GROWTH PERFORMANCE AND HISTOLOGICAL APPEARANCE OF THE HUMPBACK GROUPER JUVENILE (CROMILEPTE SALTIVELIS) AFTER TREATED WITH RECOMBINANT GROWTH HORMONE

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ABSTRACT

Despite as high price consumption fish, humpback grouper grow out take very long time so its culture considered not efficient. Therefore to accelerate its growth rate and make grow out culture more efficient, recombinant *Epinephelus lanceolatus* growth hormone (rElGH) was applied by oral route. Daily application of rough rElGH at a dose of 5 mg/100 g commercial diet for 42 days resulted significance increase in growth rate compared to control groups. No specific histological damage on kidney, liver and spleen which was attributable to rElGH administration. These results strongly suggested that growth stimulation following oral administration was due to a specific action of rElGH and recombinant GH as mentioned above save for fish consumption.

Key words: growth, histology, humpback grouper, recombinant growth hormone

INTRODUCTION

Growth hormone (GH) is one of polypeptide hormone contain nearly 200 amino acids, produced and secreted by somatotroph cells of vertebrate anterior pituitary gland. This hormone has two major actions: (1) direct metabolic effects that facilitate muscle growth and glucose sparing and (2) skeletal growth effects that are partially direct and partially mediated by IGF-I (Bolander 2004). More recently several authors stated that GH participated in nearly all main physiologic processes such as ionic and osmotic regulation; protein, lipid and carbohydrate metabolisms as well as reproduction and immune (Reinecke et al., 2005; Birzniece et al., 2009; Hattori, 2009).

Within the last 3 decades, the growth promoting effects of GH have been well documented in a variety of species of fish either as endogenous or exogenous hormone (Donaldson et al., 1979; Moriyama and Kawauchi, 2001). Studies investigating the use of recombinant growth hormone (rGH) for promoting growth in salmonids (Sekine et al., 1985; Moriyama, 1995), rainbow trout (Moriyama et al., 1993), red seabream (Bin et al., 2001), flounder (Liu et al., 2008), tilapia (Li et al., 2003; Acosta et al., 2007), rabbitfish (Funkenstein et al., 2005), goldfish (Promdonkoy et al., 2004), black seabream (Tsai et al., 1997), Indonesian eel (Handoyo, 2012), giant gourami (Irmanawati, 2013) and many others species found that rGHable to accelerate the growth. Donaldson et al. (1979) suggested that exogenous GH enhances fish growth by stimulating appetite and then improving feed and protein conversion; while Silverstein et al. (1999) suggested GH might stimulate food intake.
indirectly through metabolic changes such as increased utilization of nutrients that feedback on hypothalamic centers regulating energy balance.

Regarding to the potential food safety and environmental safety issues associated with transgenic fish (Muir and Howard, 1999), the use of transgenic organism to produce rGH for increasing yields in aquaculture has generated considerable interest (Leedom et al., 2002). One method that could be used to verify the safety of exogenous substances in target organism is histopathological examination as were done by Farris et al. (2007), Liu et al. (2008) and Irmawati (2013).

While the advantage of rGH was studied elsewhere for various species, no one study was done to humpback grouper. This species is one of major marine fish consumption culturing in Indonesia, and has high price but very low growth rate. In the present study, for the first time we studied the effect of rGH on growth, survival, feed efficiency, muscle composition and histology of liver, spleen and kidney of the humpback grouper juvenile. Liver, spleen and kidney are vital organs for vertebrate, subsequently serve as major role in metabolism, natural filter of the blood and remove wastes, and immune system. The objectives of this study were to verify the effectiveness and the safety of rGH as feed additive to promote growth of the species studied.

MATERIALS AND METHODS

1. Production of protein rElGH and preparation of feed

*Escherichia coli* strain BL21 harboring pCold-ElGH protein expression vector was used as bioreactor for producing rElGH. Bacteria culture, extraction and verification of protein rElGH were done according to Alimuddin et al. (2010). To protect rElGH from proteolitic digestive enzyme, the protein was coated by hydroxypropylmethyl cellulose phthalate (HP55, Shinetsu, Japan) as described by Moriyama et al. (1993) before it mixed with commercial diet (CD).

2. Experimental design

The experiment was conducted for 42 days with five treatment groups; (1) C- or control negative (fed with CD), (2) C+ or control positive (CD with placebo), (3) pA (CD with 0.2 mg rElGH-HP55/100 g feed), (4) pB (CD with 1 mg rElGH-HP55/100 g feed) and (5) pC (CD with 5 mg rElGH-HP55/100 g feed). The serial doses was found from preliminary test. CD was bought from Japanese Company contains approximately 48.70% crude protein and 13.30% crude lipid.

3. Fish husbandry and experimental protocol

Humpback grouper juveniles (body weight 1.44±0.30 g) obtained from Main Center for Mariculture Development, Lampung, Directorate General of Aquaculture, Ministry of Marine and Fishery, were randomly distributed into 15 buckets (45 x 45 x 45 cm), 50 fish per bucket, 3 buckets per treatment group. Each bucket in the same treatment group was fitted in 2000-L fiber glass tank respectively. They were fed with CD and acclimatized to the
experimental conditions for 1 week. Fishes were reared under natural photoperiod and supplied with sand-filtrated seawater continuously. The water temperature (28–29.0 °C), salinity (31–32‰), pH (8.0–8.5) and dissolved oxygen level (5-6 mg/l) of the water were monitored routinely and maintained within the limits. Uneaten feed, feces and other dirties were siphoned daily. Fish were hand fed three times a day (at 08:30; 12:30 and 16.30) to apparent satiation. The amount of feed consumed (FC) in each bucket according to the treatment groups and the number as well as weight of fish died were recorded daily. Total body weight of fish were measured once time per 2 weeks.

4. Chemical analysis of body muscle

At the end of the 42 days trial, body muscles of 3 fish in each aquarium were dissected and pooled together for proximate analysis of muscle. Specimens of muscle were homogenized, 5 g aliquot was oven-dried to constant weight at 105 °C to estimate the moisture content, while the remaining samples were lyophilized and stored at −20 °C for chemical analysis. Kjeldahl and HCl hydrolysis methods were employed to determine the protein and lipid content, respectively. Protein, lipid and moisture content in each sample were determined in triplicate.

The content of muscle and liver glycogen were determined by multiplication of absorbent sample with mg/ml dilution then divide by absorbent standard. Absorbent sample was obtained from a serial process which was involved heating, centrifugation, hydrolysis and measuring absorbent value by spectrophotometer. Spectrophotometer was also used for measure plasma glucose.

5. Histological study

At the end of feeding trial, samples of liver, spleen and kidney were taken from 3 fish from control groups and best dose respectively, for histopathological evaluation. Liver of all fish in each group was weighed for the determination of hepatosomatic index. All samples were fixed in fixative buffered formalin, dehydrated in a graded ethanol series, embedded in paraffin and sectioned to 4 μm with a rotary microtome. Sections were stained with haematoxylin and eosin (H&E) and observed under light microscope.

6. Calculations and statistical analysis

Specific growth rate (SGR) was calculated as \((\ln W_2 - \ln W_1) \times 100 / \text{time}\), survival (SR) was \(N_t / N_0 \times 100\), feed efficiency (FE) was calculated as \((W_2 + W_D) / W_1 / \text{feed consumed} \times 100\), \(\Delta W\) was difference between \(W_2\) and \(W_1\), protein retention (PR) was calculated as \(P_{bt} - P_{b0} / P_c \times 100\) and lipid retention (LR) was calculated as \(L_{bt} - L_{b0} / L_c \times 100\). \(W_2, W_1, W_D, N_t, N_0, P_{bt}, P_{b0}, L_{bt}\) and \(L_{b0}\) are represent final body weight, initial body weight, died body weight, final number, initial number, final protein body, initial protein body, final lipid body and initial lipid body of fish, respectively; while \(P_c\) and \(L_c\) successively represent as protein consumed and lipid consumed. Hepatosomatic index (HSI) was calculated as liver weight \(\times 100 / \text{body weight}\). All data between treatment groups were analyzed by
one-way ANOVA, followed by Duncan’s Multiple Range Test of Significance. Difference was regarded as significant when P<0.05.

RESULTS AND DISCUSSION

Fish growth performance was presented in Table 1. Although during the 42 days trial there were some fish died in C+, pB and pC but the survival rate (SR) differences among experimental groupswere not significant. Final body weight (W2), SGR, FC and FE of the rGH treatment groups were significantly different than two control groups; however the difference between two control groups were not significant. These results indicated that the growth-promoting effect was caused by rElGH, not by the placebo itself. ∆Wof pA, pB and pC subsequently were 23.69 %, 31.45 % and 40.25 % higher than ∆Wof C-. The figure of pC was highest than other experimental groups, it is indicated that the dose for pC was better than pB and pA.

Table 1. Effect of rElGH on survival, growth, feed consumed and feed efficiency of humpback grouper juvenile.

<table>
<thead>
<tr>
<th>Group</th>
<th>SR (%)</th>
<th>W2 (g)</th>
<th>∆W (g) % ∆W to Control</th>
<th>SGR</th>
<th>FC (g)</th>
<th>FE (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>C</td>
<td>100 a</td>
<td>5.92±1.1</td>
<td>4.77±0.2</td>
<td>3.92±0.0</td>
<td>217.12±3</td>
<td>109.81±5</td>
</tr>
<tr>
<td>C+</td>
<td>99.33±1</td>
<td>6.23±1.0</td>
<td>5.08±0.2</td>
<td>6.5</td>
<td>4.04±0.0</td>
<td>226.04±7</td>
</tr>
<tr>
<td>pA</td>
<td>100 a</td>
<td>7.07±0.8</td>
<td>5.90±0.5</td>
<td>23.69</td>
<td>4.33±0.0</td>
<td>245.51±1</td>
</tr>
<tr>
<td>pB</td>
<td>97.33±2</td>
<td>7.49±1.0</td>
<td>6.27±0.4</td>
<td>31.45</td>
<td>4.46±0.0</td>
<td>246.65±1</td>
</tr>
<tr>
<td>pC</td>
<td>99 ±</td>
<td>7.83±1.5</td>
<td>6.69±0.3</td>
<td>40.25</td>
<td>4.59±0.0</td>
<td>269.75±1</td>
</tr>
</tbody>
</table>

Different superscript letter in the same column indicated significant difference based on Duncan test (P<0.05).

It is the first study revealed that feeding rElGH could significantly stimulate the growth of humpback grouper. Other study related with rElGH was done by Handoyo (2012) to Indonesian eel fingerling by immersion, oral route and combination of immersion and oral; increasing growth subsequently as big as 37.4 %, 65.7 % and 102.9 % higher than control. Same application methods were used by Subaidah (2013) for whiteshrimp, the growth of rElGH treatment group were 109.9 %, 17.7 % and 40.1 % respectively higher than control. Other studies also demonstrated the efficiency of oral administrated rGHs in promoting fish growth. Moriyama et al. (1993) reported that feeding recombinant salmon GH caused significant increase in body weight and length of rainbow trout. Oral delivery of recombinant fish GH induced a 1.6-fold increase in body weight of black seabream (Tsai et al., 1997). Promdonkoy et al. (2004) used rGH from giant catfish to goldfish by oral delivery found increasing body weight 84 % higher than control. Despite the proteolytic enzymes in the gastrointestinal tract of fish could hydrolyze oral administrated GH, the present result along
with the cited reports above clearly demonstrated that dietary delivery of rGH in form of rGH coated HP55 is an effective approach to enhance fish growth.

As mentioned earlier, exogenous GH may enhance fish growth by stimulating appetite then increasing feed consume and improving feed conversion (Donaldson et al., 1979; Silverstein et al., 1999). This study showed that after consuming rGH, growth rate, and appetite/feed consume and feed efficiency of humpback grouper significantly higher than control groups (Table 1). It is consistent with study result of Markert et al. (1977) in salmonids, Wilson et al. (1988) and Silverstein et al. (2000) in channel catfish. Reversely Liu et al. (1999; 2008) found different result, feeding recombinant GH to flounder could promote fish growth and improve feed conversion, but did not affect feed intake. GH might stimulate food intake indirectly through metabolic changes such as increased utilization of nutrients that feedback on hypothalamic centers regulating energy balance (Silverstein et al., 1999). One of hypothalamic factors which may involve to regulate energy balance is ghrelin, where GH and ghrelin working together in a 2 ways mechanism (Wong et al., 2006). Further investigations are needed to verify relation between GH and ghrelin in promoting growth and feed consumption.

Table 2. Proximate composition of body muscle, liver and muscle glycogen, and plasma glucose of humpback grouper juvenile after 42 days trial.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Protein</th>
<th>Lipid</th>
<th>Carbohydrate</th>
<th>Liver Glycogen</th>
<th>Muscle Glycogen</th>
<th>Plasma Glucose</th>
</tr>
</thead>
<tbody>
<tr>
<td>C -</td>
<td>16.89±0.34&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.85±0.16&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.69±0.29&lt;sup&gt;a&lt;/sup&gt;</td>
<td>4.11±0.51&lt;sup&gt;a&lt;/sup&gt;</td>
<td>4.99±0.58&lt;sup&gt;a&lt;/sup&gt;</td>
<td>55.78±13.10&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>C +</td>
<td>16.38±0.64&lt;sup&gt;a&lt;/sup&gt;</td>
<td>6.16±0.23&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.49±0.37&lt;sup&gt;a&lt;/sup&gt;</td>
<td>4.44±0.45&lt;sup&gt;a&lt;/sup&gt;</td>
<td>3.32±1.46&lt;sup&gt;a&lt;/sup&gt;</td>
<td>41.53±5.86&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>pA</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>58.86±12.79&lt;sup&gt;a,b&lt;/sup&gt;</td>
</tr>
<tr>
<td>pB</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>-</td>
<td>59.65±15.48&lt;sup&gt;a,b&lt;/sup&gt;</td>
</tr>
<tr>
<td>pC</td>
<td>19.32±0.53&lt;sup&gt;b&lt;/sup&gt;</td>
<td>6.22±0.10&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.81±0.10&lt;sup&gt;a&lt;/sup&gt;</td>
<td>5.56±0.31&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.77±0.56&lt;sup&gt;a&lt;/sup&gt;</td>
<td>81.95±12.85&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Different superscript letter in the same column indicated significant difference based on Duncan test (P<0.05).

Table 3. Hepatosomatic index (HSI), protein retention (PR) and lipid retention (LR) of humpback grouper juvenile after 42 days trial.

<table>
<thead>
<tr>
<th>Group</th>
<th>HSI</th>
<th>PR</th>
<th>LR</th>
</tr>
</thead>
<tbody>
<tr>
<td>C -</td>
<td>2.99±0.76&lt;sup&gt;a&lt;/sup&gt;</td>
<td>3.23±0.69&lt;sup&gt;a&lt;/sup&gt;</td>
<td>-0.07±0.79&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>C +</td>
<td>3.71±0.31&lt;sup&gt;a&lt;/sup&gt;</td>
<td>2.18±1.30&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.00±0.88&lt;sup&gt;ab&lt;/sup&gt;</td>
</tr>
<tr>
<td>pA</td>
<td>4.46±1.29&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>pB</td>
<td>4.89±1.21&lt;sup&gt;b&lt;/sup&gt;</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>pC</td>
<td>5.45±0.80&lt;sup&gt;b&lt;/sup&gt;</td>
<td>8.21±1.08&lt;sup&gt;b&lt;/sup&gt;</td>
<td>2.03±0.68&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Different superscript letter in the same column indicated significant difference based on Duncan test (P<0.05).

Protein content, liver glycogen and plasma glucose from rGH treatment (represent by pC) were significantly higher than both control groups. Even though lipid and carbohydrate content have tendency higher than control but statistically not different as well as muscle glycogen (Table 2). These findings were consistent that GH has direct metabolic effects including the stimulation of lipolysis and protein synthesis which were especially pronounced in muscle. GH also has diabetogenic effect, that is induces peripheral resistance to insulin such that glucose cannot be used and blood glucose levels rise then increasing liver glycogen deposit (Bolander, 2004). Hepatosomatic index and protein retention were significantly higher
than control whereas lipid retention significantly different than C – but if compare with C + not different (Table 3). A study in coho salmon transgenic GH showed that GH able to increase HSI value 1.3 higher than control (Leggat et al., 2009). It has been known that the receptor of GH in fish mainly distributed in liver cells and liver is a main organ for GH to stimulate IGF-1 production (Moriyama and Kawauchi, 2001); therefore as a consequence of increasing IGF-1 production and other anabolic and katabolic metabolism GH treated organism need bigger size and capacity of liver.

Many studies have investigated the effect of rGH on fish body composition specifically on protein and lipid content. Silverstein et al. (2000) found that recombinant bGH injection induced protein and lipid content in USDA-103 and Norris strain of channel catfish. Whereas, Peterson et al. (2004) observed no effect of recombinant bGH injection on body composition in NWAC103 and Norris strain of channel catfish. According to Liu et al. (1999), crude protein and crude fat content of muscle in flounder were increased by feeding recombinant yeast containing salmon GH. However, Liu et al. (2008) reported that protein and lipid content in muscle of flounder were not changed by feeding GH transgenic SYN. Different result was reported on giant gourami after treated with rGH, where lipid of treated fish was significantly lower than control groups. Investigation related to the lipid content showed that lipase enzyme activity in rGH treatment fish was higher than control (Irmawati, 2012). Protein and lipid content after feeding rGH are interesting topics for discussion; generally protein content is a desirable whereas lipid content is not desirable for commercially consumption fish cultured. Although fish size, age, diet and some experimental factors can affect to the body composition, the main cause for the different results obtained by different authors may more likely be the different responses to rGH treatment from different fish species and hormone dose used in experiments (Liu et al., 2008). Silverstein and Plisetskaya (2000) stated that the potential increasing body fatness after growth hormone treatment should be viewed with caution because increased body fat in juvenile fish may reduce feed intake for long time.

Histological examination showed that no apparent differences in the appearance of liver, kidney and spleen tissue of C - and pC (Figure 1 and 2). There were some histological abnormalities found in all organs studied from both experimental groups such as inclusion body acidophilic, necrosis, hemorrhage, fatty degeneration, hydropic degeneration, hyaline degeneration and melano macrophage center. Based on routine check in our laboratory, the appearance of inclusion body acidophilic indicating that experimental fishes from all treatment infected by iridovirus. Meanwhile, another cell abnormalities caused by several infectious and non-infectious diseases had been reported (Medlineplus Encyclopedia, 2013; Kurtovic et al., 2008; Agius and Roberts, 2003). Even though not clear what is the main cause of cell abnormalities in humpback grouper, since no specific histological damage which was attributable to rGH treatment group then it could be say that rGH consumption is safe for fish. This finding was consistent with several studies conducted before, such as by Liu et al. (2008) on flounder and Irmawati (2013) on giant gourami which were found that rGH consumption
didn’t cause any specific histological damage for fish internal organs. In human clinical experience, rGH consumption replacement therapy for long time indicating a lack of convincing evidence for an increased risk of cell abnormalities (Farris et al., 2007). However, quantitative histological study still needed to elucidate the safety of rGH consumption in fish for longer duration.

Figure 1. Histological appearance of liver (a), kidney (b), and spleen (c) of control

CONCLUSION

The study result clearly demonstrated that rGH from *Epinephelus lanceolatus* gene could enhanced the growth of humpback grouper and rElGH administration by oral delivery to be an effective method and looked to be an economically viable feed additive for fish culture at commercial scale. This study also revealed that rGH consumption has no negative effects on survival, muscle composition and histology of fish studied. However, as our conclusions were based on a 6-week feeding trial, further investigations using other fish species, long-term feeding experiment and quantitative histology examination should be undertaken to further confirm the growth-promoting effect and safety as feed additive of rGH.

Acknowledgements

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REFERENCES


