

Variation in the chemical composition and the nutritive quality of different field bean UK grown cultivar samples for broiler chicks

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1 **Variation in the chemical composition and the nutritive quality of different field**
2 **bean UK grown cultivar samples for broiler chicks**

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12

13 **Abstract** 1. The chemical composition and physical characteristics of ten colour-flowered
14 different UK grown field bean cultivar samples from the same harvest year were determined.
15 Compositional variation existed between the beans.

16 2. Diets that included each bean sample at 200 g/kg were used to compare broiler growth
17 performance and determine N-corrected apparent metabolizable energy (AMEn) and
18 nutrient utilisation. The AMEn and nutrient retention coefficients for the bean samples were
19 obtained via slope ratio method. The relationships were examined between these variables
20 of nutritive value for broilers and the laboratory analysis on the bean samples.

21 3. Findings showed differences ($P < 0.05$) among the bean cultivar samples for feed
22 conversion ratio, AMEn and dry matter retention (DMR) coefficients. Further analysis
23 showed that feeding quality of different field bean cultivar samples measured as AMEn
24 highly correlates to crude protein (CP) ($P < 0.05$) contents and the colour ($P < 0.001$) of the
25 samples. Thus, beans with higher CP and pale colour have superior feeding value for
26 broilers.

27 Key words: broilers, field beans, nutrient availability, metabolisable energy, chemical
28 composition

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34 **Introduction**

35 Soybeans, which are a common source of protein in poultry diets due to having a high level
36 of protein and well balanced profile of amino acids, are an imported feed ingredient in the
37 UK. From 2002-2018 the average total of imported soybeans (as whole seed) to the UK
38 was about 791 930 tonnes per year (Statista 2020). The price of soybean meal (SBM) is
39 increasing continuously as a result of rising demand globally, particularly after prohibition of
40 animal protein inclusion in poultry diet by the European Union (O'Neill et al. 2012).
41 Additionally, large amount of the available SBM in the market is produced from genetically
42 modified crops which worries consumers and is not suitable for organic production (Vicenti
43 et al. 2009). Also, according to Gasparri et al. (2013), increasing global demand for SBM
44 caused deforestation of millions of hectares in South America, especially over last half
45 century, which have left a negative impact on Carbon Footprint and Global Environmental
46 Changes. Furthermore, recently the European Union has stimulated animal producers to
47 exploit locally grown legumes such as field beans in their diet formulations, aiming for
48 ecological and financial benefits (Fru-Nji et al. 2007). Therefore, the search for locally grown
49 alternative protein sources that can totally or at least partially replace SBM is necessary,
50 thus, decreasing or ending the dependency of the UK poultry feed industry on imported
51 SBM as a source of protein and avoiding or reducing the worries connected with SBM. High
52 concentrations of proteins in field beans and similarity of their amino acids profile to that of
53 soybeans, renders them to be considered as a desirable candidate to replace SBM, at least
54 partially, in poultry diet formulations. Field beans could possibly be produced in greater
55 amounts in the UK due to the suitability of the climate and the available cultivars. In the
56 recent years, as a consequence of breeding and increased area where field beans are being
57 planted in, the annual UK production of field beans has approximately 600 000 tonnes
58 (PGRO 2017), however, very little of it is employed in UK animal feed formulation and the

59 rest is exported. Nowadays, the demand on producing field beans is increasing and this
60 increase is expected to continue, thus they will be an available feedstuff at a high amount
61 in the UK market.

62 It has been reported that field beans can be included at 20 to 30% in broiler diets without
63 any adverse effect on performance (Farrell et al. 1999; Usayran et al. 2014; Abdulla et al.
64 2017). However, there is uncertainty about the chemical composition, which may vary
65 between different cultivars, especially with regards to the type and amount of anti-nutrients
66 that they may contain. This may also result in variation of their energy and nutrient
67 availability for broilers.

68 Adequate and precise information on the chemical composition and nutritive value of
69 different locally-grown field bean cultivars provides flexibility and constancy to the poultry
70 feed industries, allowing them to include field beans in their diet formulations.

71 The main objectives of this experiment were: To examine the differences in
72 metabolizable energy, and nutrient availability of ten UK grown field bean cultivar samples.
73 To examine the relationship between chemical composition of the field bean sample to the
74 bioavailable energy and nutrients. Differences in productive performance of broilers when
75 fed these beans are reported and compared although the nutrient variation between the
76 bean samples was not corrected in the diets.

77

78 **Material and methods**

79

80 ***Field bean cultivar samples***

81 Ten colour-flowered different UK grown field bean cultivar samples, including three spring
82 (Fuego, Fury and Maris Bead) and seven winter (Arthur, Buzz, Clipper, Divine, Honey,
83 Sultan and Wizard) grown cultivars, from 2013 harvest year were obtained from the market
84 (primarily from Askew & Barrett (Pulses) Ltd, Wisbech, UK). The beans were grown at
85 different locations and there was no information on agronomy or soil type available. All
86 harvested field bean samples were stored at ambient air temperatures in a dry store and
87 were used in broiler feeding experiment after approximately 6 months of their storage.

88 Before the animal feeding experiment, the field bean samples were hammer-milled using a
89 4 mm screen and then mixed in a horizontal mixer with the other feed ingredients. Freshly
90 milled field beans were used for analyses and in the feeding study to avoid spoilage.

91

92 ***Proximate analysis and gross energy in field bean and excreta samples***

93 Dry matter was determined by drying samples at 105°C to constant weight (AOAC 1990;
94 925.10). Crude protein (N x 6.25) concentration in the samples was determined by the dry
95 combustion method, using a Leco (FP-528 N, Leco Corp., St. Joseph, MI). Oil (as ether
96 extract) was extracted with diethyl ether by the extraction method (AOAC 2000), employing
97 a Soxtec system (Foss UK Ltd). Gross energy of the samples was measured using a Parr
98 adiabatic bomb calorimeter (Parr-6200 Calorimeter, Parr Instruments Company, Moline, IL,
99 USA), and benzoic acid was used as the standard.

100

101 ***Carbohydrate and mineral contents in field bean samples***

102 Total starch was determined by a modified version of Englyst et al. (2000), which involved
103 initial heat dispersion together with heat stable amylase followed by treatment with alkali to
104 disperse any retrograded type III resistant starch. Non-starch polysaccharides (NSP)
105 content in field beans was determined by the method of Englyst et al. (1994). In brief, starch
106 is completely dispersed, hydrolysed enzymatically, and the NSP is isolated by precipitation
107 in 80% ethanol, hydrolysed by sulphuric acid and the released sugars are measured by gas
108 chromatography.

109 The mineral contents of the field bean samples were determined according to the procedure
110 described by Tanner et al. (2002), employing inductively coupled plasma emission
111 spectrometry (Optima 4300 DV Dual View ICP-OE spectrometer, Perkin Elmer,
112 Beaconsfield, UK).

113

114 ***Phenols, tannins, phytate and trypsin inhibitors***

115 Phenolic compounds, including total phenols, non-tannin phenols, and total tannins (as
116 tannic acid equivalent), in the representative samples of the field beans were measured

117 chemically as described by Makkar et al. (1993). In brief, phenolic compounds from samples
118 were extracted with 70% aqueous acetone and measured using spectrophotometer. The
119 tannin extract containing tubes were kept on ice until all phenolic analyses were completed
120 during the same day. The phytate and phytate phosphorus contents in the field bean
121 samples were determined by HPLC as described by Kwanyuen and Burton (2005). Trypsin
122 inhibitor content in the field beans was measured by applying the assay proposed by Smith
123 et al. (1980).

124

125 ***Grain quality and viscosity of the experimental bean samples***

126 The colour score of whole fine milled bean of each cultivar sample was read in triplicate,
127 after submerging the instrument into the samples in petri dishes, employing an L* a* b*
128 colour space (Konica Minolta, Chroma Meter CR-400, Essex, UK). The instrument was
129 calibrated against a standard white-coloured reference tile and cleaned between taking
130 measurements of different samples. The L* indicates lightness, 0-100 representing dark to
131 light. The a* value gives the degree of the red-green colour, with a higher positive a* value
132 indicates more red. The b* value indicates the degree of the yellow-blue colour, with a higher
133 positive b* value indicating more yellow.

134 For the determination of hull to kernel ratio, 100 grams of clean representative grain
135 sample of each field bean variety was taken, seed coats were completely separated from
136 cotyledons with the aid of pliers, and the weights of each of cotyledons and seed coats
137 alone were measured. The weight of 1000 grains, the water holding capacity and the water
138 extracted viscosity of the field beans were determined as previously described by Pirgozliev
139 et al. (2003). Water extracted viscosity was measured with a rotating cone and cup
140 viscometer (model DV-II + LV Brookfield, Stroughton, MA, USA).

141

142

143

144 ***Diet formulation***

145 Birds were fed one of twelve mash diets. A wheat-soybean based balancer diet (control
146 diet) was formulated that had major ingredients of 533.2 g/kg wheat, 150.0 g/kg SBM, 175.0
147 g/kg full fat soya, 37.4 g/kg maize gluten meal, and 50 g/kg vegetable oil, and contained
148 231 g/kg CP and 13.71 MJ/kg metabolisable energy (ME) (Table 1). The balancer diet had
149 higher ME content than breeder's recommendation (Aviagen Ltd., Edinburgh, UK) to allow
150 the ME of the field bean containing diets to be close to the requirements. The balancer diet
151 also contained 5 g/kg of TiO₂ as an indigestible marker, although this was not used for
152 further analysis. Ten diets were then produced including 200 g/kg of one of the ten different
153 field bean cultivars and 800 g/kg of the balancer feed. To allow testing of whether there was
154 a linear relationship between the level of substitution of an individual field bean sample and
155 the determined ME or nutrient availability of the diet, another diet was formulated that
156 contained 100 g/kg of the Honey field bean sample and 900 g/kg of the balancer feed (so
157 giving three diets with three different inclusion rates of the cultivar Honey). Twelve
158 experimental diets were compared in total. Freshly milled field beans were used in the
159 formulation of the diets and were fed as mash. All diets approximately met or exceeded the
160 dietary specifications for Ross 308 broilers (Aviagen Ltd., Edinburgh, UK). Diets did not
161 contain any coccidiostat, antimicrobial growth promoters, prophylactic or other similar
162 additives.

163

164 ***Husbandry and sample collection***

165 The experiment was conducted at the National Institute of Poultry Husbandry and approved
166 by the Research Ethics Committee of Harper Adams University, UK. Approximately five-
167 hundred day-old male Ross 308 broiler chicks were obtained from a commercial hatchery
168 (Cyril Bason, Shropshire, UK). All chicks were placed in a single rear pen at 33°C and fed
169 a proprietary broiler starter feed *ad libitum* over seven days. On the first day of the
170 experiment (8 d of age), all chicks were individually weighed and unusual birds were
171 discarded, leaving 480 birds. Those birds were then randomly allocated into 96 raised-floor
172 pens (0.36 m² floor area; five birds in each pen). The pens were arranged in one tier level

173 within a controlled environment room, and each pen was equipped with a plastic feeder and
174 drinker. The floor of the pens was covered with wood shavings. Each of the twelve
175 experimental diets was fed to 8 pens following randomisation. Feed and water were
176 provided *ad libitum* throughout the experimental period.

177 The temperature was gradually reduced daily until room temperature reached 23°C
178 when birds were 21 d old. A standard lighting programme for broilers was used, decreasing
179 from 23:1 (hours light: dark) from zero day old to 18 h: 6 h at 7 days of age, which was
180 maintained until the end of the study.

181 The experiment ended when the birds were 21 d of age. The birds were group-weighted
182 on a per pen basis at the beginning (8 d old) and at the end of the study (21 day old), and
183 the average daily feed intake (FI) and bird weight gain (WG), and feed conversion ratio
184 (FCR) were determined over this time.

185 At the beginning of 18 d, the solid floor of each pen was replaced with a wire mesh and
186 all excreta were quantitatively collected daily in a plastic tray over the four final days of the
187 experiment, from 18 to 21 d age. Feed intakes were also measured for the same period.
188 The freshly collected total excreta output of each pen was immediately dried in a forced
189 draft oven at 60°C to a constant weight and then left at room temperature for three days
190 followed by weighing.

191

192 ***Determination of dietary metabolisable energy and nutrient retention coefficients***

193 The dried excreta, as well as representative balancer diet samples were ground to pass
194 through 0.8 mm screen. The dry matter, gross energy, nitrogen and fat of each dried excreta
195 and the balancer diet samples were determined in duplicate as described for the field bean
196 samples earlier.

197 The N-corrected apparent metabolisable energy (AMEn) of the diets was determined
198 using total collection technique as described by Hill and Anderson (1958). The coefficients
199 of total tract fat retention (FR), nitrogen retention (NR) and dry matter retention (DMR) were
200 determined as the difference between intake and excretion of the nutrient divided by its
201 respective intake.

202

203 Statistical analysis

204 The observational unit was the raised-floor pen with 5 birds. Statistical analyses were
205 performed using the Genstat 18th statistical software package (Genstat 17 release 3.22 for
206 Windows; IACR, Rothamstead, Hertfordshire, UK). The AMEn and the nutrient utilisation
207 coefficients of the experimental field bean samples were statistically compared using a
208 randomized block analysis of variance. The position of pens within the room was used as
209 the blocking factor. Tukey's range test was used to determine significant differences
210 between field bean treatment groups.

211 Regression analysis was used in order to test linear response of AMEn and nutrient
212 utilisation to inclusion rates of bean samples. Then the AMEn and values of the nutrient
213 retention coefficients were obtained using the slope ratio method (Finney 1978).

214 The coefficients of correlation between all studied variables were also obtained.

215

216

217 Results

218

219 Proximate analysis and gross energy in field beans

220 There were differences in the chemical composition and GE among the studied field bean
221 cultivar samples (Table 2). The overall means of protein, ash, oil and GE of the beans were
222 282.4, 35.9, 10.8 g/kg DM and 18.46 MJ/kg DM, respectively.

223 Generally, the GE contents were quite similar between different cultivars, ranged from
224 18.27 (Bazz and Sultan) to 18.60 MJ/kg DM (Divine), indicating a difference of 0.33 MJ, and
225 with a coefficient of variation (CV = 0.7%). Ether extract was the most variable nutrient (CV
226 = 10.1%), although mean levels were low. Crude protein concentration had intermediate
227 variability (CV = 6.5%), and the difference between cultivar samples was approximately 60
228 g/kg DM.

229

230 Carbohydrates and minerals

231 The carbohydrate profiles of the field bean samples are displayed in Table 3. The major
232 component of field bean carbohydrates was starch and its average content in the studied
233 cultivars was 456 g/kg DM (CV = 7.4%). There was a mean of 182 g/kg of total NSP in the
234 bean samples (CV = 15.8%) and 72% was insoluble NSP.

235 The predominant constituent sugars of NSPs were glucose, galacturonic acid, arabinose
236 and xylose, respectively. Whereas, the levels of total galactose and mannose were low, and
237 both rhamnose and fucose were scarce.

238 Soluble galacturonic acid ranged from 10.1 to 20.3 and glucose from 1.5 to 25 g/kg DM
239 in Maris Bead and Clipper, correspondingly, and soluble arabinose scored 7.6 (MB) to 17.6
240 g/kg DM (Honey). The concentrations of 62.8 to 125.7 of insoluble glucose, and 7.1 to 14.1
241 g/kg DM of insoluble galacturonic acid were found in Honey and Clipper, respectively.

242 The mineral contents of the studied field bean cultivars are summarized in Table 4. The
243 concentrations of calcium, magnesium, potassium, sodium, sulphur and boron were similar
244 among the cultivars. Phosphorus concentration varied between 4.33 to 6.87 g/kg DM for
245 Arthur and Wizard, respectively. Copper content was variable between samples with a CV
246 = 26.3%.

247

248 ***Phenols, tannins, phytate and trypsin inhibitors***

249 Total phenols, tannins, non-tannin phenols, condensed tannins, phytate and trypsin inhibitor
250 contents of the studied field bean cultivars are presented in table 5. The majority of phenolic
251 compounds in the field beans were tannins and non-tannin phenols were low. The mean
252 total tannin concentration was 5.11 mg/g with a CV of 34.3%. Non-tannin phenol contents,
253 as tannic acid equivalent, were 2.02 mg/g (CV = 35.0%). The mean of condensed tannin
254 (CT) contents, as leucocyanidin equivalents, in bean cultivars was 5.04 mg/g DM (CV =
255 30.9%). The overall mean of phytate was 14.5 mg/g (CV = 24.6%), although for trypsin
256 inhibitors it was 3.5 mg/g (CV = 19.2%).

257

258 ***Grain quality and viscosity of the experimental bean samples***

259 Color score of bean flour is illustrated in table 6. The range of lightness scores was from 88
260 to 95 (CV = 2.4%). The a* (redness-greenness degree) of bean flour varied from 0.99 to
261 1.44 (CV = 11.7%). The overall mean for b* (yellowness-blueness degree) of bean flour was
262 18 (CV% = 10.6).

263 Thousand-grain weight (TGW), water holding capacity (WHC), viscosity, cotyledon ratio
264 and seed-coat ratio of the characterized field bean samples are also presented in table 6.
265 The mean of WHC of the field bean samples was 954 g/kg DM (CV = 4.5%). The average
266 of seed-coat proportion was 136 g/kg (CV = 10.1%). Viscosity (cP) of field beans was
267 variable (CV = 25.8%) with a range from 2.07 to 5.01 cP.

268

269 ***Analysis on data from the animal phase***

270 The data from basal diet and the diet contained 10% Honey field bean cultivar were used
271 for testing linearity only and was not presented in tables with data on beans only.

272

273 ***Linearity***

274 There was a linear change in AMEn and DMR ($P < 0.001$) to the three different inclusion
275 rates of Honey bean sample, thus demonstrating the validity of the slope-ratio method that
276 was employed for determination of these variables in the field bean cultivar samples (Table
277 7).

278

279 ***Broiler growth performance, available energy and nutrient utilisation of field beans***

280 There were no mortalities and all birds survived the experiment. The variation in daily FI,
281 WG and FCR were in the expected range for broiler chickens reared from 7 to 21 d old in
282 group pens of 5 birds (coefficient of variation (CV = 5.3%, 6.0%, and 2.5%, respectively)
283 (Table 8). Compared to breeder's recommendation, daily FI was approximately 10 g/day
284 lower probably due to the feed being in a meal form rather than pelleted. There were no
285 significant differences ($P > 0.05$) in daily FI and WG. The overall FCR was in the expected
286 range (Aviagen Ltd., Edinburgh, UK), as Divine gave a better ($P < 0.001$) FCR comparing
287 to Buzz and Sultan, but did not differ ($P > 0.05$) from the rest.

288 The results on the AMEn, CPR, FD and DMR coefficients of the field bean cultivar
289 samples are presented on table 8. The AMEn ranged from 7.78 to 9.96 MJ/kg DM. The
290 large ranged AMEn was due to the AMEn of one sample (Sultan) that was highly
291 significantly lower ($P < 0.001$) than that of Divine, Fury, Fuego and Wizard, field bean
292 cultivar samples. There were no significant ($P > 0.05$) differences between the other nine
293 samples except that the AMEn of Buzz was highly significantly ($P < 0.001$) lower than that
294 of Divine. There were no differences ($P > 0.05$) in NR and FR between the studied field
295 bean cultivar samples.

296

297 ***Relationship between chemical composition and physical characteristics of the***
298 ***beans, beans available energy and nutrients, and chicken growth performance***

299 Selected correlation coefficients obtained using all the data from the laboratory analysis and
300 broiler experiments are presented in Table 9. The CP content was correlated positively (P
301 < 0.05) to determined AMEn and lightness-darkness degree. Similarly, the lightness-
302 darkness degree correlated positively ($P < 0.001$) to AMEn. There was a positive correlation
303 ($P < 0.001$) between tannins content and yellowness-blueness degree.

304

305 **Discussion**

306 The purpose of this experiment was to determine the range of variation in energy and
307 nutrient availability of ten UK grown field bean cultivar samples. In addition, it aimed to
308 determine how their AMEn and nutrient utilization relates to their chemical and physical
309 composition.

310

311 ***Broiler performance***

312 The overall final body weight of the birds in all dietary treatments was in the expected range
313 for Ross 308 broilers fed on mash diets (Pirgozliev et al. 2015; Abdulla et al. 2016 a, b) as
314 FCR was similar to breeder's recommendations (Aviagen Ltd, Edinburgh). The differences
315 in birds feed intake and growth were not statistically significant in this study, although there
316 were some differences in FCR. This is in agreement with previous reports (Metayer et al.

317 2003; Nalle et al. 2010a) when similar amount of dietary beans were fed to broilers for the
318 same feeding period. The lack of response of growth performance variables to dietary bean
319 cultivars reported by Usayran et al. (2014) may be due to the relatively short feeding period
320 (7 days only) and the use of tannin-free bean cultivars.

321

322 ***Energy availability of field beans for broilers***

323 The overall determined AMEn value of the field beans was 9.22 MJ/kg DM, which is similar
324 to other reports with broilers (Nalle et al. 2010a, b; Lacassagne et al. 1991). The AMEn for
325 Sultan was numerically the lowest and significantly lower than that of Divine, Fury, Fuego
326 and Wizard field bean cultivar samples. Sultan had one of the lowest seed sizes, the lowest
327 proportion of cotyledon, and one the greatest proportions of seed-coat in its overall seed
328 composition. However, this did not result in a lower starch content or increased NSP content
329 compared to the other samples, although protein content was low. The AMEn of Sultan was
330 1.6 MJ/kg lower than the mean of the other nine samples. The reduced protein content in
331 Sultan would only account for approximately half of this lowered AMEn. Sultan had the
332 highest polyphenol and tannin contents of the ten samples. However, it is well documented
333 that tannin content in field beans reduces their AMEn (Brufau et al. 1998; Lacassagne et al.
334 1988; Vilariño et al. 2009). Similarly, tannin content in beans was associated with reduced
335 dietary nitrogen retention (Marquardt and Ward 1979) and ileal amino acid digestibility (Ortiz
336 et al. 1993; Woyengo and Nyachoti 2012). In agreement with this report, Igbasan et al.
337 (1997) observed higher metabolizable energy contents in light (both yellow and green) pea
338 cultivars than those in dark (brown) ones when fed mature cockerels. It has been found that
339 pale legume seeds have higher nutritive value than dark seeds. It has been noted that seed-
340 coat colour has some connection with the level of one or more anti-nutrients in field beans.
341 In comparison with light coloured cultivar samples, slightly high amounts of phenols and
342 tannins (Helsper et al. 1993; Oomah et al. 2011), phytate (Rubio et al. 1992) and fibres, but
343 lower CP (Helsper et al. 1993; Duc et al. 1995) in dark coloured field beans has been
344 reported. It has been known these compounds decrease the feeding quality of feedstuffs

345 for monogastric animals. In addition, Brufau et al. (1998) and Vilariño et al. (2009) reported
346 negative relation between tannin level and metabolizable energy contents in field beans.
347 The results of this experiment have indicated that these ten field bean cultivar samples had
348 different energy and nutrient availabilities. The commercial poultry industry requires broiler
349 diets to have high energy densities. Nutritionists will only be able to incorporate significant
350 amounts of field beans in poultry diets if the beans have a high metabolizable energy. It is
351 crucial that they are able to identify and only use samples with high metabolizable energy.
352 The results of the present experiment have shown that there is an excessively large range
353 in the determined metabolizable energy of ten different UK field bean samples. There is an
354 indication from these samples that the metabolizable energy of different field bean cultivar
355 samples can be predicted by their crude protein contents and colour (L^*). These
356 characteristics of field bean samples could be used as a rapid test of their nutritive quality.
357 However, the significant relationships were predominantly influenced by the physical
358 characteristics of only one field bean sample (Sultan). The relationship was not significant
359 in the remaining nine samples, even though the large range in metabolizable energy still
360 remained. This information may be a guide to plant breeders who may be able to incorporate
361 it in the development of new field bean cultivars.

362

363

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367

368 **Disclosure statement**

369 No potential conflict of interest was reported by the authors.

370

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374

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521 **Table 1.** Ingredient composition (g/kg, as-fed basis) of the experimental broiler chicken
 522 balancer formulation.

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Ingredient	Balancer feed (g/kg)
Wheat	533.2
SBM (CP=48%)	150
Full fat Soy meal	175
Maize gluten meal	37.4
Soy oil	50
Lysine	1.9
Methionine	6.3
Threonine	1.9
Monocalcium phosphate	20
Limestone	15.5
Sodium chloride	3.8
Vitamin/mineral premix*	5
	1000
Determined composition	
ME (MJ/kg)	13.71
Protein (g/kg)	231
Lysine (g/kg)	12.4
Met + Cys (g/kg)	11.1
Calcium (g/kg)	11.1
Phosphorus available (g/kg)	8.5
Sodium (g/kg)	2.0

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525 ME = metabolisable energy.

526 This balancer was fed as a part of complete diet comprised 200 g/kg of each experimental field bean sample
 527 and 800 g/kg of the balancer. Each experimental diet met the diet specification for this strain of broiler chicken
 528 (Aviagen Ltd, Edinburgh, UK).

529 *The vitamin and mineral premix contained vitamins and trace elements to meet breeder's recommendation
 530 (Aviagen Ltd., Edinburgh, UK). The premix provided (units/kg diet) retinol, 3600 µg; cholecalciferol, 125 µg; µ-
 531 tocopherol, 34 mg; menadione, 3 mg; thiamin, 2 mg; riboflavin, 7 mg; pyridoxine, 5 mg; cobalamin, 15 µg;
 532 nicotinic acid, 50 mg; pantothenic acid, 15 mg; folic acid, 1 mg; biotin, 200 µg; iron, 80 mg; copper, 10 mg;
 533 manganese, 100 mg; cobalt, 0.5 mg; zinc, 80 mg; iodine, 1 mg; selenium, 0.2 mg; and molybdenum, 0.5 mg.

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542 **Table 2.** The chemical composition (dry matter basis) of ten UK grown studied field bean

543 cultivars.

Bean cultivar	Dry matter (g/kg)	Ash (g/kg)	Ether extract (g/kg)	Crude protein (g/kg)	Gross energy (MJ/kg)
Arthur	859	32.0	11.6	270.6	18.41
Buzz	845	38.2	10.7	276.0	18.27
Clipper	854	35.6	9.4	284.8	18.38
Divine	866	38.6	9.2	299.6	18.60
Fuego	855	34.3	12.9	269.8	18.58
Fury	856	33.8	10.5	281.0	18.56
Honey	836	34.7	10.8	293.8	18.56
Maris Bead	858	33.5	10.5	304.5	18.41
Sultan	856	39.4	11.7	244.6	18.27
Wizard	855	38.8	10.5	299.7	18.59
CV%	1.0	7.4	10.1	6.5	0.7

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Table 3. Carbohydrate composition (g/kg DM) of ten UK grown studied field bean cultivars.

Bean cultivar	Fraction	NSP constituent sugars								Total NSPs	Total starch
		<i>Rha</i>	<i>Fuc</i>	<i>Ara</i>	<i>Xyl</i>	<i>Man</i>	<i>Gal</i>	<i>Glu</i>	<i>GalA</i>		
Arthur	Soluble sugar	1.0	0.7	10.2	4.6	2.6	5.3	11.6	14.3	50.3	488
	Insoluble sugar	0.0	0.2	13.4	10.4	2.8	2.8	60.7	8.0	98.3	
	Total sugar	1.0	0.9	23.5	15.0	5.4	8.1	72.4	22.3	148.6	
Buzz	Soluble sugar	0.6	0.4	12.0	5.3	2.2	5.0	10.6	14.5	50.6	452
	Insoluble sugar	0.4	0.6	12.5	14.4	4.1	3.4	91.9	11.9	139.2	
	Total sugar	1.0	1.0	24.5	19.8	6.2	8.4	102.5	26.4	189.7	
Clipper	Soluble sugar	1.3	0.7	10.4	5.8	2.6	6.8	25.0	20.3	72.8	397
	Insoluble sugar	0.0	0.5	13.1	14.6	5.9	3.9	125.7	14.1	177.6	
	Total sugar	1.3	1.2	23.4	20.3	8.5	10.6	150.7	34.3	250.4	
Divine	Soluble sugar	1.1	0.9	9.6	5.2	2.5	5.8	8.5	12.9	46.4	434
	Insoluble sugar	0.0	0.0	11.7	15.0	3.4	3.9	89.2	10.8	134.0	
	Total sugar	1.1	0.9	21.3	20.2	5.9	9.7	103.6	23.6	180.4	
Fuego	Soluble sugar	1.0	0.5	9.7	5.5	2.2	5.0	15.8	14.4	54.1	473
	Insoluble sugar	0.0	0.5	13.1	12.1	4.1	3.8	73.2	10.1	116.9	
	Total sugar	1.0	1.0	22.9	17.5	6.3	8.8	89.0	24.4	171.0	
Fury	Soluble sugar	1.1	0.8	9.7	5.4	2.1	4.8	5.8	14.5	44.1	464
	Insoluble sugar	0.0	0.2	12.1	15.5	4.2	3.5	91.4	9.4	136.4	
	Total sugar	1.1	1.0	21.8	20.9	6.3	8.4	97.2	23.9	180.5	
Honey	Soluble sugar	1.1	0.7	17.6	5.6	6.9	6.9	7.2	16.9	62.9	517
	Insoluble sugar	0.0	0.3	11.1	10.0	2.1	2.6	62.8	7.1	95.9	
	Total sugar	1.1	1.0	28.7	15.6	8.9	9.5	70.1	23.9	158.8	
Maris Bead	Soluble sugar	0.9	0.7	7.6	2.8	1.4	4.9	1.5	10.1	30.0	443
	Insoluble sugar	0.2	0.2	12.5	11.4	4.2	3.3	80.9	12.7	125.5	
	Total sugar	1.1	0.9	20.1	14.3	5.6	8.2	82.3	22.8	155.5	
Sultan	Soluble sugar	1.0	0.4	9.7	3.7	2.1	5.4	15.4	17.1	54.8	467
	Insoluble sugar	0.0	0.5	11.4	8.2	4.6	3.1	96.1	11.6	135.4	
	Total sugar	1.0	0.9	21.0	11.9	6.6	8.5	111.5	28.7	190.2	
Wizard	Soluble sugar	0.8	0.5	11.1	3.6	2.0	5.6	4.9	14.2	42.8	424
	Insoluble sugar	0.3	0.4	11.8	15.8	5.0	3.2	101.8	12.1	150.4	
	Total sugar	1.2	0.9	23.0	19.5	6.9	8.8	106.7	26.3	193.2	
CV%	Soluble sugar	19.4	27.0	24.8	21.3	57.7	13.4	63.8	18.2	22.9	7.4
	Insoluble sugar	171.4	55.3	6.4	21.0	27.0	12.9	22.1	20.0	18.4	
	Total sugar	9.4	10.5	10.4	17.8	17.4	9.1	23.5	14.0	15.8	

Rha, rhamnose; Fuc, fucose; Ara, arabinose; Xyl, xylose; Man, mannose; Gal, galactose; Glu, glucose; GalA, galacturonic acid; Total-NSPs, total non-starch polysaccharides; Each value represents mean of duplicate analysis.

Table 4. Mineral composition (dry matter basis) of ten UK grown studied field bean cultivars.

Beans cultivar	Mineral										
	Calcium (g/kg)	Magnesium (g/kg)	Phosphorus (g/kg)	Potassium (g/kg)	Sodium (g/kg)	Sulphur (g/kg)	Boron (mg/kg)	Copper (mg/kg)	Iron (mg/kg)	Manganese (mg/kg)	Zinc (mg/kg)
Arthur	1.35	1.17	4.33	11.15	0.19	1.89	11.25	8.24	72.95	23.80	39.45
Buzz	1.34	1.41	6.64	12.75	0.12	1.58	10.35	19.50	72.60	31.10	47.55
Clipper	1.60	1.47	6.05	11.10	0.43	1.92	10.55	18.60	83.60	19.55	52.05
Divine	1.18	1.43	5.05	13.70	< 0.12	2.92	10.60	12.80	62.30	13.85	45.90
Fuego	1.07	1.34	4.96	11.85	0.30	2.12	11.50	13.90	65.05	11.10	48.65
Fury	1.00	1.17	5.07	11.75	0.31	2.12	11.25	12.75	58.75	10.70	44.70
Honey	0.82	1.37	5.26	11.80	< 0.12	2.00	11.40	11.75	48.60	11.25	45.40
Maris Bead	1.00	1.27	5.33	11.20	< 0.12	2.08	12.15	16.00	51.40	12.30	53.20
Sultan	1.19	1.42	4.61	13.10	< 0.12	1.47	10.65	9.54	67.55	23.60	64.30
Wizard	1.41	1.46	6.87	12.05	< 0.12	2.22	10.70	15.95	68.50	15.20	43.85
CV%	19.60	8.3	15.5	7.3	NA*	19.4	5.1	26.3	16.1	40.4	14.1

*NA, not applicable; Each value represents mean of duplicate analysis.

Table 5. Total phenols, tannins, non-tannin phenols (NTPH), condensed tannins, phytate and trypsin inhibitor contents (mg/g DM) of ten UK grown studied field bean cultivars.

Bean cultivar	Total phenols ^a	Tannins ^a	NTPH ^a	Condensed tannins ^b	Phytate	Trypsin inhibitors
Arthur	4.5	3.5	1.0	2.8	9.86	3.1
Buzz	4.7	2.2	2.5	2.9	20.84	2.6
Clipper	7.1	4.6	2.5	5.3	16.62	3.3
Divine	7.1	4.8	2.4	6.2	13.35	4.2
Fuego	8.3	6.1	2.3	6.8	12.90	4.4
Fury	6.3	4.3	2.0	4.7	13.77	3.7
Honey	7.3	4.4	2.8	3.9	13.51	3.4
Maris Bead	6.9	6.1	0.8	4.5	13.90	3.8
Sultan	10.9	8.3	2.6	7.3	10.63	2.3
Wizard	8.1	6.8	1.4	6	19.80	3.8
CV%	25.7	34.3	35.0	30.9	24.6	19.2

^aAs tannic acid equivalents; ^bAs leucocyanidin equivalents. Each value represents mean of triplicate analysis.

Table 6. Weight of 1000 grains (TGW), water holding capacity (WHC), water extract viscosity (WEV), cotyledon and seed-coat ratio and colour score (L*, a* and b*) of flour of ten UK grown field bean cultivars*.

Bean cultivar	TGW (g DM)	WHC (g/kg)	WEV (cP)	Cotyledon (g/kg)	Seed-coat (g/kg)	L*	a*	b*
Arthur	685	915	2.07	866.4	133.6	93.59	1.07	17.72
Buzz	693	871	2.41	868.9	131.1	91.49	1.27	14.69
Clipper	539	943	4.52	843.4	156.6	91.76	1.17	18.94
Divine	444	935	4.18	863.2	136.8	94.66	0.99	17.59
Fuego	466	1005	3.58	858.1	141.9	94.25	1.14	17.96
Fury	483	1010	4.59	863.7	136.3	95.16	1.21	18.22
Honey	754	956	4.81	889.8	110.2	94.63	1.06	17.04
Maris Bead	311	961	5.01	876.7	123.3	93.18	1.01	19.05
Sultan	407	997	4.04	844.5	155.5	87.71	1.44	22.29
Wizard	681	947	3.40	867.3	132.7	94.04	1.18	19.34
CV%	27.1	4.5	25.8	1.6	10.1	2.4	11.7	10.6

Each value of WHC and WEV represents mean of triplicate analysis.

* L*, lightness-darkness degree of bean flour; a*, redness-greenness degree of bean flour; b*, yellowness-blueness degree of bean flour; Each value represents mean of triplicate analysis.

Table 7. Linearity table.

Variable	Bean rate in the diets			SEM	P value		
	0.0%	10.0%	20.0%		Treatment	Linear	Quadratic
Total collection							
AMEn (MJ/kg_DM)	14.27	13.86	13.30	0.096	< 0.001	< 0.001	0.540
NR	0.625	0.623	0.621	0.0033	0.697	0.404	0.998
FR	0.749	0.756	0.771	0.0110	0.400	0.193	0.779
DMR	0.716	0.702	0.683	0.0048	< 0.001	< 0.001	0.644

SEM = pooled standard errors of mean; AMEn = N-corrected apparent metabolisable energy; NR = nitrogen retention coefficient; FR = fat retention coefficient; DMR = dry matter retention coefficient.

Table 8. Daily feed intake (FI), weight gain (WG) and feed conversion ratio (FCR) of broiler chickens fed on diets containing 200 g/kg of one of the ten different UK grown field bean cultivar samples. Nitrogen-corrected apparent metabolisable energy (AMEn), total tract fat (FR) retention, nitrogen (NR) and dry matter (DMR) retention coefficients (obtained with slope ratio method) of ten UK grown field bean cultivar samples fed to broiler chickens.

Diet	FI (g DM/b/d)	WG (g/b/d)	FCR (g:g)	AMEn bean (MJ/kg DM)	AMEn diet (MJ/kg DM)	NR	FR	DMR
Arthur	57.4	46.2	1.244 ^{abc}	9.19 ^{abc}	13.25 ^{abc}	0.596	0.752	0.546 ^{ab}
Buzz	59.3	47.3	1.254 ^{bc}	8.20 ^{ab}	13.06 ^{ab}	0.556	0.740	0.491 ^{ab}
Clipper	58.4	47.1	1.240 ^{abc}	9.16 ^{abc}	13.25 ^{abc}	0.594	0.903	0.528 ^{ab}
Divine	60.0	50.0	1.201 ^a	9.96 ^c	13.41 ^c	0.624	0.916	0.571 ^b
Fuego	57.7	46.9	1.233 ^{abc}	9.78 ^{bc}	13.37 ^{bc}	0.606	0.861	0.562 ^b
Fury	58.5	48.2	1.212 ^{ab}	9.84 ^{bc}	13.38 ^{bc}	0.572	0.868	0.566 ^b
Honey	58.8	48.3	1.220 ^{ab}	9.43 ^{abc}	13.30 ^{abc}	0.604	0.858	0.550 ^b
Maris Bead	59.5	48.7	1.221 ^{ab}	9.35 ^{abc}	13.29 ^{abc}	0.558	0.855	0.554 ^b
Sultan	57.4	45.1	1.274 ^c	7.78 ^a	12.97 ^a	0.538	0.850	0.461 ^a
Wizard	56.5	46.4	1.217 ^{ab}	9.52 ^{bc}	13.32 ^{bc}	0.556	0.879	0.547 ^{ab}
Mean	58.3	47.4	1.232	9.22	13.26	0.580	0.848	0.538
CV%	5.3	6.0	2.5	11.3	10.2	10.3	13.8	9.8
SEM	1.10	1.01	0.0110	0.370	0.074	0.0211	0.0415	0.0186
<i>P</i> value	0.455	0.060	< 0.001	< 0.001	<0.001	0.091	0.061	0.001

Each value represents mean of eight replicate pens of 5 chicks each; Bird performance was determined from 7 to 21 d age; AMEn and retention coefficients were determined from 17 to 21 d age; ^{a,b,c}Values within a column with different superscripts differ significantly at $P \leq 0.05$.

Table 9. Selected correlation coefficients between determined AMEn and laboratory analysis of field bean cultivars.

	<i>AMEn</i>	<i>L</i>	<i>a</i>	<i>b</i>	<i>Starch</i>	<i>CP</i>	<i>NSP tot</i>	<i>NSP n</i>	<i>NSP s</i>
<i>L</i>	0.924								
<i>a</i>	-0.762	-0.768							
<i>b</i>	-0.220	-0.500	0.400						
<i>Starch</i>	-0.040	0.159	-0.048	-0.186					
<i>CP</i>	0.658	0.683	-0.772	-0.338	-0.294				
<i>NSP tot</i>	-0.197	-0.364	0.396	0.189	-0.757	-0.073			
<i>NSP n</i>	-0.138	-0.314	0.354	0.238	-0.916	0.080	0.918		
<i>NSP s</i>	-0.202	-0.250	0.246	-0.028	0.026	-0.345	0.571	0.197	
<i>Tannins</i>	-0.111	-0.371	0.312	0.894	-0.129	-0.201	0.050	0.137	-0.162

$df = 7$; $P < 0.05$ ($r^2 \geq 0.632$; $0.765 \leq r^2$); $P < 0.001$ ($r^2 \geq 0.765$).

AMEn, nitrogen-corrected apparent metabolisable energy; *L**, lightness-darkness degree of bean flour; *a**, redness-greenness degree of bean flour; *b**, yellowness-blueness degree of bean flour; *Starch*, (g/kg DM); *CP*, crude protein in beans (g/kg DM); *NSP tot*, *NSP n* and *NSP s*, is respectively total, non-soluble and soluble non-starch polysaccharide contents in beans (g/kg DM); *Tannins*, as tannic acid equivalents, content in beans (mg/g DM).