



## Rice Distillers Dried Grain Is a Promising Ingredient as a Partial Replacement of Plant Origin Sources in the Diet for Juvenile Red Seabream (*Pagrus major*)

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**ABSTRACT:** This study was designed to test the effects of dietary distillers dried grain (DDG) level on the growth performance, feed utilization, body composition and antioxidant activity of juvenile red seabream (*Pagrus major*). Six isonitrogenous and isocaloric diets were formulated to contain 0%, 5%, 10%, 15%, 20%, and 25% DDG from rice (designated as DDG0, DDG5, DDG10, DDG15, DDG20, and DDG25), respectively. Juvenile red seabream averaging  $10.1 \pm 0.05$  g were randomly distributed into 400-L tanks in a flow through systems. Three replicate groups of fish were fed one of the experimental diets to visual satiation two times a day for 10 weeks. Survival, weight gain, feed efficiency, protein efficiency ratio and hepatosomatic index of fish were not affected by dietary DDG levels ( $p > 0.05$ ). Proximate and amino acid composition of whole body in juvenile red seabream were not affected by dietary DDG levels ( $p > 0.05$ ). Plasma content of total protein, glucose, cholesterol, glutamic-pyruvic transaminase, phospholipid and triglyceride were not affected by dietary DDG levels ( $p > 0.05$ ). 1, 1-Diphenyl-2-picryl-hydrazyl radical and alkyl radical scavenging activities in plasma and liver of fish were not affected by dietary DDG levels ( $p > 0.05$ ). The results of this experiment suggest that DDG has the potential to replace plant origin ingredients such as wheat flour and corn gluten meal and could be used up to 25% in diet without incurring negative effects on the growth performance of juvenile red seabream. (**Key Words:** *Pagrus major*, Distillers Dried Grain, Growth, Antioxidant Activity)

### INTRODUCTION

In aquaculture diets, feed is generally the single largest expenditure for most aquatic species and it accounts for up to 60% of the total farm production cost (Cheng and Hardy, 2004). Due to the shortage and expense of many imported ingredients such as wheat flour (Rahman et al., 2013), there is additional merit in identifying a new alternative ingredient for target fish. Replacement of wheat flour with less expensive ingredients would be useful in minimizing feed cost. The success of reducing the expense of the prepared diet, without lowering fish performance, leads to a positive effect on the profitability of commercial fish

production.

Distillers dried grain (DDG) is a cereal by-product which is fermented and distilled to obtain alcoholic beverage (Hertrampf and Piedad-Pascual, 2000). The DDG has been documented as a promising feed ingredient in the poultry and livestock industries (Cheng and Hardy, 2004; Jacob et al., 2008) because it is highly nutritious and economical. Researchers have worked with incorporation of DDG in aquafeed (Chevanan et al., 2007,2010; Kannadhason et al., 2009). The enhanced availability and potential cost-benefit of DDG in aquafeed presents a substantial economic value as it is less expensive than other protein/energy sources like soybean meal (He et al., 2013). However, little information is available on the potential use of DDG from rice for red seabream. Our earlier study recommended that rice-based DDG may well offer an economical ingredient to make lower-cost feed and allow a

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Submitted Mar. 19, 2014; Revised May 31, 2014; Accepted Jul. 8, 2014

reasonable amount of wheat flour substitution in juvenile olive flounder, *Paralichthys olivaceus*, diets (Rahman et al., 2013). Research has shown that corn-based DDG is considered an acceptable ingredient in diets for some fish species such as rainbow trout (Cheng and Hardy, 2004), channel catfish (Lim et al., 2009; Li et al., 2010), tilapia (Shelby et al., 2008) and sunshine bass (Thompson et al., 2008).

Red seabream (*Pagrus major*) is one of the most valuable and widely farmed marine fish species in Asia and its production was established on a commercial scale more than 20 years ago (Foscarini, 1988). To sustain a productive and cost-effective red seabream culture operation, a reliable feed must be produced which is both acceptable and improves growth and survival of the fish under culture. Based on our findings, we conclude that DDG in the diet might be able to maintain fish growth while reducing feed expenditure. Therefore, the present study was conducted to examine the effects of DDG on growth performance, body composition and antioxidant activity of juvenile red seabream.

## MATERIALS AND METHODS

### Experimental diets

Essential amino acid and proximate composition of ingredients used in the experimental diets are presented in Table 1. Ingredients and chemical composition of the experimental diets are presented in Table 2. The six experimental diets were formulated to contain 0%, 5%, 10%,

15%, 20%, and 25% DDG from rice (designated as DDG0, DDG5, DDG10, DDG15, DDG20, and DDG25) with iso-nitrogenous and isocaloric diets. Pollack fish meal was used as the primary protein source. Fish oil was used as lipid source. The DDG used in this study was produced by filtration of an aqueous mixture of fermented rice with *Aspergillus oryzae* and yeasts in manufactured by Incheon Makgeolli factory (Incheon, Korea). The ingredient was dried at 60°C for 24 h and finely ground prior to incorporating in the experimental diets. All ingredients were carefully combined with 30% distilled water and pellets were prepared using laboratory moist pelleting equipment. The pellets were dried at room temperature for 48 h and finely grounded into desirable particle sizes. All diets were stored at -30°C until utilized.

### Experimental fish and feeding conditions

Juvenile red seabream were transported from a private hatchery (Namhae, Korea) to the Marine Biology Center for Research and Education at Gangneung-Wonju National University. The fish were acclimated to laboratory conditions by feeding commercial pellets for 2 weeks before starting the feeding trial. After this conditioning period, juvenile red seabream (mean body weight 10.1±0.05 g) were randomly distributed in 18 tanks (400-L rectangular plastic tanks) at a density of 30 fish per tank, respectively. Each experimental diet was fed to three groups of fish to apparent satiation two times per day (9:00 and 17:00 h for 6 days per week) for 10 wks. Filtrated seawater was supplied at a flow rate of 4 L/min in each tank and the mean water temperature and salinity were 20.4±1.98°C and 34±0.1 ppt, respectively. The photoperiod was left under natural conditions during the feeding trail. Records were kept of daily feed consumption, mortalities, and feeding behavior.

### Sample collections and analytical methods

At the end of the feeding trial, all fish in each tank were collectively weighed immediately after anesthetizing with tricaine methanesulfonate (MS222, Sigma, St. Louis, MO, USA) at a concentration of 100 ppm, and after starvation for 24 h. Proximate composition was analyzed according to standard methods (AOAC, 1995). Ten fish per tank at the end of the feeding trials were sampled and stored at -25°C for proximate composition analysis. Crude protein was determined by the Kjeldahl method using an auto Kjeldahl System (Buchi, Flawil, Switzerland). Crude lipid was analyzed with ether extraction in a soxhlet extractor (SER 148, VELP Scientifica, Milano, Italy). Moisture was determined using a dry oven at 105°C for 6 h and also the ash content was determined after combustion at 600°C for 4 h in a muffle furnace. Amino acid composition in the experimental diets and whole body of fish was determined by acid hydrolysis with 6 N HCL (reflux for 23 h at 110°C)

**Table 1.** Composition of proximate and essential amino acids of the ingredients of experimental diets

	Ingredients		
	Fish meal	Wheat flour	Distillers dried grain powder <sup>1</sup>
Proximate composition (% , dry matter basis)			
Dry matter	95.8	89.3	98.2
Crude protein	75.3	19.3	19.1
Crude lipid	8.8	3.9	7.8
Ash	14.6	2.2	0.5
Essential amino acid composition (% protein)			
Arg	6.7	5.7	6.9
His	2.3	2.9	2.0
Ile	4.5	2.3	3.6
Leu	8.3	6.0	8.0
Lys	8.8	3.7	3.1
Met+Cys	5.1	2.8	3.4
Phe+Tyr	8.1	6.8	10.8
Thr	4.8	3.5	4.7
Val	4.5	3.2	5.8

<sup>1</sup>Residue obtained by filtration of an aqueous mixture of fermented rice with *Aspergillus oryzae* and yeasts produced from Incheon Makgeolli factory (Incheon, Korea).

**Table 2.** Ingredients and chemical composition of experimental diets

Items	Diets					
	DDG0	DDG5	DDG10	DDG15	DDG20	DDG25
Ingredients (%)						
Pollack fish meal	60.0	60.0	60.0	60.0	60.0	60.0
Distillers dried grain powder <sup>1</sup>		5.0	10.0	15.0	20.0	25.0
Wheat flour	23.0	18.5	14.0	9.5	5.0	1.5
Corn gluten meal	4.0	3.5	3.0	2.5	2.0	0.5
$\alpha$ -Potato-starch	5.0	5.0	5.0	5.0	5.0	5.0
Squid liver oil	5.0	5.0	5.0	5.0	5.0	5.0
Vitamin premix <sup>2</sup>	1.0	1.0	1.0	1.0	1.0	1.0
Mineral premix <sup>3</sup>	1.0	1.0	1.0	1.0	1.0	1.0
Vitamin C (50%) <sup>4</sup>	0.5	0.5	0.5	0.5	0.5	0.5
Choline salt (50%)	0.3	0.3	0.3	0.3	0.3	0.3
Taurine	0.2	0.2	0.2	0.2	0.2	0.2
Nutrient content (dry matter basis)						
Crude protein (%)	48.9	48.4	48.3	48.0	47.3	47.2
Crude lipid (%)	9.6	10.1	10.4	10.6	10.9	10.8
Ash (%)	13.3	13.5	13.7	13.5	13.6	14.9
N-free extract <sup>5</sup>	28.2	28.0	27.6	27.9	28.2	27.1
Essential amino acid composition (% protein)						
Arg	6.5	6.5	6.4	6.7	6.6	6.7
His	2.2	2.2	2.2	2.1	2.2	2.2
Ile	3.3	3.6	3.9	2.9	3.4	3.7
Leu	8.2	8.5	8.7	8.4	8.4	8.1
Lys	7.2	7.1	6.9	6.7	6.7	6.8
Met+Cys	3.7	3.9	4.0	3.7	4.0	4.2
Phe+Tyr	7.1	7.2	7.4	7.3	7.3	7.1
Thr	4.7	4.9	4.6	4.7	4.8	4.8
Val	4.0	4.3	4.5	3.6	4.2	4.5

DDG, distillers dried grain.

<sup>1</sup> Residue obtained by filtration of an aqueous mixture of fermented rice with *Aspergillus oryzae* and yeasts produced from Incheon Makgeolli factory (Incheon, Korea).

<sup>2</sup> Vitamin premix contained the following amount which were diluted in cellulose (g/kg premix): DL- $\alpha$ -tocopheryl acetate, 18.8; thiamin hydrochloride, 2.7; riboflavin, 9.1; pyridoxine hydrochloride, 1.8; niacin, 36.4; Ca-D-pantothenate, 12.7; myo-inositol, 181.8; D-biotin, 0.27; folic acid, 0.68; p-aminobenzoic acid, 18.2; menadione, 1.8; retinyl acetate, 0.73; cholecalciferol, 0.003; cyanocobalamin, 0.003.

<sup>3</sup> Mineral premix contained the following ingredients (g/kg premix): MgSO<sub>4</sub>·7H<sub>2</sub>O, 80.0; NaH<sub>2</sub>PO<sub>4</sub>·2H<sub>2</sub>O, 370.0; KCl, 130.0; Ferric citrate, 40.0; ZnSO<sub>4</sub>·7H<sub>2</sub>O, 20.0; Ca-lactate, 356.5; CuCl, 0.2; AlCl<sub>3</sub>·6H<sub>2</sub>O, 0.15; KI, 0.15; Na<sub>2</sub>Se<sub>2</sub>O<sub>3</sub>, 0.01; MnSO<sub>4</sub>·H<sub>2</sub>O, 2.0; CoCl<sub>2</sub>·6H<sub>2</sub>O, 1.0.

<sup>4</sup> ROVIMIX STAY-C 35. DSM Nutrition Ltd., Seoul, Korea.

<sup>5</sup> Nitrogen-free extract (NFE) = 100-(crude protein+crude lipid+ash).

followed by using an automatic amino acids analyzer (Hitachi, Tokyo, Japan).

### Blood chemistry

Blood samples were taken from the caudal veins of five fish per tank using 1-mL heparinized syringes. The collected blood was centrifuged (3,500 g for 10 min) and the plasma was separated and stored in a -75°C freezer. Plasma total protein, glucose, total cholesterol, glutamic-pyruvic transaminase (GPT), phospholipid and triglyceride concentrations were determined using a clinical investigation commercial kit (Asan Pharmaceutical Co., Seoul, Korea).

### Radical scavenging activities

At the end of the feeding trials, five fish per tank were sampled and stored at -75°C for antioxidant activity analysis. Samples were extracted from plasma and liver using extract buffer in 5 mM Tris-HCl and 35 mM glycine by pH 8.4 with a homogenizer (Wiggenhauser, Berlin, Germany), followed by centrifugation (13,000 g for 10 min at 4°C). The supernatant was then collected and analyzed for its radical scavenging activity.

### Assays of 1, 1-Diphenyl-2-picryl-hydrazyl and Alkyl radicals scavenging activity on electron spin resonance spectrometer

1, 1-Diphenyl-2-picryl-hydrazyl (DPPH) radical

scavenging activity was evaluated using the method described by Nanjo et al. (1995). A 30  $\mu$ L peptide solution (or ethanol itself as control) was added to 30  $\mu$ L of DPPH (60  $\mu$ M) in ethanol solution. After mixing vigorously for 10 s, the solution was moved into a 100  $\mu$ L quartz capillary tube, and the scavenging activity of peptide on DPPH radical was determined using spectrometer (JEOL Ltd., Tokyo, Japan). After 2 min, the spin adduct was determined using an electron spin resonance (ESR) spectrometer. The measurement conditions were magnetic field, 336.5 $\pm$ 5 mT; power, 5 mW; modulation frequency, 9.41 GHz; amplitude; 1 $\times$ 1,000 and sweep time; 30 s.

Alkyl radicals were determined by 2, 2-azobiz-(2-amidinopropane)-hydrochloride (AAPH). The phosphate buffered saline (pH 7.4) reaction mixtures included 10 mM AAPH, 10 mM 4-POBN and identified concentrations of samples, which were incubated at 37°C in a water bath for 30 min (Hiramoto et al., 1993) and then moved to a capillary tube. The spin adduct was recorded on a spectrometer (JEOL Ltd., Japan). The measurement conditions were as follows: modulation frequency, 100 kHz; microwave power, 10 mW; microwave frequency, 9,441 MHz; magnetic field, 336.5 $\pm$ 5 mT and sweep time, 30 s.

The DPPH and alkyl radical scavenging activities (RSA) were computed following equation in which  $H$  and  $H_0$  were the relative peak height of radical signals with and without sample, respectively.

$$\text{RSA (\%)} = \frac{(1 - H)}{H_0} \times 100$$

### Statistical analysis

The data were subjected to one-way analysis of variance using SPSS version 19.0 (SPSS Inc., Chicago, IL, USA).

Significant differences ( $p < 0.05$ ) among the means were determined using a Duncan's multiple range test (Duncan, 1955).

## RESULTS

Growth performance, feed utilization and morphological parameters of juvenile red seabream fed the experimental diets containing different levels of DDG are presented in Table 3. No significant differences were identified in survival, weight gain, feed efficiency, daily protein intake, protein efficiency ratio and daily feed intake among the groups ( $p > 0.05$ ). Also there was no significant difference in morphological parameters, such as condition factor and hepatosomatic index. The results of body composition and essential amino acid composition of juvenile red seabream fed the experimental diets are presented in Table 4. The DDG level did not affect the proximate and amino acid composition of whole body juvenile red seabream ( $p > 0.05$ ). The results of hematological parameters of plasma in juvenile red seabream are shown in Table 5. Plasma contents of total protein, glucose, total cholesterol, GPT, phospholipid and triglyceride were not affected by dietary DDG levels ( $p > 0.05$ ). The results of RSA of plasma and liver in juvenile red seabream are presented in Table 6. DPPH and alkyl RSA in the plasma of fish did not show significant differences among treatments. Also, DPPH and alkyl RSA in the liver of fish were not affected by dietary DDG levels ( $p > 0.05$ ).

## DISCUSSION

The present results show that up to 25% DDG in the formulated diets did not have a negative effect on the

**Table 3.** Growth performances, feed utilization and morphological parameters of juvenile red seabream fed the experimental diets for 10 wks<sup>1</sup>

	Diets						SEM	p-value
	DDG0	DDG5	DDG10	DDG15	DDG20	DDG25		
IBW (g)	10.1	10.1	10.2	10.1	10.1	10.2	0.08	0.732
Survival (%)	72	89	72	82	78	78	5.09	0.135
WG (%) <sup>2</sup>	269	270	273	279	241	238	16.18	0.449
FE (%) <sup>3</sup>	91	92	96	96	87	84	4.96	0.655
DPI (%) <sup>4</sup>	0.79	0.86	0.81	0.83	0.83	0.92	0.05	0.724
PER (%) <sup>5</sup>	2.01	1.90	2.15	1.99	1.85	1.76	0.13	0.538
DFI (%) <sup>6</sup>	1.62	1.78	1.58	1.70	1.73	1.93	0.11	0.424
CF <sup>7</sup>	1.71	1.77	1.70	1.73	1.74	1.82	0.06	0.735
HSI <sup>8</sup>	2.40	2.53	2.37	2.42	2.30	2.37	0.05	0.159

DDG, distillers dried grain; SEM, standard error of mean; IBW, initial body weight; WG, weight gain; FE, feed efficiency; DPI, daily protein intake; PER, protein efficiency ratio; DFI, daily feed intake; CF, condition factor; HSI, hepatosomatic index.

<sup>1</sup> Values are means of triplicate groups. <sup>2</sup> WG = (final fish wt. - initial fish wt.) $\times$ 100/initial fish wt.

<sup>3</sup> FE = wet weight gain $\times$ 100/feed intake. <sup>4</sup> DPI = protein intake $\times$ 100/[(initial fish wt.+final fish wt.+dead fish wt.) $\times$ days reared/2].

<sup>5</sup> PER = wet weight gain/protein intake. <sup>6</sup> DFI = feed intake $\times$ 100/[(initial fish wt.+final fish wt.+dead fish wt.) $\times$ days reared/2].

<sup>7</sup> CF = fish weight (g) $\times$ 100/fish length (cm)<sup>3</sup>. <sup>8</sup> HSI = liver weight $\times$ 100/body weight.

**Table 4.** Proximate and essential amino acid composition of the whole body in juvenile red seabream fed the experimental diets for 10 wks<sup>1</sup>

Items	Diets						SEM	p-value
	DDG0	DDG5	DDG10	DDG15	DDG20	DDG25		
Proximate composition (%)								
Moisture	65.5	65.6	65.3	65.6	66.8	66.6	0.78	0.703
Crude protein	19.0	18.6	18.9	18.4	18.3	18.4	0.34	0.612
Crude lipid	8.3	9.1	8.6	8.7	8.3	7.5	0.53	0.478
Ash	5.7	5.2	5.7	5.6	5.6	5.7	0.24	0.635
Essential amino acids (% protein)								
Arg	6.9	7.0	7.1	6.9	6.9	7.0	0.11	0.874
His	2.3	2.3	2.2	2.3	2.3	2.3	0.03	0.656
Ile	3.2	3.0	2.8	3.2	3.1	2.9	0.35	0.930
Leu	7.4	7.4	7.3	7.5	7.5	7.3	0.14	0.809
Lys	8.1	8.1	7.9	8.2	8.0	8.0	0.10	0.644
Met+Cys	3.8	3.8	3.7	3.8	3.8	3.8	0.07	0.868
Phe+Tyr	6.8	6.7	6.7	6.8	6.8	6.7	0.11	0.991
Thr	4.6	4.5	4.5	4.6	4.5	4.5	0.12	0.968
Val	3.6	3.4	3.2	3.7	3.6	3.3	0.33	0.909

DDG, distillers dried grain; SEM, standard error of mean.

<sup>1</sup> Values are means of triplicate groups.

growth performance, morphological parameters, body composition and amino acid profile of juvenile red seabream. Results of this research indicate that rice-based DDG can be considered to be a candidate as a feed ingredient for juvenile red seabream. Rice-based DDG might also be a good ingredient in diets for olive flounder (Rahman et al., 2013). Several studies observed a good growth rate of tilapia when fed diets containing corn-based

DDG (Wu et al., 1996,1997; Coyle et al., 2004; Abo-state et al., 2009). This generally agrees with a number of studies showing that 30% to 40% corn-based DDG may be included in diets for channel catfish (Tidwell et al., 1990; Webster et al., 1993; Robinson and Li, 2008), rainbow trout (Cheng and Hardy, 2004; Stone et al., 2005; Barnes et al., 2012), hybrid catfish (Zhou et al., 2010) and yellow perch (Schaeffer et al., 2011) without incurring negative effects on

**Table 5.** Hematological parameters of the plasma in juvenile red seabream fed the experimental diets for 10 wks<sup>1</sup>

	Diets						SEM	p-value
	DDG0	DDG5	DDG10	DDG15	DDG20	DDG25		
Total protein (g/dL)	4.0	3.4	3.9	3.7	3.8	3.7	0.21	0.527
Glucose (mg/dL)	69	53	64	71	52	55	6.85	0.301
Total cholesterol (mg/dL)	337	269	351	333	346	367	29.24	0.498
GPT (IU/L)	0.7	0.7	1.0	1.3	0.7	0.3	0.30	0.334
Phospholipid (mg/dL)	677	694	755	750	792	864	49.81	0.393
Triglyceride (mg/dL)	162	163	182	193	161	208	18.22	0.598

DDG, distillers dried grain; SEM, standard error of mean; GPT, glutamic-pyruvic transaminase.

<sup>1</sup> Values are means of triplicate groups.**Table 6.** Radical scavenging activity of the plasma and liver in juvenile red seabream fed the experimental diets for 10 wks<sup>1</sup>

	Diets						SEM	p-value
	DDG0	DDG5	DDG10	DDG15	DDG20	DDG25		
Plasma								
DPPH radical	63.7	77.6	77.9	83.7	80.3	84.3	4.82	0.250
Alkyl radical	93.2	93.9	94.0	94.5	93.6	93.4	0.46	0.092
Liver								
DPPH radical	74.2	77.5	79.6	81.3	81.6	81.7	1.80	0.226
Alkyl radical	74.1	73.0	75.6	71.6	74.3	68.6	3.55	0.454

DDG, distillers dried grain; SEM, standard error of mean; DPPH, 1, 1-Diphenyl-2-picryl-hydrazyl.

<sup>1</sup> Values are means of triplicate groups.

growth performance. Furthermore, the inclusion of corn-based DDG in the diet may improve palatability of sunshine bass (Thompson et al., 2008). The acceptable growth performance of freshwater fishes fed diet containing DDG is definitely related to various aspects such as improving digestibility (Randall and Drew, 2010) and decreased exposure to anti-nutritional factors (Borgeson et al., 2006). Tidwell et al. (1990) reported that there were no significant differences in fish weight, survival and protein efficiency ratio of channel catfish fed diets containing up to 40% DDG for 11 weeks. In the present study, survival, weight gain, feed efficiency and protein efficiency ratio of fish fed diets containing up to 25% DDG were not different among the treatments. We consider the main reason for this result was removing anti-nutritional factors in DDG by means of fermentation. It has been reported that the nutritional profile of palm kernel meal as a constituent in a diet for tilapia *Oreochromis sp.* could possibly be enhanced by means of fermentation (Ng et al., 2002). Ramachandran et al. (2005) reported that fermentation might considerably reduce the anti-nutritional factors and crude fibre content. Slater et al. (2011) reported that fermentation associated with carbohydrate sources previous to feeding may improve digestibility as well as convenience in juvenile fish diets. During production of DDG from rice, yeasts are widely used to facilitate the fermentation method. The DDG includes a substantial quantity of yeast cells (Zohu et al., 2010) which might be abundant in proteins, B-complex vitamins and  $\beta$ -glucans. Scientific studies with sea bass (Oliva-Teles and Goncalves, 2001) and sunshine bass (Gause and Trushenski, 2011) reported growth improvement with dietary inclusion of yeast. Li et al. (2011) also ascribed the positive effect of DDG on fish growth to be due to the presence of yeast in DDG. That there was no negative effects due to the high inclusion of DDG in diets for red seabream is probably associated with beneficial effect on DDG during fermentation by *Aspergillus oryzae* and yeasts with rice.

Li et al. (2011) reported that different levels of corn-based DDG did not significantly influence the hematological parameters of Nile tilapia, *Oreochromis niloticus*. Related findings were observed the hematological parameters of juvenile red seabream in the present study. The effect of DDG was not significantly different between treatments in whole body proximate composition of juvenile red seabream. Recently studies with Nile tilapia (Li et al., 2011) and rainbow trout (Barnes et al., 2012) revealed that whole body proximate composition was not affected by dietary levels of DDG.

Many studies have shown that there was a direct relationship between antioxidant activity and total phenolic content in vegetables, fruits and grain items (Velioglu et al., 1998; Siriwardhana et al., 2003; Heo et al., 2005). DPPH

has been utilized for the determination of free radical scavenging effects on pure antioxidant chemical substances, plant and fruit extracts and food materials (Matsukawa et al., 1997; Park et al., 2005; Wong et al., 2006). However, to date, there is no available information on scavenging activities in fish fed diets containing DDG as an ingredient. Pham and Lee (2007) reported that nutritional dietary supplements of Cheongkukjang (Korean fermented soybean with rice straw) exhibited high DPPH free RSA in parrot fish. Kim et al. (2010) suggested that DPPH radical scavenging activity in juvenile olive flounder was gradually increased by the inclusion of *Meju* which is fermented soybean meal. Alkyl radicals had been measured applying ESR, generally known as a technique capable of recognizing huge reactive free radicals. Park et al. (2005) reported alkyl radical scavenging effects and noticed the positive impacts using extract from *Sargassum thunbergii*. In the present study, DPPH and alkyl RSA in plasma and liver of red seabream were not different among all groups. The results indicate that phenolic compounds in DDG in the diets did not influence in radical scavenging activity of juvenile red seabream.

Results of this experiment suggest that DDG has the potential to replace material of plant origin such as wheat flour and corn gluten meal and could be used up to 25% in diets without incurring negative effects on the growth performance of juvenile red seabream.

#### ACKNOWLEDGMENTS

This research was supported by the Fishery Commercialization Technology Development Program (110077-3) and Korea Sea Grant Program funded by Ministry of Oceans and Fisheries in Korea.

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