



Nutritive evaluation, metabolisable energy and digestible amino acid contents of different indigenous feedstuff for Nile tilapia (*Oreochromis niloticus*)

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Abstract

Three trials were executed to examine the nutritive profile, metabolisable energy and digestible amino acid (AA) contents of four indigenous feed ingredients including wheat (W), wheat middling (WM), canola meal (CM) and rapeseed meal (RSM) in Nile tilapia. Three samples of each test ingredient were collected from three different locations of Multan (MUL) and Sukkar (SKR), of Pakistan. The collected three samples were pooled thereafter to make a homogenous/ representative sample of each test ingredient from a particular study site. Nutrients composition, AA and energy digestibility of these indigenous ingredients were evaluated by using laboratory analyses and fish studies. Proximate analysis indicated variations in some of the nutrients due to location ($p < 0.05$). Differences were also observed in some AA including arginine, lysine, serine, cysteine, glutamic and aspartic acids, histidine, valine and glycine contents of these ingredients ($p < 0.05$). Digestibility of leucine, glycine and glutamic acid was higher ($p < 0.05$) in RSM from MUL. Among W samples from MUL, AA digestibility for lysine, threonine, and aspartic acid was higher ($p < 0.05$). Crude protein, arginine, alanine, serine, and aspartic acid had higher digestibility ($p < 0.05$), whereas digestibility was lower ($p < 0.05$) for threonine, valine and tyrosine in RSM from MUL. Metabolisable energy contents did not differ among W, WM, CM and RSM regarding their origin ($p > 0.05$). The results indicated that nutritional profiles and their digestibility indices vary with the location for Nile tilapia.

Keywords: Indigenous ingredients, amino acid, energy, digestibility, Nile tilapia.

Avaliação nutricional, energia metabolizável e conteúdo de aminoácidos digestíveis de diferentes alimentos indígenas para tilápia do Nilo (*Oreochromis niloticus*)

Resumo

Três experimentos foram executados para examinar o perfil nutritivo, a energia metabolizável e o conteúdo de aminoácidos digestíveis (AA) de quatro ingredientes alimentícios, incluindo trigo (W), farelo de trigo (WM), farelo de canola (CM) e farelo de colza (RSM) em tilápia do Nilo. Três amostras de cada ingrediente do teste foram coletadas de dois locais diferentes (Multan (MUL) e Sukkar (SKR), do Paquistão) e assim agrupadas. A composição nutricional, AA e digestibilidade energética desses ingredientes indígenas foram avaliadas por meio de análises laboratoriais e estudos de peixes. A análise imediata indicou variações ($p < 0,05$) em alguns dos nutrientes devido à localização. Variações ($p < 0,05$) também foram observadas em alguns teores de AA desses ingredientes. A digestibilidade da leucina, glicina e ácido glutâmico foi maior ($p < 0,05$) em RSM de MUL. Entre as amostras de W da MUL, a digestibilidade de AA para Lys, Thr e Asp foi maior ($p < 0,05$). Proteína Crud, arginina, alanina, serina e ácido aspártico apresentaram maior digestibilidade ($p < 0,05$), e menor ($p < 0,05$) para treonina, valina e tirosina em MRS. Nenhuma diferença ($p > 0,05$) relacionada a energia metabolizável foi observada entre esses ingredientes em relação à sua origem. Os resultados indicaram que os perfis nutricionais e sua digestibilidade variam com a localização.

Palavras-chave: ingredientes indígenas, aminoácidos, energia, digestibilidade, tilápia do Nilo.

1. Introduction

In aquaculture production, feed cost accounts for about 50 to 80% of the total production cost (FAO, 2017), where crude protein and energy being the most expensive ones.

The major energy sources used in fish feed are cereal grains including corn that contribute up to 50% of the fish diets (Marković et al., 2016). The higher inclusion

level of corn in fish diets is not economical because of its higher production cost with seasonal fluctuation, less availability in shortage periods (Dec-Jan and May-June) and more demand by the feed industry (NARC, 2017). Additionally, corn is also used for human food as corn flour, corn oil and also in silage production for animals. For least cost feed formulation, therefore, there is a dire need of alternative, cheaper and readily available energy sources including wheat, sorghum, wheat middling and barley for aquaculture feed.

The use of highly digestible protein sources including fish, shrimp and soybean meals in aquaculture feed may lead to a higher feed cost. There is a tremendous pressure on nutritionists, therefore, to find out substitute of the mentioned protein sources in aquaculture production. The possible low-cost alternative energy and protein sources of corn and SBM includes wheat (W), wheat middling (WM), canola meal (CM) and rapeseed meal (RSM) in fish diets (Toghyani et al., 2015). Canola is an improved variety of rapeseed with lower concentration of erucic acid (<2% in oil) and glucosinolates (< 30µmol/g) in the defatted meal (Maison and Stein, 2014). This improved variety is called “double zero” and “canola” in Europe and Australia, and North America, respectively (Newkirk, 2009). High fiber content (12%) in CM decreases energy and protein digestibility resulting in a lower AMEn (Khajali and Slominski, 2012). Nutritional values of different cereal grains are determined primarily by chemical composition (Fairbairn et al., 1999), and these results may be applied to forecast nutritional profile precisely. The nutritional profile of an ingredient is interrelated with several factors including variety, agronomic practices, geographical locations, environmental circumstances, harvesting conditions and processing of the seed and meal (Daun et al., 2011). Nutritional contents, however, differ between various ingredients, with different digestibility and availability to aquatic animals. Data regarding the nutritional profile including digestible energy and other essential nutrients including protein as well as AA is essential for precise feed formulation and to control the aquaculture wastes (Zhou et al., 2004; Liu et al., 2009). Measuring the total, as well as bioavailable protein and AA, are important to assess the quality of an ingredient. The bioavailable AA is described as the amount of AA that will be digested, absorbed and used in the animal or birds body (Ravindran and Bryden, 1999).

There is a scarcity of published data regarding the apparent digestibility coefficients of amino acids and metabolisable energy of indigenous protein and energy sources in tilapia fish. The objectives of the study were, therefore, to evaluate the nutritive profiles, especially gross energy, protein and AA contents, apparent metabolisable energy (AME), and apparent digestibility of protein and AA of indigenous wheat, wheat middling, canola meal and rapeseed meal from different origin in tilapia fish.

2. Materials and Methods

In total, 3 samples of each indigenous energy (wheat (W) and wheat middling (WM)) and protein source (canola meal (CM) and rapeseed meal (RSM)), were collected from two diverse topographical sites i.e. Multan (MUL), Punjab and Sukkur (SKR), Sindh of Pakistan and pooled thereafter.

2.1. Fish stock for AME of indigenous energy sources

A total of 300 sex-reversed Nile tilapia (*Oreochromis niloticus*) of about 5 g body weight were reared on a commercial diet for four weeks (28 days) with feed at 10% of their body weight. The fish were maintained in 15 cylindrical fiber glass tanks (20 fish in each) with 450 L volume containing individual aeration system, water supply and central drainage with 100% water renewal of at least two times per day. Tanks were used as a flow-through metabolic chambers. To minimize environmental fluctuation, tanks were placed in an indoor room and the water temperature was maintained at 26 to 28 °C by using water submerged heaters. The pH (7.2-8.5) and dissolved oxygen (DO), maintained above 5.0 mg L⁻¹, in water were measured daily through portable pH meter (STARTER 300pH portable, Ohaus company) and DO meter (STARTER 300D portable, Ohaus company).

2.2. Diets and excreta collection

For four weeks, four experimental diets of W and WM from two different origins, with one reference diet were provided to fish (Table 1). Each diet was randomly assigned to three tanks with 20 fish/ tank in a completely randomized design. All the experimental diets were offered in mash for seven days as an adaptation period for fish. After an adaptation period, fish were off feed for 24 hrs and fecal material from each tank was completely washed. The fecal samples were collected on daily basis in a 200 mL capacity plastic container and immediately stored. At the end of the collection period, the samples from experimental diets and excreta were analysed for DM and gross energy (GE) contents with the help of a bomb calorimeter. The determination of metabolisable energy of feedstuff was carried out using AOAC (2000).

2.3. Fish stock for amino acid digestibility of indigenous energy sources

A total of 300 sex-reversed tilapia (*Oreochromis niloticus*) individuals of about 33 gm on an average weight were randomly distributed among 15 tanks containing 20 fish each on a commercial diet for three days. The physical and chemical properties of water were same as mentioned above in section 2.1 of the manuscript.

2.4. Diets and excreta collection

The experimental diets (Table 2), comprising of W and WM from two different origins, and a protein-free diets were offered to fishes. All the diets were randomly assigned to 15 tanks with three replicates of each. After an adaptation period of 3-days, the fishes were fasted

for 24 hrs and randomly assigned to experimental and a protein-free diet. The experimental diets were provided at 5% live weight to all experimental groups three times daily with equal intervals for 14 days.

The fecal samples were collected on daily basis in plastic bottles. The collected fecal samples, of each experimental unit comprising of 20 fish, were grouped into a plastic container of 200 mL. All the collected samples were stored in air tight containers at -4 °C for laboratory analysis.

2.5. Fish stock for protein and amino acid digestibility of indigenous protein sources

In this trial, 20 fish were maintained in each of the 15 cylindrical fiberglass tanks with 450 L capacity and containing individual aeration system, water supply, and bottom outflow. Tanks were used as a flow-through metabolic chambers.

2.6. Diet and excreta collection

Four diets of indigenous protein sources (Table 3) comprising of RSM and CM of two different origins, and one reference protein-free diet were formulated. Chromium oxide, an indigestible marker, was used in all the diets with 1% inclusion level. The diets were allocated to tanks, fecal samples were collected and stored as described above in section 2.4 of the manuscript.

2.7. Proximate analysis

All chemical analyses of the ingredients, feed and excreta samples were done in duplicate. Dry matter, crude fiber, ether extract, ash and energy (bomb calorimetry) were evaluated according to the procedures described in AOAC (2000). Kjeldhal or semi-micro Kjeldhal method (AOAC 2000) was used for nitrogen content, and CP contents were determined by multiplying nitrogen with 6.25. Chromic

Table 1. Composition of the reference and experimental diets for apparent metabolisable energy measurements in tilapia.

Ingredients (%)	RD	W (MUL)	W (SKR)	WM (MUL)	WM (SKR)
SBM	60.0	39.0	39.0	39.0	39.0
Corn	28.0	19.0	19.0	19.0	19.0
Wheat	0.00	30.0	30.0	0.00	0.00
Wheat Middling	0.00	0.00	0.00	30.0	30.0
Oil	10.0	10.0	10.0	10.0	10.0
Premix	1.00	1.00	1.00	1.00	1.00
DCP	1.00	1.00	1.00	1.00	1.00
Total	100.00	100.00	100.00	100.00	100.00

RD: reference diet; W: wheat; MUL: Multan; SKR: Sukkur; WM: wheat middling; SMB: soybean meal; DCP: di-calcium phosphate.

Table 2. Dietary ingredients and calculated nutrient composition for protein and AA digestibility in tilapia.

Ingredients (%)	RD	W (MUL)	W (SKR)	WM (MUL)	WM (SKR)
SBM	55.5	36.39	36.39	36.39	36.39
Corn	35.0	24.1	24.1	24.1	24.1
Wheat	0.00	30.0	30.0	00.0	0.00
Wheat Middling	0.00	0.00	0.00	30.00	30.0
Oil	8.00	8.00	8.00	8.00	8.00
Choline	0.50	0.50	0.50	0.50	0.50
Salt	0.01	0.01	0.01	0.01	0.01
Premix	1.00	1.00	1.00	1.00	1.00
Total	100.00	100.00	100.00	100.00	100.00

RD: reference diet; W: wheat; MUL: Multan; SKR: Sukkur; WM: wheat middling; SMB: soybean meal.

Table 3. Dietary ingredients and calculated nutrient composition for protein and AA digestibility in tilapia.

Ingredients (%)	RD	CM (MUL)	CM (SKR)	RSM (MUL)	RSM (SKR)
SBM	55.5	36.39	36.39	36.39	36.39
Corn	34.9	24.1	24.1	24.1	24.1
Wheat	0.00	30.0	30.0	00.0	0.00
Wheat Middling	0.00	0.00	0.00	30.00	30.0
Oil	8.00	8.00	8.00	8.00	8.00
Choline	0.50	0.50	0.50	0.50	0.50
Salt	0.01	0.01	0.01	0.01	0.01
Premix	1.00	1.00	1.00	1.00	1.00
Total	100.00	100.00	100.00	100.00	100.00

RD: reference diet; MUL: Multan; SKR: Sukkur; SMB: soybean meal.

oxide was evaluated by the Scott (1978) method. Amino acids contents were analyzed following acid hydrolysis using an AA analyzer (Biochrom 30+, Biochrom Limited, Cambridge, UK).

2.8. Digestibility measurements

The apparent digestibility coefficients (ADCs) of energy, protein and AA for the test ingredients and diets were calculated as described by Cho and Slinger (1979):

$$ADC = 100 \times [1 - (F / D) \times (D_i / F_i)]$$

$$ADC_I = [ADC_T - (0.7 \times ADC_R)] / 0.3$$

Where, D = % nutrient or energy of a diet; F = % nutrient or energy of feces; Di = % marker (Cr₂O₃) in the diet; Fi = % marker (Cr₂O₃) in the feces, ADCT = % apparent digestibility coefficient of protein, amino acids or energy in the test diet; ADCR = apparent digestibility coefficient of nutrient or energy in the reference diet; I = test ingredient under investigation.

2.9. Data analysis

The data were analyzed statistically by ANOVA using PROC MIXED procedure of SAS (version 9.1, SAS Institute, 1985). Differences were considered significant at P < 0.05 and the significant differences between the means were separated by least significant difference test.

4. Results

4.1. Proximate analysis

The findings of proximate analysis for W, WM, CM, and RSM are shown in Table 4. There were significant variances between CP contents of energy sources (W and WM). Crude protein contents were significantly higher in W (p = 0.026) and WM (p = 0.036) from MUL than that from SKR. Similarly, significant variations were observed in ash contents of energy contents. Ash contents were significantly higher in SKR W (p = 0.01) and MUL WM (p = 0.01). Among protein sources, SKR CM (p = 0.042) and RSM (p = 0.023) has significantly greater DM compared with

that from MUL. Between protein sources, CP content of CM from SKR was significantly greater (p = 0.036) than those from MUL. Moreover, ash contents of protein sources were statistically greater in MUL than from SKR (p < 0.05). Canola meal (p = 0.021) and RSM (p = 0.026) from MUL had significantly greater ash contents compared with those from SKR. The GE contents of indigenous energy and protein sources were not affected by difference in origin except for CM. Gross energy contents of CM was found significantly greater (p = 0.022) in SKR compared with that from MUL.

4.2. Total amino acid contents of indigenous energy sources

The results of total amino acid contents of indigenous energy sources (W and WM) are summarized in Table 5. Among the essential amino acids (EAAs), Arg (p = 0.031) and Lys (p = 0.021) contents were significantly higher in W from SKR compared with that from MUL. Similarly, among non-essential amino acids (NEAAs) in W, Cys (p = 0.015), Glu (p = 0.030) and Asp (p = 0.012) was found significantly higher, whereas Ser (p = 0.04) contents were significantly lower in SKR that from MUL. Likewise, in WM from different origins, among different EAAs, Arg (p = 0.027), His (p = 0.033) and valine (p = 0.020) were found significantly higher in MUL compared with SKR. All other EAAs were statistically the same in two different origins of WM. Among NEAAs in WM, Gly (p = 0.017), Ser (p = 0.036) and Glu (p = 0.010) were significantly higher in MUL samples compared with those from SKR.

4.3. Digestible CP and AA contents of indigenous energy sources

The results of protein and amino acids digestibility of indigenous energy sources are shown in Table 6. Digestibility of CP was significantly influenced by the origin (p < 0.05) of the ingredients in WM. Wheat middling from MUL (p = 0.031) had a higher digestibility of CP compared with that from SKR. Wheat from MUL showed a significantly greater digestibility of Lys (p = 0.033) and Thr (p = 0.010) compared with those from SKR among essential AA.

Table 4. Proximate analysis (%) and gross energy contents (Kcal/kg) of the indigenous protein and energy sources for fish diet.

Item ¹	Canola Meal				Rapeseed Meal				Wheat				Wheat Middling			
	MUL	SKR	SEM	P-value	MUL	SKR	SEM	P-value	MUL	SKR	SEM	P-value	MUL	SKR	SEM	P-value
DM	89.0	90.1	0.3	0.042	89.1	90.7	0.2	0.023	92.3	92.4	0.21	0.24	92.1	92.0	0.11	0.65
CF	12.2	11.7	0.8	0.68	11.9	12.6	0.4	0.22	3.0	3.1	0.08	0.43	3.2	3.3	0.07	0.48
CP	35.2	36.8	0.5	0.036	36.4	37.2	1.1	0.25	11.3	10.1	0.11	0.026	17.3	16.4	0.22	0.036
EE	2.4	3.1	0.7	0.24	1.6	1.4	0.5	0.39	5.3	5.4	0.12	0.64	9.3	9.2	0.11	0.61
Ash	6.9	5.8	0.4	0.021	9.1	8.2	0.2	0.026	1.71	2.21	0.13	0.01	6.6	5.4	0.19	0.01
GE	4100	4250	15.6	0.022	4210	4260	20.8	0.171	4179	4190	12.4	0.36	4390	4430	20.3	0.37

¹DM: Dry matter; CF: Crude fiber; CP: Crude protein; EE: Ether extract; GE: Gross energy. Results described as means of triple observation. MUL: Multan; SKR: Sukkur; SEM: standard error of mean.

The digestibility of only Asp was significantly greater ($p = 0.04$) in MUL compared with that from SKR, among NEAAs. Similarly, in WM, among EAAs, digestibility of valine ($p = 0.01$) and Thr ($p = 0.030$) were significantly higher in SKR, whereas Arg digestibility was significantly

lower ($p = 0.021$) compared with that from MUL. Among NEAAs in WM, the digestibility of Ala ($p = 0.014$), Ser ($p = 0.010$) and Asp ($p = 0.034$) was significantly greater, whereas Tyr digestibility was significantly lower ($p = 0.030$) in MUL compared with that from SKR

Table 5. Amino acid profile of indigenous energy sources of different origins commonly used in tilapia fish feed.

Item	Wheat				Wheat middling			
	MUL	SKR	SEM	P-value	MUL	SKR	SEM	P-value
Essential AA								
Arginine	4.3	4.7	0.1	0.031	6.1	5.8	0.1	0.027
Isoleucine	3.1	3.2	0.2	0.44	3.0	2.6	0.2	0.32
Leucine	6.2	6.0	0.2	0.39	5.8	5.5	0.2	0.41
Lysine	2.7	3.1	0.1	0.021	3.7	3.5	0.2	0.56
Histidine	2.0	1.9	0.2	0.61	2.2	1.9	0.1	0.033
Methionine	1.5	1.4	0.1	0.53	1.2	1.1	0.2	0.62
Phenylalanine	4.3	4.0	0.2	0.24	3.2	3.0	0.2	0.47
Threonine	2.7	2.8	0.1	0.45	3.1	2.9	0.1	0.32
Valine	4.0	3.9	0.1	0.38	4.2	3.8	0.1	0.02
Nonessential AA								
Glycine	3.5	3.3	0.1	0.26	4.5	4.0	0.2	0.017
Alanine	3.2	3.0	0.2	0.71	4.1	3.8	0.2	0.42
Tyrosine	2.4	2.3	0.2	0.85	2.3	2.1	0.1	0.47
Serine	4.4	4.1	0.1	0.04	4.0	3.6	0.1	0.036
Cysteine	1.9	2.3	0.1	0.015	1.8	1.7	0.2	0.72
Glutamic acid	25.6	26.3	0.2	0.030	17.8	16.9	0.2	0.01
Aspartic acid	4.8	5.6	0.1	0.012	6.2	6.0	0.2	0.76

Results described as means of triple observation. Essential AA: essential amino acids. Nonessential AA: non-essential amino acids. MUL: Multan. SKR: Sukkur. SEM: standard error of mean.

Table 6. Apparent digestibility (%) of energy, crude protein and Amino acid indices of the indigenous energy sources of different origins in tilapia fish.

Item ¹	Wheat				Wheat Middling			
	MUL	SKR	SEM	P-value	MUL	SKR	SEM	P-value
Energy	68.9	67.2	1.8	0.45	65.7	67.2	1.4	0.620
CP	94.1	93.4	1.3	0.38	86.4	83.1	1.2	0.031
Essential AA								
Arginine	89.1	86.7	1.2	0.86	88.9	85.7	1.2	0.02
Isoleucine	94.2	95.4	1.5	0.35	82.8	84.7	1.5	0.62
Leucine	96.5	96.2	1.3	0.71	84.3	82.1	1.4	0.81
Lysine	93.4	91.8	1.1	0.033	85.7	84.3	1.5	0.93
Histidine	94.8	93.9	1.5	0.45	86.4	87.5	1.2	0.64
Methionine	96.2	95.7	1.8	0.62	84.0	83.5	0.9	0.73
Phenylalanine	95.2	94.8	1.4	0.51	81.3	83.8	1.4	0.80
Threonine	93.4	90.7	1.2	0.010	76.8	80.2	1.3	0.030
Valine	93.4	95.6	1.3	0.47	78.2	83.1	1.5	0.01
Nonessential AA								
Glycine	97.2	96.8	1.4	0.56	84.1	82.2	1.3	0.671
Alanine	96.4	97.2	1.7	0.81	81.8	78.2	1.6	0.014
Tyrosine	94.3	95.1	1.5	0.62	80.2	83.1	1.0	0.021
Serine	95.7	96.5	1.6	0.83	85.3	82.1	0.9	0.012
Cysteine	94.5	96.8	1.2	0.67	80.7	79.2	1.2	0.871
Glutamic acid	97.2	95.1	1.4	0.54	90.2	89.7	1.4	0.071
Aspartic acid	96.8	93.1	1.7	0.04	84.3	81.2	1.1	0.034

¹CP: Crude protein. AA: Amino acids. Results described as means of triple observation. Essential AA: essential amino acids. Nonessential AA: non-essential amino acids. MUL: Multan. SKR: Sukkur. SEM: standard error of mean.

4.4. Aggregate amino acid contents of indigenous protein sources

Table 7 represents the aggregate amino acid contents of commonly used indigenous protein sources (CM and RSM) of different origin. Among EAAs in CM from MUL, contents of Arg ($p = 0.031$) and Phe ($p = 0.035$) were significantly higher, whereas Ile ($p = 0.041$) was significantly lower compared with that from SKR. Among NEAAs of CM, Gly ($p = 0.031$) and Ser ($p = 0.021$) contents were statistically higher in SKR than that from MUL. Contents of all other AAs remained unaffected by origin. Likewise, in RSM, Met

($p = 0.022$) and valine ($p = 0.031$) contents were significantly higher among EAAs in SKR than that from MUL. Among NEAAs, Glu ($p = 0.021$) content was significantly higher in RSM from MUL, whereas Ser ($p = 0.010$) was significantly lower compared with that from SKR.

4.5. Digestible CP and AA contents of indigenous protein sources

Apparent digestibility coefficients of CP and AAs of indigenous protein sources (CM and RSM) are presented in Table 8. There was non-significant effect of origin on digestibility of essential and non-essential AAs of CM

Table 7. Amino acid profile of the indigenous protein sources of different origins commonly used in tilapia fish feed.

Item	Canola Meal				Rapeseed Meal			
	MUL	SKR	SEM	P-value	MUL	SKR	SEM	P-value
Essential AA								
Arginine	6.8	6.4	0.1	0.031	6.1	5.6	0.2	0.67
Histidine	2.8	2.6	0.2	0.68	2.5	2.8	0.2	0.53
Isoleucine	4.3	4.8	0.1	0.041	3.8	3.2	0.3	0.49
Leucine	6.9	7.1	0.2	0.52	6.5	6.1	0.1	0.83
Lysine	5.4	5.2	0.2	0.81	5.3	6.1	0.2	0.59
Methionine	2.4	2.5	0.1	0.75	1.98	2.21	0.1	0.022
Phenylalanine	5.8	5.2	0.1	0.035	5.6	6.2	0.1	0.63
Threonine	4.6	4.4	0.2	0.63	3.9	4.2	0.2	0.49
Valine	5.2	5.0	0.2	0.51	4.7	5.3	0.1	0.031
Nonessential AA								
Alanine	4.10	3.98	0.2	0.62	4.2	3.8	0.3	0.43
Aspartate	3.21	3.10	0.1	0.73	2.1	1.9	0.1	0.31
Cysteine	2.43	2.48	0.2	0.81	2.0	2.3	0.2	0.67
Glycine	4.51	4.67	0.1	0.031	4.9	5.2	0.2	0.61
Glutamic acid	6.42	6.18	0.3	0.49	16.5	14.3	0.3	0.021
Serine	4.02	4.23	0.1	0.021	4.1	4.8	0.2	0.010

Results described as means \pm SE of triple observation. Essential AA: essential amino acids. Nonessential AA: non-essential amino acids. MUL: Multan. SKR: Sukkur. SEM: standard error of mean.

Table 8. Apparent digestibility (%) of crude protein and amino acids of indigenous protein sources from different origin in tilapia fish.

Item	Canola Meal				Rapeseed Meal			
	MUL	SKR	SEM	P-value	MUL	SKR	SEM	P-value
CP	64.3	66.1	1.5	0.62	61.3	62.8	1.3	0.65
Essential AA								
Arginine	90.1	91.5	1.8	0.83	89.5	87.2	1.8	0.73
Histidine	90.4	88.4	2.1	0.89	88.3	87.8	1.6	0.81
Isoleucine	82.3	83.1	1.3	0.54	79.4	77.9	2.3	0.62
Leucine	85.4	85.0	1.1	0.67	84.3	80.1	1.2	0.021
Lysine	84.3	83.7	1.2	0.59	87.5	85.4	2.4	0.43
Methionine	88.4	88.0	0.9	0.72	84.1	83.2	2.0	0.59
Phenylalanine	86.1	85.9	1.8	0.91	80.3	77.9	1.7	0.63
Threonine	85.7	85.2	2.1	0.83	81.8	79.5	2.1	0.83
Valine	81.9	80.8	1.4	0.79	78.9	76.4	1.3	0.71
Nonessential AA								
Alanine	85.4	85.1	1.6	0.62	83.4	82.4	2.4	0.65
Aspartate	80.6	78.9	2.3	0.73	82.1	80.7	2.1	0.39
Cysteine	77.8	76.9	1.8	0.77	72.3	70.2	1.5	0.43
Glycine	83.1	82.8	2.7	0.81	80.1	76.1	1.3	0.023
Glutamic acid	90.7	90.1	2.0	0.92	88.5	80.7	2.3	0.015
Serine	84.3	84.0	1.8	0.75	82.7	80.7	1.8	0.72

Results described as means \pm SE of triple observation. Essential AA: essential amino acids. Nonessential AA: non-essential amino acids. MUL: Multan. SKR: Sukkur. SEM: standard error of mean.

from MUL and SKR. Significant increase, however, was observed in digestibility of Leu ($p = 0.021$), Gly ($p = 0.023$) and Glu ($p = 0.015$) in MUL RSM compared with SKR. Digestibility of all other AAs were not affected by origin ($p > 0.05$).

5. Discussion

5.1. Proximate analysis

Proximate composition of the evaluated WM is within the range reported in literature (Cromwell et al., 2000). Variations in CP and ash of W and WM from two different origins may be correlated with variety, agronomic characteristics, topographical positions and environmental conditions during crop development, harvesting conditions and processing of the seeds (Daun et al., 2011). Similarly, proximate composition of CM was in close agreement with values described in the literature (Selle and Ravindran, 2007; Rogiewicz et al., 2012). Significant discrepancies were, however, found between CM from different origins for their chemical constituents. In agreement with published data (Nyirenda et al., 1987), season and variety affects CP, CF, starch (energy) and ash contents in grains. The higher CP and energy contents in SKR samples compared with MUL samples, may be correlated with variety, different geographical locations, environmental conditions (Newkirk, 2009; Daun et al., 2011), season and site of cultivation (Conan et al., 1992; Metayer et al., 1993).

5.2. Contents and apparent digestibility of crude protein and amino acids

The evaluated aggregate AA contents of indigenous W were within range reported in literature (Abdel-Aal and Hucl, 2002; Ling et al., 2008). In the review by McNab (1996), it was reported that fertilizer application increases the total AA contents in wheat, which may be related to higher AAs contents in SKR sample. The CP digestibility of W was in close relation with the findings stated by Cheng and Hardy (2002). Digestibility of EAAs (Lys and Thr) of MUL W was observed to be greater than those from SKR. The digestibility of Lys was lowest despite a markedly high CP and TAA digestibility in wheat samples from SKR. This low total Lys content indicates that some of Lys and other AA may have been reduced because of grains drying process (Widyaratne and Zijlstra, 2008).

Variations in amino acids composition of WM samples may be attributed with variety, agronomic conditions, locations and environmental impact during crop development (Daun et al., 2011). According to Furuya et al. (2001) and Adewole et al. (2017), high CF and non-starch polysaccharides contents reduce time of contact between enzyme and substrate, hence reducing the digestibility that may be related to low digestibility values for WM compared with W.

In the present study, AA contents of indigenous CM samples were within range described in the literature (Guimarães et al., 2008). The AA digestibility of CM showed no difference regarding their origin. In addition,

according to published literature, average AA digestibility of CM was high (Forster et al., 1999), with Arg being the most digestible AA, whereas Cys being the poorer one. These higher digestibility values could be due to reduced erucic acid and glucosinolates contents in CM by applying heat treatment after oil extraction by hexane (Newkirk, 2002; Drew et al., 2007).

The AAs composition of RSM was in line with the findings stated in literature (Kasprzak et al., 2017; Ullah et al., 2017). Reduced AAs contents of RSM from MUL could be due to varietal or seasonal differences. Application of high temperature during the processing of rapeseeds to extract oil could also be a reason for lower AAs contents in RSM (González-Vega et al., 2011). In addition, existence of anti-nutritional factors (erucic acid and glucosinolates) in RSM may result in reduced digestibility of CP (Zhou et al., 2004) and AAs (Khajali and Slominski, 2012). The greater hemicellulose, cellulose and pectin in rapeseed hulls may bind with AA released during protein digestion and thereby decrease the AA absorption in the small intestine (Bjergegaard et al., 1991).

6. Conclusions

The findings of the current study showed various nutritional differences in the ingredients from different origins. These variations were, however, small in most of the nutrients. The variations were also observed in the digestibility values of different nutrients due to their origin. This generated database can be helpful for nutritionist formulating diets by using indigenous ingredients, for Nile tilapia and other fish species.

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