	0.003	0.03
Marginal means for the two-way ANOVA model									
Saponin supplementation									
No saponin	-4.1 ^a	42.4 ^a	-13.1	-263.6 ^a	27.2 ^b	2.7	-0.6	18.9	26.8
With saponin	-15.8 ^b	17.6 ^b	-30.8	-314.2 ^b	32.4 ^a	2.6	-1.9	19.6	26.3
Basal diet									
MG	-9.7 ^b	55.6 ^a	-24.2	-350.2 ^c	37.1 ^a	3.6 ^b	3.3 ^a	24.6 ^a	32.4 ^a
PPC	-17.2 ^c	-13.2 ^d	-29.3	-371.6 ^c	36.0 ^a	0.0 ^b	0.3 ^a	17.2 ^b	29.6 ^a
SFM	-9.3 ^b	34.3 ^b	-29.3	-274.8 ^a	15.5 ^c	-13.1 ^b	-8.4 ^b	13.3 ^c	21.8 ^b
RSM	-16.9 ^c	21.9 ^c	-34.3	-276.0 ^b	28.6 ^b	1.8 ^b	-4.8 ^b	17.2 ^b	20.8 ^b
HBM	3.3 ^a	51.3 ^a	7.2	-231.7 ^a	31.6 ^b	21.1 ^a	3.3 ^a	23.9 ^a	28.2 ^a
Means of the diets for the one-way ANOVA model									
MG-0	-11.3 ^c	51.6 ^{b,c}	-28.3	-377.4 ^c	31.8 ^{c,d}	-3.5 ^{c,d}	1.1 ^{b,c,d}	21.2 ^d	29.6 ^{b,c}
MG-S	-8.1 ^{b,c}	59.7 ^{a,b}	-20.0	-323.3 ^{b,c}	42.4 ^b	10.7 ^{b,c}	5.5 ^b	28.1 ^{b,c}	35.1 ^{a,b}
PP-0	6.3 ^a	34.7 ^{d,e}	-16.1	-246.6 ^a	33.0 ^{c,d}	2.6 ^{b,c}	2.8 ^{b,c,d}	14.9 ^{e,f}	30.6 ^{a,b,c}
PP-S	-40.7 ^e	-61.0 ^g	-42.6	-496.6 ^d	39.0 ^b	-2.6 ^{c,d}	-2.1 ^{d,e}	19.5 ^{d,e}	28.6 ^{b,c}
SF-0	-9.6 ^c	40.8 ^{c,d}	-33.9	-233.5 ^a	11.8 ^f	-22.7 ^d	-10.3 ^f	12.7 ^f	20.7 ^{d,e}
SF-S	-9.1 ^c	27.7 ^e	-24.8	-206.0 ^a	19.2 ^e	-3.5 ^{c,d}	-6.4 ^{e,f}	13.9 ^f	23.0 ^{c,d}
RS-0	-7.2 ^{b,c}	32.6 ^{d,e}	-25.5	-247.0 ^a	29.5 ^{c,d}	2.7 ^{b,c}	-1.3 ^{c,d,e}	22.8 ^{c,d}	26.3 ^{c,d}
RS-S	-26.6 ^d	11.1 ^f	-43.1	-305.0 ^b	27.7 ^d	0.8 ^c	-8.3 ^f	11.7 ^f	15.2 ^e
HB-0	1.1 ^{a,b}	52.2 ^{b,c}	38.1	-223.6 ^a	29.7 ^{c,d}	34.7 ^a	4.7 ^{b,c}	23.0 ^{c,d}	27.0 ^{c,d}
HB-S	5.5 ^a	50.4 ^{b,c}	-23.7	-239.9 ^a	33.6 ^c	7.6 ^{b,c}	1.9 ^{b,c,d}	24.8 ^{c,d}	29.5 ^{b,c}
WG-S	3.7 ^a	71.1 ^a	-15.0	-229.9 ^a	54.3 ^a	23.8 ^{a,b}	13.3 ^a	36.4 ^a	37.5 ^a
SBM	-7.2 ^{b,c}	23.9 ^e	-31.4	-306.6 ^b	40.3 ^b	19.0 ^{a,b,c}	-1.3 ^{c,d,e}	30.7 ^b	25.6 ^{c,d}

MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.
^{a,b,c,d,e,f,g} Mean values within a column with unlike superscript letters were significantly different (*P* < 0.05).

significant reduction. There was no significant saponin–basal diet interaction.

Significant basal diet differences were discernible for the MI, DI and liver OSI. The OSI for MI in fish fed the HBM was significantly higher than the rest of the basal diets, which did not differ significantly from each other. The DI OSI of fish fed HBM was the highest and was significantly different from PPC and SFM but not RSM and SFM. Fish fed the MG and HBM had the highest liver OSI while fish fed the SFM and PPC diets had the lowest. Intermediate values were observed for fish fed RSM.

OSI for the ST, spleen and kidney showed no apparent effects of either saponin inclusion or composition of the basal diet.

Histology. The scores from the visual analogue scale evaluation of the DI histology are presented in Table 15. In fish fed PPC, saponin inclusion seemed to affect all evaluated histological variables except vacuole size variation. These fish showed higher degrees of mucosal fold fusion, increased width and degree of cellular infiltration of the lamina propria and submucosa, abnormal nucleus position and reduced vacuolisation in enterocytes, and increased numbers of goblet cells. For other basal diets, no saponin effects were

observed except for significantly shorter mucosal folds and a wider lamina propria in fish fed RSM and higher numbers of goblet cells in fish fed SFM.

Comparison of the results for the unsupplemented diets did not reveal clear differences between the basal diets for any of the histological parameters.

Plasma parameters. No significant effects of either the saponin inclusion or basal diet were apparent for the plasma indicators of lipid metabolism analysed that included cholesterol, TAG and NEFA (results not presented). Saponin inclusion to most basal diets showed a negative trend on glucose levels ($P=0.08$; pooled SEM = 0.2; two-way ANOVA marginal means, without saponin = 5.2, with saponin = 5.0 mmol/l) with the exception of the SFM.

Summary of saponin effects. The effects of saponin supplementation were as follows: For all basal diets, there seemed to be negative effects on AD of lipid and individual fatty acids and on FDM, but to different degrees. For other variables, however, the saponin effects were dependent on the composition of the basal diet, and the predominant picture showed major negative effects for PPC on all, except FER and nutrient retention. For the other basal diets, the effects were minor. There was a trend towards positive effect of saponins

Table 10. Retention of crude protein (CP) and energy in Atlantic salmon fed the experimental diets

	Retention % of ingested		Retention % of digested	
	CP	Energy	CP	Energy
Two-way ANOVA model				
P (model)	0.03	0.06	0.09	0.32
Pooled SEM	2.4	3.4	2.8	4.1
P values effect tests in the two-way ANOVA model				
Saponin	0.22	0.54	0.19	0.60
Basal diet	0.01	0.02	0.03	0.16
Interaction	0.57	0.36	0.63	0.50
Marginal means for the two-way ANOVA model				
Saponin supplementation				
No saponin	49.0	54.0	56.9	64.0
With saponin	51.0	52.6	59.4	62.6
Basal diet				
MG	50.1 ^a	59.4	56.1	66.3
PPC	52.4 ^a	55.0	60.8	64.6
SFM	52.8 ^a	52.9	61.5	64.2
RSM	42.4 ^b	44.7	51.7	56.0
HBM	52.4 ^a	54.5	60.5	65.2
Means of the diets for the one-way ANOVA model				
MG-0	48.1 ^{a,b,c}	55.9 ^{a,b,c}	54.3 ^{b,c,d}	62.8
MG-S	52.0 ^{a,b}	62.8 ^a	57.9 ^{a,b,c,d}	69.7
PP-0	51.5 ^{a,b}	58.3 ^{a,b}	59.0 ^{a,b,c,d}	67.6
PP-S	53.4 ^{a,b}	51.8 ^{b,c,d}	62.7 ^{a,b}	61.6
SF-0	53.9 ^a	52.9 ^{a,b,c,d}	62.7 ^{a,b}	64.4
SF-S	51.8 ^{a,b}	52.8 ^{a,b,c,d}	60.2 ^{a,b,c}	63.9
RS-0	42.0 ^c	45.6 ^{c,d}	50.9 ^d	56.4
RS-S	42.8 ^c	43.8 ^d	52.4 ^{c,d}	55.6
HB-0	49.6 ^{a,b,c}	57.2 ^{a,b}	57.4 ^{a,b,c,d}	68.5
HB-S	55.2 ^a	51.8 ^{b,c,d}	63.5 ^a	61.9
WG-S	55.1 ^a	62.7 ^a	59.6 ^{a,b,c,d}	68.4
SBM	45.8 ^{b,c}	53.2 ^{a,b,c,d}	52.7 ^{c,d}	61.7

MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.

^{a,b,c,d}Mean values within a column with unlike superscript letters were significantly different ($P<0.05$).

on feed intake, significant for SFM, with corresponding results for body weight and TGC. The RSM showed reduced AD of all nutrients, minerals included, when saponins had been added; which was also the case for mineral AD in SFM. Supplementation of saponins resulted in increased AD of amino acids in MG; reduced brush-border LAP activity in the DI for RSM, with a similar trend for MG, SFM and HBM. Only the PPC showed negative effects of saponin supplementation on growth, trypsin activity in chyme of the DI, and on morphology and histology of the DI.

Summary regarding the basal diets. The basal diet with PPC, as indicated by the PP-0 diet, generally showed high AD values for all the nutrition variables, with only the AD of Cys and Trp not ranking highest. Regarding the gut variables, however, PP-0 did less well and some of the variables indicated tendencies towards enteritis-like symptoms. The HB-0 diet overall showed somewhat lower nutritional values than the PP-0, with lower values for protein and amino acid digestibility and protein retention. However, the HB-0 generally showed the best values for many gut variables, but with somewhat reduced bile salt concentration in the intestinal content. The MG-0 diet was intermediate for both the nutrient and gut variables characterised by somewhat lower feed intake and growth, lower AD of Trp, lipid and ash, and CP retention, and some indication of gut reactions. The SF-0 diet showed relatively lower values for all nutrition variables except for

the AD of Trp and CP retention, but variables regarding the gut condition were quite good. Overall, the RS-0 showed the lowest values for the AD variables, and the values for the gut variables also ranked low, but somewhat better than those of the PP-0 diet.

Discussion

Effects of saponin supplementation

The negative effect of saponins on lipid digestibility was most probably related to their ability to form insoluble complexes with 3-β-hydroxysteroids. They also interact with and form large, mixed micelles with bile salts and cholesterol, accounting for increased bile salt and cholesterol excretion^(11,23). The association with bile salts may have implications for lipid digestion and uptake by the intestinal enterocytes. Both a direct effect on lipid emulsification and micelle formation, and an indirect effect secondary to effects on the body pool of bile salts could reduce lipid digestibility.

The observation that saponin supplementation affected feed intake to a variable degree is in agreement with results of other studies. Bureau *et al.*⁽¹²⁾ observed reduced feed intake as an effect of an SBM-derived saponin-rich alcohol extract when fed to Chinook salmon, but not when fed to rainbow trout. Twibell & Wilson⁽²⁴⁾ also observed negative saponin

Table 11. Retention of absorbed essential amino acids and cystein in Atlantic salmon fed the experimental diets (%)

	Lys	Met	Cys	Arg	His	Ile	Leu	Thr	Phe	Val
Two-way ANOVA model										
<i>P</i> (model)	0.10	0.04	<0.0001	0.01	0.50	0.57	0.004	0.66	0.11	0.50
Pooled SEM	3.7	3.6	4.5	2.6	3.0	3.9	2.8	3.7	3.0	4.3
<i>P</i> values effect tests in the two-way ANOVA model										
Saponin	0.33	0.23	0.67	0.14	0.32	0.20	0.23	0.45	0.51	0.58
Basal diet	0.02	0.01	<0.0001	0.002	0.46	0.50	0.001	0.38	0.03	0.20
Interaction	0.70	0.53	0.32	0.48	0.45	0.66	0.60	0.82	0.70	0.87
Marginal means for the two-way ANOVA model										
Saponin supplementation										
No saponin	57.4	55.1	72.5	44.7	48.3	49.7	44.2	57.1	46.1	50.3
With saponin	59.8	58.1	73.8	47.3	50.3	53.0	46.4	59.0	47.4	51.8
Basal diet										
MG	67.5	57.8 ^b	54.3 ^b	55.8 ^a	51.2	52.7	33.0 ^b	58.6	41.5	53.3
PPC	55.0	50.0 ^b	80.2 ^a	41.5 ^b	49.3	51.3	49.0 ^a	60.2	45.8	51.9
SFM	58.2	55.6 ^b	81.9 ^a	44.6 ^b	48.4	51.0	49.3 ^a	59.1	46.2	51.3
RSM	52.9	53.1 ^b	60.4 ^b	44.6 ^b	46.3	47.4	45.2 ^a	53.1	46.5	44.0
HBM	59.3	66.4 ^a	89.0 ^a	43.4 ^b	51.4	54.4	50.0 ^a	59.3	53.8	54.7
Means of the diets for the one-way ANOVA model										
MG-0	66.0 ^{a,b}	55.7 ^{b,c,d}	54.6 ^{c,d}	53.2 ^{a,b}	50.5	50.9	32.2 ^d	57.3	41.0	51.8
MG-S	69.0 ^a	60.0 ^{a,b,c,d}	54.0 ^{c,d}	58.3 ^a	52.0	54.5	33.7 ^{c,d}	59.9	42.0	54.7
PP-0	55.4 ^{b,c,d}	50.5 ^{c,d}	82.9 ^{a,b}	41.8 ^c	49.6	51.5	49.1 ^{a,b}	60.6	46.5	53.4
PP-S	54.5 ^{b,c,d}	49.6 ^{c,d}	77.5 ^b	41.1 ^c	49.0	51.0	48.9 ^{a,b}	59.8	45.0	50.4
SF-0	58.9 ^{a,b,c,d}	57.0 ^{b,c,d}	79.7 ^b	45.3 ^{b,c}	49.4	51.7	49.7 ^{a,b}	60.1	46.6	51.7
SF-S	57.4 ^{a,b,c,d}	54.2 ^{b,c,d}	45.5 ^d	43.9 ^c	47.4	50.3	48.9 ^{a,b}	58.1	45.7	50.9
RS-0	48.8 ^d	48.7 ^d	58.4 ^{c,d}	41.3 ^c	41.7	43.1	41.5 ^{b,c}	50.2	43.2	42.2
RS-S	57.0 ^{b,c,d}	57.4 ^{b,c,d}	62.5 ^c	47.9 ^{b,c}	50.8	51.7	48.9 ^{a,b}	56.1	49.8	45.9
HB-0	57.6 ^{a,b,c,d}	63.8 ^{a,b}	82.7 ^{a,b}	41.7 ^c	50.3	51.3	48.4 ^{a,b}	57.6	53.1	52.2
HB-S	61.0 ^{a,b,c}	69.1 ^a	95.2 ^a	45.1 ^{b,c}	52.4	57.6	51.6 ^a	61.0	54.5	57.2
WG-S	65.1 ^{a,b}	61.5 ^{a,b,c}	77.1 ^b	52.3 ^{a,b}	48.9	49.8	45.5 ^{a,b}	62.7	41.0	52.2
SBM	52.8 ^{c,d}	56.0 ^{b,c,d}	84.2 ^{a,b}	43.2 ^c	46.5	47.9	45.7 ^{a,b}	56.8	43.3	50.6

MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.
^{a,b,c,d} Mean values within a column with unlike superscript letters were significantly different (*P*<0.05).

Table 12. Bile salt concentration and trypsin activity in intestinal content from the experimental fish

	Bile salts (mg/g DM)					Trypsin (U/mg DM)				
	PI1	PI2	MI	DI1	DI2	PI1	PI2	MI	DI1	DI2
Two-way ANOVA model										
P (model)	0.02	0.11	0.01	0.21	0.23	0.51	0.19	0.20	<0.0001	<0.0001
Pooled SEM	15	13	9	7	2	46	23	27	9	7
<i>P</i> values effect tests in the two-way ANOVA model										
Saponin	0.24	0.29	0.003	0.02	0.02	0.75	0.46	0.52	0.59	0.0003
Basal diet	0.17	0.27	0.03	0.65	0.32	0.48	0.45	0.06	<0.0001	<0.0001
Interaction	0.01	0.06	0.02	0.40	0.86	0.36	0.08	0.71	0.0002	<0.0001
Marginal means for the two-way ANOVA model										
Saponin supplementation										
No saponin	146	131	116 ^a	45	14	248	183	175	64	18 ^b
With saponin	135	122	94 ^b	33	9	238	172	164	61	35 ^a
Basal diet										
MG	148	127	126 ^a	44	13	282	203	230	54 ^{b,c}	11 ^b
PPC	125	109	89 ^b	38	14	224	170	160	106 ^a	87 ^a
SFM	159	138	102 ^b	40	10	261	180	143	48 ^c	9 ^b
RSM	126	125	102 ^b	38	10	248	176	167	71 ^b	17 ^b
HBM	145	132	105 ^b	33	11	201	160	148	35 ^c	10 ^b
Means of the diets for the one-way ANOVA model										
MG-0	138 ^{a,b,c}	117	139 ^a	54	16 ^{a,b}	242	190	254	87 ^c	13 ^c
MG-S	159 ^{a,b}	137	114 ^{a,b,c}	33	10 ^{b,c,d}	321	216	205	21 ^f	8 ^c
PP-0	177 ^a	137	124 ^{a,b}	48	15 ^{b,c}	277	188	176	70 ^{c,d}	37 ^b
PP-S	72 ^d	80	54 ^d	28	13 ^{b,c,d}	170	151	144	141 ^a	136 ^a
SF-0	156 ^{a,b}	147	108 ^{b,c}	49	13 ^{b,c,d}	260	191	144	58 ^{d,e}	12 ^c
SF-S	161 ^{a,b}	128	97 ^{b,c}	32	7 ^d	262	170	142	38 ^{e,f}	5 ^c
RS-0	129 ^{a,b,c}	130	110 ^{a,b,c}	40	12 ^{b,c,d}	243	216	156	73 ^{c,d}	21 ^{b,c}
RS-S	122 ^{b,c}	120	95 ^{b,c}	36	8 ^{c,d}	254	135	179	70 ^{c,d}	13 ^c
HB-0	131 ^{a,b,c}	121	100 ^{b,c}	33	12 ^{b,c,d}	170	131	147	34 ^{e,f}	8 ^c
HB-S	159 ^{a,b}	143	109 ^{a,b,c}	34	9 ^{b,c,d}	231	189	149	37 ^{e,f}	11 ^c
WG-S	103 ^{c,d}	103	94 ^{b,c}	42	22 ^a	174	133	197	65 ^{c,d}	22 ^{b,c}
SBM	117 ^{b,c,d}	100	88 ^c	31	11 ^{b,c,d}	280	227	187	115 ^b	129 ^a

PI, pyloric intestine; MI, mid-intestine; DI, distal intestine; MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.
^{a,b,c,d,e,f} Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).

effects on juvenile channel catfish (*Ictalurus punctatus*). However, the results of Knudsen *et al.*^(6,7) showed no effects on feed intake when supplemented to diets with lupin meal. It may be suggested that the varying effects of saponin supplementation are due to varying doses of saponins in the diets, being positive at low levels and negative at higher levels⁽²⁴⁾. However, our chances of evaluating this suggestion are hampered by lack of solid information on saponin content of various plant ingredients. No standardised methods for quantification of saponins are available, and reliability and reproducibility of the methods in use are not satisfactory.

The mechanism behind the negative effects of saponin supplementation to the diets with PPC and to some degree to the diet with RSM on nutrient digestibility is not clear, but interactions with one or more of the many ANF are likely. Several ANF are present in peas including protease inhibitors, phytic acid, oligosaccharides, lectins, tannins and saponins^(25,26). Many of the same types are found in RS which are characterised by a very high fibre and phytate content, presence of condensed tannins, a potent trypsin inhibitor, anthocyanidins, glucosinolates and lectins^(27–29). All, possibly except the glucosinolates, have the potential to interfere with digestive processes. However, hardly any information exists regarding interactions between antinutrients and effects on digestion.

Questions regarding their interaction with saponins cannot be answered without further investigations.

The negative effect of saponin inclusion on AD of Na in the PPC, SFM and RSM diets, and for Mg in PPC and RSM basal diets may also have been the result of complex formation between the ANF of these protein sources and minerals in the intestinal lumen, effects on intestinal uptake mechanisms, or on mineral recirculation between the intestine and the body. Na uptake in marine fish is a key in osmoregulation and has been demonstrated to occur along the entire GIT but more significantly in the anterior segments (reviewed by Grosell⁽³⁰⁾). However, the magnitude of the alterations, and the fact that many of the values were negative, indicate that altered drinking behaviour may have been a major factor in these effects. The low FDM in the PP-S and the SBM diets suggests that the fish fed these diets had diarrhoea, which could, at least partly, explain the lower Na AD. The difference in AD of Na between the PP-0 and PP-S diets was remarkable (35% compared to –61%), and supports this suggestion. However, the fact that the Na AD in fish fed the PP-0 diet, which did not show low FDM, was not different from that of SBM, indicates that other mechanisms are involved. The positive effect of saponin supplementation on AD of Cu, Fe, Mn, Zn and P for the MG and SFM diets may be related to

Table 13. Leucine aminopeptidase activity in intestinal tissue homogenates from pyloric, mid and distal intestine from Atlantic salmon fed the experimental diets

	Total capacity (mm/kg BW (mmol/h per kg fish weight))			Specific activity, mm/g protein (μ mol/h per mg tissue protein)		
	Pyloric	Mid	Distal	Pyloric	Mid	Distal
Two-way ANOVA model						
<i>P</i> (model)	0.60	0.09	0.0003	0.73	0.28	<0.0001
Pooled SEM	39	2	9	44	25	26
<i>P</i> values effect tests in the two-way ANOVA model						
Saponin	0.72	0.46	0.0004	0.61	0.90	0.0002
Basal diet	0.57	0.02	0.0001	0.39	0.58	0.0001
Interaction	0.69	0.49	0.29	0.49	0.43	0.01
Marginal means for the two-way ANOVA model						
Saponin supplementation						
No saponin	275	17	95 ^a	357	223	409 ^a
With saponin	266	18	66 ^b	342	225	314 ^b
Basal diet						
MG	257	17	77 ^b	322	243	352 ^c
PPC	276	19	39 ^c	351	234	227 ^d
SFM	247	16	85 ^b	329	234	376 ^{b,c}
RSM	265	13	91 ^{a,b}	367	231	414 ^{a,b}
HBM	309	21	110 ^a	378	178	437 ^a
Means of the diets for the one-way ANOVA model						
MG-0	261	14	83 ^{b,c}	342	238	369 ^{b,c}
MG-S	253	19	70 ^{b,c,d}	303	249	336 ^{b,c}
PP-0	312	19	63 ^{c,d}	402	240	360 ^{b,c}
PP-S	241	19	13 ^e	299	229	94 ^d
SF-0	225	16	396 ^{a,b}	303	216	393 ^{a,b,c}
SF-S	268	17	74 ^{b,c,d}	355	252	358 ^{b,c}
RS-0	271	15	110 ^a	360	258	458 ^a
RS-S	258	12	71 ^{b,c,d}	374	204	371 ^{b,c}
HB-0	307	20	122 ^a	376	165	465 ^a
HB-S	310	23	98 ^{a,b}	379	192	409 ^{a,b}
WG-S	224	16	53 ^d	302	209	311 ^c
SBM	288	14	20 ^e	364	189	129 ^d

BW, body weight; MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.
^{a,b,c,d,e} Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).

Table 14. Organo-somatic indices (SI) from the Atlantic salmon fed the experimental diets*

	Stomach SI	Intestine SI			Liver SI	Spleen SI	Kidney SI	
		Pyloric	Mid	Distal			Trunk	Head
Two-way ANOVA model								
<i>P</i> (model)	0.35	0.49	0.0004	0.01	0.02	0.19	0.94	0.44
Pooled SEM	0.02	0.13	0.01	0.03	0.05	0.01	0.02	0.01
<i>P</i> values effect tests in the two-way ANOVA model								
Saponin	0.32	0.80	0.32	0.05	0.75	0.54	0.50	0.25
Basal diet	0.12	0.14	<0.0001	0.003	0.004	0.05	0.83	0.51
Interaction	0.91	0.99	0.17	0.20	0.64	0.73	0.88	0.36
Marginal means for the two-way ANOVA model								
Saponin supplementation								
No saponin	0.48	1.93	0.22	0.51 ^a	1.22	0.08	0.55	0.14
With saponin	0.49	1.95	0.22	0.47 ^b	1.21	0.08	0.56	0.14
Basal diet								
MG	0.47	2.16	0.20 ^b	0.48 ^b	1.28 ^{a,b}	0.10	0.54	0.14
PPC	0.49	1.87	0.21 ^b	0.40 ^c	1.16 ^c	0.08	0.56	0.13
SFM	0.47	1.86	0.21 ^b	0.51 ^b	1.11 ^c	0.07	0.55	0.14
RSM	0.46	1.81	0.19 ^b	0.49 ^b	1.17 ^{b,c}	0.08	0.56	0.15
HBM	0.52	2.00	0.29 ^a	0.57 ^a	1.35 ^a	0.08	0.55	0.13
Means of the diets for the one-way ANOVA model								
MG-0	0.47	2.16	0.19 ^b	0.46 ^{b,c}	1.27 ^{a,b,c,d}	0.10	0.53	0.137
MG-S	0.47	2.15	0.21 ^b	0.49 ^{a,b,c}	1.29 ^{a,b,c}	0.10	0.56	0.149
PP-0	0.49	1.85	0.20 ^b	0.46 ^{b,c}	1.19 ^{b,c,d}	0.08	0.57	0.135
PP-S	0.49	1.89	0.22 ^b	0.35 ^d	1.13 ^{d,c}	0.09	0.56	0.132
SF-0	0.46	1.88	0.22 ^b	0.52 ^{a,b}	1.13 ^{d,c}	0.07	0.55	0.142
SF-S	0.48	1.85	0.20 ^b	0.50 ^{a,b,c}	1.09 ^d	0.08	0.55	0.136
RS-0	0.45	1.79	0.20 ^b	0.54 ^{a,b}	1.19 ^{b,c,d}	0.08	0.56	0.159
RS-S	0.47	1.82	0.19 ^b	0.45 ^{b,c}	1.16 ^{b,c,d}	0.08	0.57	0.133
HB-0	0.50	1.96	0.28 ^a	0.58 ^a	1.32 ^{a,b}	0.08	0.54	0.136
HB-S	0.54	2.05	0.30 ^a	0.57 ^a	1.39 ^a	0.09	0.56	0.128
WG-S	0.43	2.06	0.22 ^b	0.42 ^{c,d}	1.22 ^{b,c,d}	0.10	0.54	0.140
SBM	0.47	2.02	0.21 ^b	0.41 ^{c,d}	1.18 ^{b,c,d}	0.09	0.55	0.138

MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.

^{a,b,c,d} Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).

* Organ weight as a percentage of fish body weight.

an increase in permeability of the intestine^(7,10,31) which has been an observed effect of soyasaponins.

A mechanism for the positive effect of saponins in the MG diet on AD of amino acids may be related to stimulatory effects of saponins on trypsin activity in the intestinal chyme. Although not significant, there was a numerical increase in the activity of trypsin in PI1 of 30%, and the activity was elevated also in PI2. Elevated trypsin activities were observed also for HBM, but without the beneficial effect on amino acid AD. The explanation for the difference is possibly related to ANF in the HBM that may have hindered a beneficial effect.

The negative effects of saponins on LAP activity of the distal intestinal brush border of fish for all basal diets, significant for PPC and RSM, are in agreement with results of an earlier study with Atlantic salmon⁽³²⁾ in which effects of the same saponin preparation, supplemented at 1 and 2 g per kg in a diet with 30% soya protein concentrate were investigated. The explanation for this reduction in enzyme activity can only be speculative as current knowledge is weak. Saponins have many recorded biological and pharmacological activities and a large number of them are ascribed to their cell membrane effects⁽⁷⁾ which may affect BBM enzymes directly or indirectly by increasing cell proliferation. The effects on LAP may have been attributed to a loss of differentiated intestinal epithelial

cells and their subsequent replacement with immature cells expressing less BBM enzymes⁽¹⁹⁾.

Supplementation of saponins to the basal diet with PPC caused effects in addition to those shared by one or more of the other basal diets: negative effect on growth, elevation of trypsin activity of chyme in the distal intestinal compartments and enteritis in the DI. These clear disruptive effects were very similar to the symptoms typical for soyabean-induced enteritis observed also in fish fed the SBM. In a recent study, similar histological observations were seen in fish fed a diet with somewhat higher PPC inclusion (35%) compared to the present work⁽²⁰⁾ and without saponin supplementation. This indicates that the saponin level in PPC could be high enough to induce enteritis when the dietary inclusion level is high or when combined with SBM or other saponin-containing ingredients. Whether saponins can cause these effects alone is a matter of discussion. There is a possibility that development of enteritis requires another alcohol-soluble component in addition to the saponins, a component that is present in soybeans, peas, lupins and possibly other legumes. This was also suggested by Knudsen *et al.*⁽⁷⁾ based on information gained from saponin supplementation of a diet with lupin kernel meal, using a 65% purified soyasaponin extract, which induced enteritis. The work of Knudsen *et al.*⁽⁷⁾ showed a dose-dependency of the saponin effects. In the present

Table 15. Details of the score-based evaluation of the intestinal histology of fish fed the experimental diets*

	Mucosal folds		Lamina propria		Submucosa		Enterocytes			Other cell types	
	Height	Fusion	Width	Cellularity infiltration	Width	Cellularity infiltration	Nuclear position	Vacuole size	Vacuole size variation	IEL numbers	Goblet cell numbers
Two-way ANOVA model											
<i>P</i> (model)	0.01	0.003	<0.0001	<0.0001	0.02	0.01	0.001	0.0002	0.13	0.45	0.001
Pooled SEM	0.5	0.7	0.3	0.4	0.4	0.5	0.5	0.3	0.1	0.8	0.6
<i>P</i> values effect tests in the two-way ANOVA model											
Saponin	0.01	0.003	<0.0001	0.0001	0.06	0.01	0.001	0.001	0.68	0.15	0.0003
Basal diet	0.01	0.01	<0.0001	<0.0001	0.01	0.01	0.003	0.001	0.14	0.86	0.01
Interaction	0.05	0.03	<0.0001	0.0002	0.08	0.14	0.004	0.002	0.11	0.27	0.01
Marginal means for the two-way ANOVA model											
Saponin supplementation											
No saponin	7.8 ^a	1.9 ^b	1.1 ^b	2.0 ^b	1.7	3.2 ^b	1.5 ^b	7.9 ^a	2.1	2.3	3.1 ^b
With saponin	6.6 ^b	3.5 ^a	2.6 ^a	3.5 ^a	2.3	4.2 ^a	3.0 ^a	7.1 ^b	2.0	3.1	5.1 ^a
Basal diet											
MG	7.6 ^{a,b}	2.7 ^b	1.3 ^b	2.4 ^{b,c}	2.0 ^b	3.9 ^b	2.0 ^{b,c}	7.6 ^a	1.5	2.4	4.1 ^{a,b}
PPC	5.9 ^c	4.8 ^a	4.1 ^a	5.2 ^a	3.2 ^a	5.0 ^a	3.9 ^a	6.2 ^b	1.7	3.1	5.5 ^a
SFM	7.8 ^a	2.0 ^b	1.1 ^b	1.7 ^c	1.6 ^b	3.3 ^b	1.7 ^b	7.9 ^a	1.9	2.7	3.7 ^{b,c}
RSM	6.6 ^{b,c}	2.4 ^b	1.7 ^b	2.7 ^b	1.9 ^b	3.7 ^b	2.5 ^{b,c}	7.7 ^a	2.9	3.1	4.7 ^{a,b}
HBM	8.1 ^a	1.5 ^b	1.1 ^b	1.6 ^c	1.4 ^b	2.7 ^b	1.2 ^c	7.9 ^a	2.2	2.5	2.5 ^c
Means of the diets for the one-way ANOVA model											
MG-0	7.5 ^a	2.4 ^b	1.2 ^c	2.1 ^{b,c}	1.8 ^b	3.4 ^{c,d}	2.0 ^{b,c}	7.4 ^a	1.8 ^{b,c}	3.0 ^{a,b,c}	3.7 ^{b,c,d,e}
MG-S	7.6 ^a	3.1 ^b	1.5 ^{b,c}	2.7 ^{b,c}	2.1 ^b	4.4 ^{b,c}	1.9 ^{b,c}	7.9 ^a	1.2 ^{b,c}	1.9 ^{b,c}	4.6 ^{b,c}
PP-0	7.5 ^a	2.4 ^b	1.4 ^{b,c}	2.6 ^{b,c}	2.1 ^b	3.7 ^{c,d}	1.4 ^c	7.6 ^a	1.6 ^{b,c}	1.7 ^{b,c}	3.0 ^{c,d,e}
PP-S	4.3 ^{b,c}	7.2 ^a	6.8 ^a	8.0 ^a	4.3 ^a	6.3 ^a	6.5 ^a	4.9 ^b	1.9 ^{b,c}	4.5 ^a	8.0 ^a
SF-0	7.9 ^a	1.7 ^b	0.8 ^c	1.4 ^c	1.6 ^b	3.2 ^{c,d}	1.3 ^c	8.1 ^a	2.8 ^{a,b}	2.2 ^{b,c}	2.4 ^{d,e}
SF-S	7.8 ^a	2.3 ^b	1.3 ^{b,c}	2.0 ^{b,c}	1.7 ^b	3.4 ^{c,d}	2.2 ^{b,c}	7.8 ^a	1.1 ^c	3.1 ^{a,b,c}	5.1 ^b
RS-0	7.4 ^a	1.5 ^b	1.1 ^c	2.0 ^{b,c}	1.7 ^b	3.2 ^{c,d}	1.7 ^{b,c}	8.2 ^a	2.2 ^{a,b}	2.6 ^{a,b,c}	4.4 ^{b,c,d}
RS-S	5.7 ^b	3.3 ^b	2.3 ^b	3.4 ^b	2.2 ^b	4.3 ^{b,c}	3.3 ^b	7.2 ^a	3.6 ^a	3.5 ^{a,b,c}	5.1 ^b
HB-0	8.3 ^a	1.5 ^b	1.1 ^c	1.7 ^c	1.6 ^b	2.9 ^{c,d}	1.0 ^c	8.1 ^a	2.2 ^{a,b,c}	2.1 ^{b,c}	2.1 ^e
HB-S	7.7 ^a	1.6 ^b	1.1 ^c	1.5 ^c	1.1 ^b	2.7 ^{c,d}	1.4 ^c	7.8 ^a	2.2 ^{a,b,c}	2.7 ^{a,b,c}	2.9 ^{c,d,e}
WG-S	7.4 ^a	2.6 ^b	1.5 ^{b,c}	2.4 ^{b,c}	1.0 ^b	2.4 ^d	2.6 ^{b,c}	7.3 ^a	1.5 ^{b,c}	1.4 ^c	5.1 ^b
SBM	3.8 ^c	7.0 ^a	6.7 ^a	7.7 ^a	3.9 ^a	5.8 ^{a,b}	6.6 ^a	5.0 ^b	2.0 ^{b,c}	3.7 ^{a,b}	8.2 ^a

IEL, intra-epithelial lymphocyte; MG, maize gluten; PPC, pea protein concentrate; SFM, sunflower meal; RSM, rapeseed meal; HBM, horsebean meal; -0, non-inclusion; -S, inclusion; PP, pea protein; SF, sunflower; RS, rapeseed; HB, horsebean; WG, wheat gluten; SBM, soyabean meal.

^{a,b,c,d,e} Mean values within a column with unlike superscript letters were significantly different ($P < 0.05$).

* Scores represent means using a visual analogue scale scoring system as described by Penn *et al.*⁽²⁰⁾. The score range was arbitrarily set as 0–10. Scores are based on the actual appearance of each tissue characteristic (i.e. high scores are normal for mucosal fold height and enterocyte vacuolisation; low scores are normal for mucosal fold fusion, lamina propria and submucosa width and cellularity, and enterocyte nucleus position (basal); and intermediate scores are normal for frequency of intraepithelial lymphocytes and goblet cells.

study, the varying effects of saponin supplementation that depended on basal diets were probably, at least partly, due to varying saponin levels in the diets. However, our chances of evaluating this suggestion are hampered by lack of solid information on saponin content of various plant ingredients, as previously noted. The present diets were analysed for saponins at a well-equipped laboratory, but without reliable results. The different matrices seemed to give different artifacts. We have also made efforts in our own laboratory to analyse saponogenins A and B, the two aglycones of the saponins, with equally inaccurate results. The difficulties seem to be related to variable binding of saponins to the matrices and compounds of the protein sources, as recovery of supplemented saponins differed greatly. It is known that most legumes, peas and horsebeans included, contain similar saponins as soyabeans⁽³³⁾ but in different proportions and at lower levels than that found in soyabeans⁽²⁶⁾. The saponin content in soyabean seed has been reported at levels from 2 to 60 g/kg^(34,35). Knudsen *et al.*⁽⁵⁾ suggest that commercial batches of SBM can be expected to contain 5–7 g/kg. In comparison, the reported content in peas^(36–38) varies between 0.7 and 2.5 g/kg. Up-concentration of saponins during pea meal processing, specifically air classification, may be expected. *Vicia faba* bean varieties differ in their saponin content, with 0.3 g/kg reported for broad beans⁽³³⁾ and 0.1 g/kg for field beans⁽³⁹⁾, but no reports specifically mention the horsebean variety.

Regarding the basal diets

The potential of the plant ingredients investigated in the present study as part of multisource feed formulations has been shown for a number of fish species^(28,40). In Atlantic salmon, focus is growing and knowledge accumulating on performance of plant ingredients besides soyabeans as sole or multisource formulation replacements of fishmeal as the dietary protein source. However, as mentioned, due to varying levels of fibre and other ANF, the maximum level that can be included in salmonid diets varies greatly.

Previous studies on Atlantic salmon in our group corroborate the good performance observed in the present study with the PPC and HBM basal diets. Aslaksen *et al.*⁽¹⁶⁾ found that diets with whole peas, whole and dehulled faba beans (closely related to horsebeans) at dietary inclusion rates of 18, 22 and 19%, respectively, all performed similar to an FM-based control diet. The high retention of digested methionine and cysteine for the HBM in the present study was remarkable as the digestibility for these amino acids was also high. Lysine retention was also among the highest for HBM, only second to the value observed for GC, and further strengthens the potential of HBM as a valuable protein source in feed for salmonids. In the study by Aslaksen *et al.*⁽¹⁶⁾ two pea products produced by air classification with 35% and 50% CP content also performed comparably to the FM-based control when fed to Atlantic salmon at 20% inclusion. However, in a more recent study, we have observed that, as with SBM, a PPC included at 35% in the diet was associated with negative effects such as reduced growth,

reduced lipid digestibility and induction of distal intestinal enteritis in Atlantic salmon⁽²⁰⁾. In the present study, fish performed well with no signs of the negative effects with an inclusion level of 31%. Thus, the present state of the art indicates that an inclusion level of about 30% is safe whereas 35% may be too high to prevent antinutrient-induced consequences, although some products seemingly can be included at 50% without negative consequences⁽⁴¹⁾. Furthermore, when pea products are combined with other ingredients containing saponins and possibly other antinutrients such as soya-bean products, negative nutritional effects may be expected.

Investigations of cereal glutes partially replacing FM in diets of carnivorous farmed fish species such as Atlantic cod⁽⁴²⁾, European seabass⁽⁴³⁾, gilthead seabream^(44,45), turbot⁽⁴⁶⁾ and rainbow trout^(14,47), have shown good performance regarding nutrient utilisation, growth, flesh quality and absence of negative effects on health. WG, replacing up to 50% of CP from FM, was associated with increased AD of most amino acids except alanine and lysine in diets for Atlantic salmon⁽¹⁵⁾. However, in a study with large (2.4 kg initial weight) Atlantic salmon fed a diet incorporating 30% MG showed lower SGR than fish fed the control FM diet⁽²⁰⁾. The present results agree with previous investigations in showing values of the basal diet with MG typically falling within the middle range of values observed for other ingredients regarding both nutritional and gut health indicators. In combination, MG and WG supplemented with crystalline amino acids have supported acceptable growth performance and better energy utilisation compared to an FM-based control⁽⁴⁸⁾ in feeds for Atlantic salmon. MG and WG, both possess good properties as alternative plant-based protein sources for fish feeds; i.e. low levels of fibre, starch and antinutrients, relatively high protein content with a favourable amino acid profile, high nutrient digestibility and good palatability⁽⁴⁰⁾. MG is widely used in fish feeds for salmon⁽⁴⁹⁾. Cost of the ingredients has been a major limitation to their use in fish feeds, but sharp increases in the relative price of fishmeal will eventually make these refined products increasingly more economical to use⁽⁴⁰⁾.

The comparatively low performance of the SFM basal diet may be related to the SFM content of ANF such as a high fibre content, oligosaccharides, phytic acid and arginase inhibitor⁽²⁷⁾. This is somewhat in contrast to the general consideration that SFM is a valuable protein source with good availability and relatively few ANF. Dehulling and extrusion nutritionally improve the ingredient as shown by Gill *et al.*⁽⁵⁰⁾ who recommended SFM use in Atlantic salmon feed formulations at inclusion levels of up to 23% of digestible dietary protein based on results showing no adverse effects on fish performance. However, in another study, and in line with the present results, feeding extruded diets incorporating 20% CP from SFM to Atlantic salmon reduced protein digestibility⁽¹⁶⁾. Other possibly beneficial dietary effects of SFM have been observed associated with reduced sea lice infestation on Atlantic salmon⁽⁵¹⁾. Further studies on the nutritional and health aspects of SFM are, therefore, needed for better understanding of the effects of SFM in diets for Atlantic salmon and other farmed fish species.

In the present study, the lowest fish performance was associated with the RSM basal diet. This finding was in agreement with the low performance of RSM diets also observed by Aslaksen *et al.*⁽¹⁶⁾. This poor performance can be attributed to the presence of heat-stable glucosinolates, high fibre and a potent trypsin inhibitor^(27,28). Removal of antinutrients seems necessary for RS to become a useful protein source for Atlantic salmon.

Conclusions

Dietary soyasaponin supplementation at a level of 0.2%, a level expected in diets with approximately 20–30% SBM, caused minor to moderate effects on fish growth, nutrient utilisation and intestinal physiology when supplemented to diets with MG, SFM, HBM and RSM. In the diet with PPC, the saponin supplementation induced major effects that were similar to the typical effects induced by SBM diets. Our present study therefore confirms earlier suggestions that saponins are contributors to the development of diet-induced enteritis in salmonids as seen with SBM and high levels of PPC. Care should be taken when combining plant ingredients containing saponins such as peas and SBM and possibly also other saponin-containing ingredients to avoid high levels of saponins.

Acknowledgements

The present work was carried out under an industry-driven project led by BioMar AS and partly funded by The Research Council of Norway (project no. 187294) and the Aquaculture Protein Centre (project no. 145949/120). There were no conflicts of interest for any of the authors involved in this work. The authors' contributions were as follows: E. M. C. contributed in the areas of experimental design, sampling, statistics and manuscript write-up; F. F. V. contributed to the experimental design, sampling, sample analyses and manuscript write-up; M. H. P., to the experimental design, sampling, histology and manuscript review; J. R., to the experimental design, sampling and manuscript review; S. R., to the experimental design, feeding and husbandry of experimental fish, sampling and manuscript review; A. G., to the experimental design, feed production, sampling and manuscript review; M. H., to the experimental design, feed formulation and production, sampling and manuscript review; and Å. K. contributed to the experimental design, sampling, statistics, manuscript write-up and review. Thanks are due to the animal technicians at Nofima Marin at Sunndalsøra for excellent fish care and management and to the laboratory technicians at Nofima Marin and the Gut and Health Group of the Aquaculture Protein Centre for skilful performance of all the necessary analyses.

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