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1 A study of the discrete and interactive effects of different polysaccharides on the digestibility of
2 diets fed to barramundi (*Lates calcirifer*).

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23 **Abstract**

24

25 This study examined the single, paired and combined inclusion effect of a range of
26 different polysaccharides types on the dry matter, protein and energy digestibility of diets fed to
27 barramundi (*Lates calcarifer*). The different polysaccharides included pregelatinised starch,
28 cellulose, lignin and pectin. There were significant differences among the digestibility
29 parameters of the diets with the different inclusion levels of each of the different polysaccharide
30 types. Using a MANOVA analysis effects were noted for polysaccharide type, inclusion level
31 and interaction terms on the digestibilities of dry matter, protein and energy. Cellulose addition
32 resulted in a reduction in both dry matter and energy that was largely commensurate with its
33 inclusion level, but its effect on protein digestibility was marginal. Starch had the least effect on
34 any of the digestibility parameters of all the polysaccharide types examined. Pectin had the
35 largest effect on dry matter, while lignin had the greatest impact on diet protein and energy
36 digestion. In the diets with paired combinations of polysaccharides, lignin and pectin were
37 responsible for negatively synergistic interactions in all digestibility parameters. These results
38 show that different polysaccharide classes can have distinctly different effects on diet
39 digestibility parameters. The results also show that some classes of polysaccharide have greater
40 interactive effects than others.

41

42 1. **Introduction**

43 Aquaculture production is becoming increasingly reliant on the replacement of fishmeal
44 with a combination of plant meals in feeds (Gatlin, et al., 2007). Already the use of terrestrial
45 plant-derived raw materials in fish feeds is common place (Aslaksen et al., 2007; Gatlin et al.,
46 2007; Hardy, 2010). Plant materials have recognised benefits including a reduction in
47 formulation costs and improved functional characteristics (Krogdahl et al., 2010; Glencross et
48 al., 2010). Less understood are the anti-nutritional effects caused by the introduction of non-
49 starch polysaccharides (NSP) that are present in the carbohydrate fraction of many of these plant
50 materials. The NSP content of many plant materials is recognised as a key factor which has
51 been shown to affect the nutritional variability among these materials being fed to fish
52 (Glencross et al., 2008, 2012a; 2012b).

53 Fish, like most monogastrics are generally capable of some level of starch digestion
54 (Bergot and Breque, 1983; Amirkolaie et al., 2006; Enes et al., 2008; Moreira et al., 2008;
55 Glencross et al., 2012a). However, it is the presence of the plant cell wall derived NSP's which
56 has caused many complications due largely to their indigestible nature (Kraugerud et al., 2007;
57 Hansen and Storebakken, 2007; Glencross, 2009). The non-nutritive value of most NSP's means
58 they act largely as a bulking agent, which generally results in a reduction in feed digestibility
59 with increased content in the feed. Although the non-nutritive value of many NSP's is well
60 understood, what is less understood is the anti-nutritional and or interactive effects of the more
61 complex NSP's which lead to a reduction in digestibility which exceeds their level of inclusion
62 (Glencross et al., 2008; 2012b). An example of this is seen with the presence of lignin in diets of
63 rainbow trout either via inclusion of some plant proteins or as a specific additive (Glencross et
64 al., 2008; 2012b). With the common practice of blending different plant proteins and
65 carbohydrates, it is therefore becoming increasingly important to consider not only the discrete
66 effects but also the interactive of NSP's.

67 There have been various reports on the discrete effects of different NSP classes in fish
68 diets (Glencross et al., 2003; Leenhouders et al., 2004; 2007; Amirkolaie et al., 2005; Hansen
69 and Storebakken, 2007; Glencross, 2009, Glencross et al., 2012b). Studies with rainbow trout
70 have shown cellulose (insoluble NSP) inclusion in diets cause a reduction in dry matter and
71 energy digestibility, but only a nominal effect on protein digestibility of the diet. (Hansen and
72 Storebakken, 2007; Glencross, 2009, Glencross et al., 2012b). In a study by Glencross (2009)
73 assessing insoluble and soluble NSP classes in lupins in rainbow trout diets, the classes were
74 reported to cause different effects on diet digestibility. However, a further study detected only
75 marginal differences in across soluble NSP classes of pectin and mannan (Glencross et al
76 2012b). The inclusion of lignosulphonate had the largest negative effect on diet digestibility,

77 particularly on protein digestibility (Glencross et al., 2012b). It was suggested that the cause of
78 the larger variability in the earlier study may be due to an interaction effect and that the testing
79 of blended NSP's should be considered. Therefore this study aims to examine the effects of the
80 discrete and blended inclusion levels of different classes of NSP (cellulose, lignin and pectin)
81 and starch on the digestible value of diets fed to barramundi, *Lates calcarifer*.

82 2. Materials and Methods

83 2.1 Diet preparation

84 The experiment design was based on a diet formulation strategy that used a diet-
85 substitution approach, although the assessment of the digestible value of those ingredients was
86 not the intent of this experiment (Glencross et al., 2007). To achieve this, a basal diet was
87 formulated and prepared to include approximately 600 g/kg DM protein, 185 g/kg DM fat and
88 an inert marker (yttrium oxide at 1 g/kg) (Table 1). Each polysaccharide ingredient or blend was
89 added at to the test diets at 200 g/kg inclusion to a reciprocal-sample of the basal mash (see
90 Table 1). The diets were made by the addition of water (about 25% of mash dry weight) to the
91 mash whilst mixing to form a dough which was subsequently screw pressed using a pasta maker
92 through a 4 mm diameter die. The moist pellets produced were then oven dried at 60°C for
93 around 12 h before being allowed to cool to ambient temperature in the oven. The basal diet was
94 prepared in a similar manner, but without the addition of any test ingredient. The source and
95 composition of all ingredients is presented in Table 2.

97 2.3 Fish handling and faecal collection

98 Hatchery-reared barramundi (*Lates calcarifer*) were kept in a series of experimental
99 tanks (300 L) supplied with aeration and heated flow-through seawater (salinity =35 PSU) at a
100 flow rate of about 3 L/min maintaining dissolved oxygen at 6.2 ± 0.2 mg/L and temperature at
101 $28.6 \pm 0.2^\circ\text{C}$ (mean \pm S.D.). Each of the tanks were stocked with 10 fish of 397.7 ± 68.9 g
102 (mean \pm S.D.; n = 40 from a representative sample of the population). Treatments were
103 randomly assigned amongst 24 tanks, with each treatment having two replicates. The
104 experiment was repeated and blocked by time to achieve a total of four replicates per treatment.
105 The same batch of fish was used for both blocks, but with a randomisation of the fish between
106 each block. The fish were allowed to acclimatise to their allocated dietary treatment for at least
107 seven days before faecal collection commenced (Blyth et al., 2012).

108 The barramundi were manually fed once daily to apparent satiety, as determined over
109 three separate feeding events between 0800 and 0900 each day. Faeces were collected the same
110 afternoon (1600 – 1700) from each fish within each tank using stripping techniques based on
111 those reported by Glencross (2011). Fish were anaesthetised using AQUI-S (0.002 mL/L) in a
112 separate smaller aerated tank. Once loss of equilibrium was observed, close attention was paid
113 to the relaxation of the ventral abdominal muscles of the fish to ensure the fish were removed
114 from the water before they defecated in the anaesthetic tank. The faeces were then expelled
115 from the distal intestine using gentle abdominal pressure while the ventral abdominal muscles
116 were relaxed. Hands were rinsed between handling each fish to ensure that the faeces were not

117 contaminated with urine or mucous. The faecal sample was placed in a small plastic vial and
118 stored in a freezer at -20°C. Fish were not stripped on consecutive days in order to minimise
119 stress on the animal and maximise feed intake prior to faecal collection. Faeces were collected
120 over a six-day period, with each fish being stripped three times, once every second day. Faecal
121 samples from different days were pooled within tank, and kept frozen at -20°C before being
122 freeze-dried in preparation for analysis.

123

124 2.4 *Chemical and digestibility analysis*

125 Diet and faecal samples were analysed for dry matter, yttrium, nitrogen and gross
126 energy content. Diets and ingredients were analysed for these same parameters and in addition
127 for ash, total lipid, lignin, neutral-detergent fibre and acid-detergent fibre. Dry matter was
128 calculated by gravimetric analysis following oven drying at 105°C for 24 h. Total yttrium
129 concentrations were determined after mixed acid digestion using inductively coupled plasma
130 mass spectrophotometry (ICP-MS) based on the method described by (McQuaker et al., 1979).
131 Protein levels were calculated from the determination of total nitrogen by CHNOS auto-
132 analyser, based on N x 6.25. Gross ash content was determined gravimetrically following loss of
133 mass after combustion of a sample in a muffle furnace at 550°C for 12 h. Total lipid content of
134 the diets was determined gravimetrically following extraction of the lipids using
135 chloroform:methanol (2:1). Gross energy was determined by ballistic bomb calorimetry. Dietary
136 fibres were determined by digesting the defatted sample with multiple washes of acetone and
137 ethanol. The resulting residue was corrected for undigested protein and ash according to the
138 method of the Champ et al. (1998). Neutral-detergent fibre (NDF) samples were boiled with
139 buffered NDF solution. The residue was collected on a coarse sintered glass crucible (Van
140 Soest and Robertson, 1981). The acid-detergent fibre (ADF) was determined following a
141 sample being reacted in 0.5M acid detergent solution and the residue was collected on a coarse
142 sintered glass crucible after, the method of Van Soest and Goering (1970). Lignin was
143 determined by reacting the ADF residue with cold 72% sulphuric acid. The sample was ashed
144 and the residue measured gravimetrically (Van Soest and Robertson, 1981). Total carbohydrate
145 content was determined based on dry matter – (protein+lipid+ash) content. Cellulose content
146 was determined based on the ADF – Lignin. Hemicellulose content was determined based on
147 NDF – ADF content.

148 Differences in the ratios of the parameters of dry matter, protein or gross energy to
149 yttrium, in the feed and faeces in each treatment were calculated to determine the apparent
150 digestibility (AD_{diet}) for each of the nutritional parameters examined in each diet based on the
151 following formula (Maynard and Loosli, 1979):

152
$$AD_{diet} = \left(1 - \frac{Y_{diet} \times Parameter_{faeces}}{Y_{faeces} \times Parameter_{diet}} \right) \times 100$$

153

154 where Y_{diet} and Y_{faeces} represent the yttrium content of the diet and faeces respectively, and
155 $Parameter_{diet}$ and $Parameter_{faeces}$ represent the nutritional parameter of concern (organic matter,
156 protein or energy) content of the diet and faeces respectively.

157

158 2.5 *Statistical analysis*

159 All values are means unless otherwise specified. Effects of each of the discrete
160 treatments on the digestibility of dry matter, protein and gross energy in each of the diets
161 were initially examined by a one-way ANOVA (Table 3). Levels of significance were
162 determined using a Least Significant Difference (LSD) test. Limits for all critical ranges were
163 set at $P < 0.05$. Further analysis of the effects of each NSP type were examined using Multiple
164 regression to identify magnitude, vectors and significance of effects related to inclusion level of
165 each NSP (Table 3). Additionally a stepwise forward regression was also employed to
166 determine the additive effects of the different NSP classes which influenced each digestibility
167 parameter (Table 5). These tests were undertaken using the Statistica™ v6.0 software (Statsoft®,
168 Tulsa, OA, USA). Regression analysis was undertaken using the data analysis tools of
169 Microsoft excel.

170 **3. Results**

171

172 *3.1 Ingredient composition*

173 There were substantial differences in the chemical characteristics of the four different
174 polysaccharide ingredients used in this study. The measurement of ADF, NDF and lignin
175 allowed the determination of lignin, cellulose and hemicellulose contents of each sample (Table
176 1). Lignin content was highest in the liginosulphate (99 g/kg DM) and lowest in the pectin (0
177 g/kg DM). Cellulose content was highest in the purified cellulose ingredient (827 g/kg DM) and
178 lowest in the starch (1 g/kg DM) and pectin (0 g/kg DM). Similarly, hemicellulose was also
179 highest in the purified cellulose ingredient (129 g/kg DM) and lowest in the starch (2 g/kg DM)
180 and pectin (0 g/kg DM). Protein content in all test ingredients was low (<30 g/kg DM) as was
181 also the total lipid content (<62 g/kg DM).

182

183 *3.2 Diet digestibilities*

184 There were distinct differences between the diets in terms of their dry matter and energy
185 digestibility (Table 3), with differences among protein digestibilities of the diets generally being
186 somewhat less than the other two parameters. A two-way ANOVA analysis (MANOVA)
187 identified that there were significant effects of the different polysaccharide types (P=0.001) and
188 also the inclusion levels (P=0.001) of each of the polysaccharide types on the digestibility of dry
189 matter and energy of each diet (Table 3). A significant (P=0.001) interaction term was also
190 defined between the type of polysaccharide and the level of its inclusion (Table 3). Regression
191 analysis of each polysaccharide type and inclusion level clearly shows the effect of each
192 polysaccharide type on the digestibilities of dry matter, protein and energy (Table 1 and 4).

193 The addition of cellulose caused a reduction in dry matter digestibility that was almost
194 directly commensurate with the level of its addition (Table 1 and 4). Lignin caused a slightly
195 larger reduction on dry matter digestibility (Table 1 and 4). Pectin caused a reduction in dry
196 matter digestibility almost double that of its inclusion level (Table 1 and 4). Starch inclusion
197 showed a nominal effect on dry matter digestibility.

198 The addition of the cellulose + starch blend was observed to make a reduction in dry
199 matter digestibility that was close to commensurate with the level of cellulose addition (Table 1
200 and 4). The cellulose + pectin + starch + lignin blend was observed to make a reduction in dry
201 matter digestibility that was near commensurate with the combined level of addition. The starch
202 + lignin and cellulose + pectin + starch + lignin blends also had a similar such effect on dry
203 matter digestibility. The pectin + lignin blend caused a reduction to dry matter digestibility
204 almost double that of its inclusion level (Table 1 and 4). The cellulose + lignin blend had a

205 similar such effect on dry matter digestibility. The pectin + starch blend was also observed to
206 cause a reduction in the dry matter digestibility which exceeded the level of NSP addition, but
207 not to the same degree as observed with the pectin + lignin or cellulose + lignin blend.

208 The addition of starch had almost no impact on protein digestibility (Table 1 and 4).
209 The cellulose and cellulose blends containing starch or pectin had a similar effect on protein
210 digestibility. The pectin + starch blend caused only a minor reduction in protein digestibility.
211 However, the addition of pectin, cellulose + lignin and the cellulose + pectin + starch + lignin
212 blend caused a reduction in protein digestibility which exceeded the pectin + starch blend, but
213 was lower than total inclusion level. The lignin and starch + lignin blend were observed to cause
214 a reduction protein matter digestibility that was close to commensurate with the level addition.
215 The pectin + lignin blend was observed to make a reduction in protein matter digestibility which
216 exceeds the level of addition.

217 The cellulose blends containing starch or pectin caused a reduction in energy
218 digestibility (Table 1 and 4), slightly less than the level of addition. The addition of cellulose or
219 starch was observed to make a reduction in energy matter digestibility that was close to
220 commensurate with the level of addition. Pectin, had a similar such effect on energy
221 digestibility. The starch + lignin, starch + pectin and cellulose + pectin + starch + lignin blends
222 caused a reduction in energy digestibility which exceeded total addition. The pectin + lignin and
223 cellulose + lignin blend had a reduction effect on dry matter digestibility almost double the
224 inclusion level. Lignin caused a reduction in protein digestibility greater than the pectin + starch
225 blend, but lower than the pectin + lignin blend.

226 Step wise regression indicated that pectin was the dominant fibre affecting dry matter
227 digestibility, followed by lignin and then cellulose (Table 5). The presence of pectin, lignin and
228 cellulose in the model explained 36.9% of the variation in diet dry matter digestibility. The
229 addition of starch did not add to the model for diet dry matter or protein digestibility. Lignin
230 was the dominant fibre class affecting diet protein digestibility, followed by pectin and then
231 cellulose (Table 5) accounting for 80.3% of the variation in digestibility. Lignin was also the
232 dominant fibre class affecting diet energy digestibility, followed by pectin, cellulose and then
233 starch (Table 5) accounting for 31.2% of the variation in digestibility.

234

235

236 4. Discussion

237

238 It has been demonstrated that different polysaccharide types cause varying effects on
239 nutrient and energy digestibilities when included in fish diets (Glencross, 2009; Glencross et al.,
240 2012b). It is now understood that the presence of these complex carbohydrates has a significant
241 bearing on the variability in many of the nutritional responses by fish to diets (Aslaksen et al.,
242 2007). While the discrete effects of different polysaccharides have been studied, little
243 information is available on the potential interactive effects among the different polysaccharides
244 classes. As raw materials contain different polysaccharide types and are typically added in a
245 combination rather than added in isolation to aquaculture feeds, it is important to consider the
246 implications of the addition of several different types of these polysaccharides to the nutritional
247 value of feeds. It has been suggested that it is the interactive effect of these different
248 polysaccharides that contributes to the variable effects seen in the digestibility values of many
249 raw materials (Glencross et al., 2008; 2012b).

250

251 4.1 Polysaccharide class discrete effects

252 Each of the dietary non-starch-polysaccharide classes had a clear effect on the
253 digestibility of dry matter content of diets fed to barramundi. This is a clear contrast to the effect
254 seen with the pregelatinised wheat starch, which had little effect on dry matter digestibilities.
255 This difference clearly shows that although this species of fish can digest pregelatinised starch
256 that it has almost no ability to digest any of the non-starch-polysaccharide (NSP) classes tested
257 in this study.

258 With the majority NSP classes the decline in dry matter digestibility is directly related
259 to the level of inclusion on of each NSP sample. This effect was seen directly with cellulose and
260 to a lesser degree with lignin. These observations are similar to those observed by Glencross et
261 al. (2012b), who also reported a significant decline in organic matter digestibility of diets with
262 different levels of cellulose inclusion when fed to rainbow trout. Similar such effects were also
263 reported by Hansen and Storebakken (2007), who also reported a significant decline in organic
264 matter digestibility of diets with high cellulose inclusion when fed to rainbow trout. Glencross
265 (2009) also noted the same effects with cellulose on dry matter and energy digestibilities and
266 also the insoluble NSP content of lupin kernel meals. These observations combined with the
267 present ones support that in most cases the NSP are simply acting as non-nutritive filler and
268 have limited greater interactive effect on diet dry matter digestibility.

269 However, in the present study, the single or blended addition of pectin or lignin resulted
270 in reductions in digestibility at levels which exceed their inclusion level and are indicative of an

271 interactive anti-nutritional factor. Pectin is a soluble fibre, commonly used as a gelling agent in
272 products for human consumption. It has been suggested that NSP's in soluble form are most
273 detrimental to fish (Reftsie, et al., 1999), due to their ability to disrupt digestive function by
274 increasing the viscosity of intestinal contents (Francis et al., 2001). When pectin is blended with
275 any of the fibre types it is possible this interaction is occurring and leading to reduced diet
276 digestibility. A similar interaction was observed with the cellulose + lignin blend diet.

277 A multivariate approach by Glencross et al (2008) assessed over seventy lupin meals
278 with rainbow trout. It was shown in that study that lignin content negatively affected nitrogen
279 digestibility and that in combination lupin protein level and lignin content was the strongest
280 predictor of protein digestibility. This result was supported by a subsequent study with rainbow
281 trout in which lignin inclusion (as liginosulphate) in particular was observed to have a significant
282 effect on diet protein digestibilities (Glencross et al., 2012b). That study assessed three inclusion
283 levels 25, 50 and 100g/kg and interestingly, the effect on protein digestibility ceased to be linear
284 after the 50 g/kg inclusion level. It was suggested that this could be due to a saturation effect of
285 the liginosulphate on whatever loci it is disrupting in the digestion process. These findings are
286 consistent with observations from other studies on the lignin fibre class, with liginosulphanate
287 also found to have significant effects on diet digestibility even at very low inclusion levels
288 (Glencross., 2009; Glencross et al., 2008; 2012b).

289 The different dietary NSP types had clear effects on the digestibility of the energy
290 content of diets fed to barramundi. The response of diet energy digestibility to inclusion of the
291 different NSP's, either singly or blended, is consistent with the fact that these fish are obtaining
292 nominal energetic value from the presence of NSP's in the diet. This is also consistent with
293 recent findings that show a limited energetic value from the inclusion of NSP's in diets for trout
294 (Glencross et al., 2012b). Interestingly, when lignin (100g/kg) was pair combined with any of
295 the fibre classes it resulted in an exacerbated reduction in diet energy digestibility indicative of a
296 negatively synergistic interaction. However, this was not observed in the diet which contained a
297 50g/kg inclusion of each of the fibre classes, suggesting that providing the inclusion of lignin is
298 $\leq 50\text{g/kg}$ it is unlikely to significantly disrupt energy digestibility in barramundi.

299

300 4.2 Conclusions

301 This study demonstrates that there are both discrete and interactive effects of the
302 different polysaccharides on the nutrient and energy digestibilities of diets fed to barramundi.
303 While the direct manipulative approach of the present study provides some clear indications on
304 which NSP from different feed grains might affect their own digestibility, it would be useful to
305 follow up this work with an assessment of a broad suite of feed grains and cross reference these

306 observations using a multivariate analysis approach, similar to that done by Glencross et al
307 (2008) with lupin meals, but instead the proposed study should examine a cross section of the
308 different feed grains available. Being able to corroborate the results from the present study with
309 that from a multivariate analysis of actual feed grains will help consolidate the hypothesis that it
310 is the NSP complexity in these raw materials that is a key cause of nutritional value variability.
311 Secondly further research to identify why both pectin and lignin have the effects that the do
312 would be useful to understand the mode of action of these NSP on the digestion of diets by fish.

313

314

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318

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Tables and Figures

Table 1. Formulations, composition and digestibility coefficients of the experiment diets (all values except ADC's are g/kg).

Ingredient	Basal	A	B	C	D	E	F	G	H	I	J	K	Pooled SEM
Fishmeal	764	611.2	611.2	611.2	611.2	611.2	611.2	611.2	611.2	611.2	611.2	611.2	
Fish oil	100	80	80	80	80	80	80	80	80	80	80	80	
Wheat gluten	130	104	104	104	104	104	104	104	104	104	104	104	
Cellulose	-	200	-	-	-	100	100	100	-	-	-	50	
Pectin	-	-	200	-	-	100	-	-	100	100	-	50	
Pregelld starch	-	-	-	200	-	-	100	-	100	-	100	50	
Lignin	-	-	-	-	200	-	-	100	-	100	100	50	
Vitamin premix	5	4	4	4	4	4	4	4	4	4	4	4	
Yttrium oxide	1	0.8	0.8	0.8	0.8	0.8	0.8	0.8	0.8	0.8	0.8	0.8	
Dry matter	937	944	920	944	932	937	937	941	929	919	934	923	
Protein	637	499	488	513	518	501	515	519	501	515	474	519	
Lipid	197	189	157	177	174	171	150	171	167	151	163	161	
Ash	155	124	130	121	157	129	125	141	129	128	143	135	
Total Carbohydrates	11	188	226	190	151	200	209	170	204	206	220	185	
- Starch	49	10	15	268	0	28	138	67	224	87	300	156	
- ADF	45	181	34	51	79	106	135	125	58	74	77	99	
- NDF	33	177	26	39	63	96	128	115	45	63	68	92	
- Lignin	6	11	8	3	29	5	0	16	1	9	11	9	
- Cellulose	39	170	26	47	50	100	135	109	57	65	66	90	
- Hemicellulose	12	4	8	12	15	10	7	10	12	11	9	6	
Gross Energy	22.07	20.56	20.95	20.78	22.18	20.38	21.18	20.65	20.60	20.83	21.74	20.59	
Dry matter ADC	0.742	0.591	0.456	0.744	0.550	0.592	0.655	0.458	0.508	0.443	0.598	0.591	0.021
Protein ADC	0.908	0.941	0.810	0.916	0.692	0.875	0.932	0.790	0.848	0.650	0.710	0.767	0.015
Energy ADC	0.814	0.651	0.628	0.640	0.562	0.686	0.676	0.487	0.608	0.456	0.456	0.620	0.021

* Vitamin and mineral premix includes (IU/kg or g/kg of premix): Vitamin A, 2.5MIU; Vitamin D3, 0.25 MIU; Vitamin E, 16.7 g; Vitamin K,3, 1.7 g; Vitamin B1, 2.5 g; Vitamin B2, 4.2 g; Vitamin B3, 25 g; Vitamin B5, 8.3; Vitamin B6, 2.0 g; Vitamin B9, 0.8; Vitamin B12, 0.005 g; Biotin, 0.17 g; Vitamin C, 75 g; Choline, 166.7 g; Inositol, 58.3 g; Ethoxyquin, 20.8 g; Copper, 2.5 g; Ferrous iron, 10.0 g; Magnesium, 16.6 g; Manganese, 15.0 g; Zinc, 25.0 g. **estimated from the Carbohydrate content of each test ingredient and its inclusion level in each diet.

Table 2. Measured nutrient composition of the experimental ingredients (all values are g/kg DM unless otherwise indicated)

	Fishmeal	Gluten	Cellulose	Starch	Lignin	Pectin
Dry Matter	954	912	930	891	891	887
Protein	682	801	7	7	7	30
Lipid	101	80	9	16	6	6
Ash	183	9	1	3	669	53
Total Carbohydrates*	35	110	983	975	318	911
- Starch	28	302	0	997	31	23
- ADF	72	91	928	0	0	7
- NDF	44	91	925	0	0	5
- Lignin	6	187	290	0	613	0
- Cellulose	66	0	637	0	0	7
- Hemicellulose	28	0	2	0	0	3
Gross Energy	18.3	22.8	17.2	16.8	21.0	14.9

^a Pregelatinised wheat starch: Manildra, Auburn, NSW, Australia. ^b Cellulose and Pectin: Sigma Chemical Company, St Louis, MO, USA. ^b Calcium ligninosuphate: Dustex, Canningvale, WA, Australia. *calculated based on dry matter – (protein+ash+fat)

Table 3. Multiple regression analyses of ingredient digestible values

	Parameter	Vector	Std.err.	Coefficients	Std.err	T(43)	p-value
ADC-DM	Intercept			0.753	0.061	12.364	0.000
	Cellulose	-0.347	0.166	-0.008	0.004	-2.097	0.042
	Pectin	-0.658	0.166	-0.015	0.004	-3.972	0.000
	Starch	-0.087	0.166	-0.002	0.004	-0.528	0.600
	Lignin	-0.555	0.166	-0.013	0.004	-3.351	0.002
ADC-PRO	Intercept			0.914	0.023	39.263	0.000
	Cellulose	0.094	0.092	0.001	0.001	1.019	0.314
	Pectin	-0.413	0.092	0.007	0.001	-4.480	0.000
	Starch	-0.070	0.092	0.001	0.001	-0.757	0.453
	Lignin	-0.894	0.092	0.014	0.001	-9.711	0.000
ADC-E	Intercept			0.838	0.062	13.433	0.000
	Cellulose	-0.417	0.173	-0.009	0.004	-2.405	0.021
	Pectin	-0.523	0.173	-0.012	0.004	-3.106	0.004
	Starch	-0.386	0.173	-0.009	0.004	-2.226	0.031
	Lignin	-0.750	0.173	-0.017	0.004	-4.327	0.000

Table 4. Step wise regression analyses of ingredient digestible values

Parameter	Variable	Step	Multiple-R	Multiple-R ²	R ² Change	F – to enter	P-value
ADC-DM	Pectin	1	0.392	0.154	0.154	8.346	0.006
	Lignin	2	0.546	0.298	0.145	9.290	0.004
	Cellulose	3	0.607	0.369	0.071	4.921	0.032
ADC-PRO	Lignin	1	0.790	0.624	0.624	76.393	0.000
	Pectin	2	0.888	0.789	0.165	35.154	0.000
	Cellulose	3	0.896	0.803	0.014	3.245	0.078
ADC-E	Lignin	1	0.394	0.155			
	Pectin	2	0.451	0.203	0.048	2.722	0.106
	Cellulose	3	0.483	0.233	0.030	1.713	0.197
	Starch	4	0.559	0.312	0.079	4.957	0.031

