

Associations between Feed Efficiency, Body Growth and Serum Insulin-like Growth Factor-I Level for Korean Native Ogol Chickens

W. K. Kim, M. H. Kim, D. S. Seo, C. Y. Lee¹, Y. O. Suk² and Y. Ko*

Dept. of Animal Science, College of Life and Environmental Sciences, Korea University, Seoul 136-701, Korea

ABSTRACT : Increasing of body weight has been one of the important economic factors in the poultry industry. Insulin-like growth factor (IGF)-I is a polypeptide that serves to regulate muscle development and body growth. Moreover, IGF-I is related to feed efficiency. However, there are few studies regarding the regulatory roles of chicken IGF-I/-II compared with that of mammals. Especially, the Korean Native Ogol Chicken (KNOC) has a lean body growth and its body weight is generally lighter than the broiler chicken. Therefore, this study was conducted to investigate associations among serum IGF-I/-II concentration, feed efficiency, and body growth in KNOC. The body weight and feed intake of KNOC were recorded from 20 to 36 weeks at 2 weeks intervals, and blood was taken every 2 weeks. Serum IGF-I/-II were measured by RIA. Chickens were divided into two groups, high and low serum IGF-I concentration. Generally, feed efficiency and growth performance (body weight and weight gain) in the high serum IGF-I group were higher than those of the low group during the experimental period. In particular, the body weight of the IGF-I high group were significantly different from those of the IGF-I low group at 34 and 36 weeks, respectively ($p < 0.05$). Moreover, body weight, weight gain, and feed efficiency had a significant correlation with serum IGF-I at several weeks ($p < 0.05$ and $p < 0.01$). These results show that IGF-I plays an important role in body growth and suggests a possibility that serum IGF-I could be used as a selection marker for body growth in KNOC. (*Asian-Aust. J. Anim. Sci.* 2005. Vol 18, No. 4 : 532-537)

Key Words : IGF-I, IGF-II, Body Growth, Feed Efficiency, Korean Native Ogol Chicken

INTRODUCTION

Korean Native Ogol Chicken (KNOC) is a protected species by the Korean government (Protected Species Act No. 265), which is a dual purpose (egg and meat) chicken with lean body growth. The body weight of KNOC is generally lighter than that of broiler chicken.

Body growth has been one of the most important economic traits in the domestic animal industry and is regulated by many factors including endocrine factors. Consequently, many endocrinological studies have been conducted to improve body growth in many species (Oudin et al., 1998; Auchtung et al., 2001; Seo et al., 2001; Yoon et al., 2001; Lee et al., 2002; Yun et al., 2003a). In particular, growth factors, such as insulin-like growth factor (IGF)-I have been shown to stimulate body growth in mammals (Bass et al., 1999; Ohlsson et al., 2000). However, there are few studies regarding the regulatory roles of chicken IGFs in body growth.

Chicken IGF-I is a 7 kDa nonglycoprotein composed of 70 amino acids. Although there is a structural similarity between mammalian and avian IGF-I (Ballard et al., 1990), some unique differences have been reported. For example,

free form of IGF-I is more present in chickens and biological responses to IGFs are different in several metabolic pathways (McMurtry et al., 1997; Duclos et al., 1999). Moreover, several reports suggested that IGF-I expression in the chicken muscle could be largely independent of growth hormone, contrary to mammals (Rosselot et al., 1995; Tanaka et al., 1996). Generally, mammalian IGF-I mediates growth hormone action (Rotwein, 1991) and regulates cellular proliferation and differentiation by endocrine as well as auto/paracrine manners in many species (Jones and Clemmons, 1995; Simmen et al., 1998; Liu and LeRoith, 1999).

Similarly, avian IGFs play important roles in the uptake of amino acid and glucose, muscle protein synthesis, feed efficiency, and posthatch development (Kocamis et al., 1998; Buyse and Decuyper, 1999; Colon and Kita, 2002). There were reports regarding a negative correlation between chicken IGF-I and body growth (Pym et al., 1991; Bacon et al., 1993). In addition, Huybrechts et al. (1992) reported that IGF-I did not appear to stimulate growth rate. However, recent studies have shown that circulating IGF-I concentrations have a positive correlation with muscle growth rate and decreased fatness (Guernec et al., 2003; Tesseraud et al., 2003; Yun et al., 2003b)

Collectively, due to such contradictory findings, studies are required to clarify regulatory roles of IGF-I in avian growth physiology. Therefore, this study was conducted to elucidate the association among serum IGF-I/-II concentration, body growth, and feed efficiency, and to investigate the possibility of improving both body weight and feed efficiency by serum IGF-I concentration in KNOC.

* Corresponding Author: Y. Ko. Tel: +82-2-3290-3054, Fax: +82-2-925-1970, E-mail: yongko@korea.ac.kr

¹ Regional Animal Industry Research Center, Jinju 660-758, Korea.

² Department of Applied Animal Science, Sahmyook University, Seoul 139-742, Korea.

Received May 19, 2004; Accepted November 26, 2004

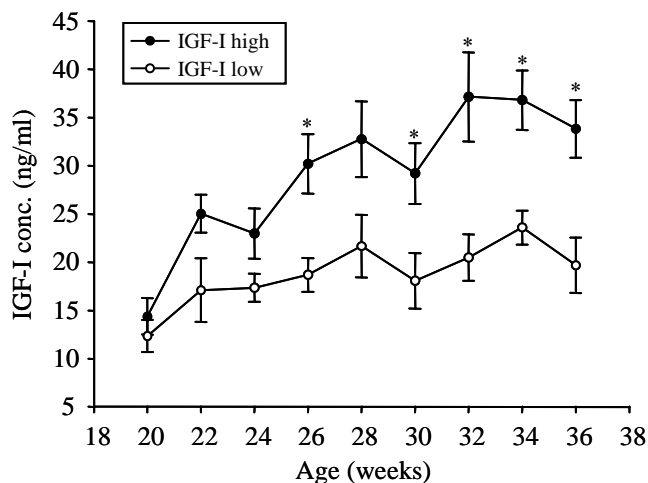


Figure 1. Comparison of serum IGF-I concentrations between high and low IGF-I expression groups. Values are the mean \pm SE. Means with different superscripts differ significantly ($p < 0.05$).

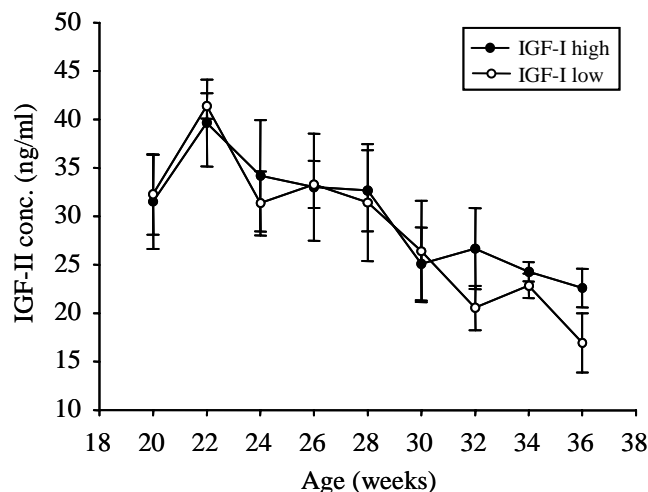


Figure 2. Comparison of serum IGF-II concentrations between high and low IGF-I expression groups. Values are the mean \pm SE.

MATERIALS AND METHODS

Animals

KNOCs were purchased from Korean Native Ogol Chicken Breeding Farms (Yeonsan, Korea). After acclimation, they were raised at the Sahmyook University Animal Breeding Center. A total of 50 female KNOC were kept in individual cages (40 cm high, 25 cm wide and 35 cm deep), and fed a commercial boiler diet (Kang et al., 2003). The body weight and feed intake of KNOC were recorded biweekly and blood was taken from the wing vein from 20 to 36 wks every 2 wks. Serum was made according to the standardized procedure.

Selection for experimental groups

KNOCs were divided into high and low groups based on serum IGF-I concentrations at the age of 36 weeks. Selection between high (>28.3 ng/ml, $n=10$) and low (<17.9 ng/ml, $n=10$) groups was made as 20% in both the upper and lower classes, respectively.

IGF Radioimmunoassay (RIA)

Recombinant human IGF-I/-II (GroPep, Pty Ltd., North Adelaide, Australia) were iodinated by the chloramine-T method (Lee and Henricks, 1990). The concentrations of serum chicken IGF-I/-II were measured by heterologous RIA (Radecki et al., 1997) with minor modifications, using anti-human IGF-I and anti-mouse IGF-II antiserum (GroPep, Pty Ltd.). The intra and inter assay variations of IGF-I were 7.6 and 13.2%, and IGF-II assay had an intra assay variation of 8.3% and inter assay variation of 14.8%.

Feed efficiency

Feed intake and feed efficiency were measured by standardized methods from 20 to 36 wks every 2 wks.

Statistical analysis

Data were statistically analyzed using the Duncan method of one-way ANOVA and the Pearson's correlation coefficients procedure in SAS (SAS Inst. Inc., Cary, North Carolina, USA)

RESULTS

Serum IGF-I/-II concentrations

The changes in serum IGF-I/-II concentrations of the selected group during 20-36 wk are shown in Figure 1 and 2. In general, serum IGF-I concentrations progressively increased with age in both high and low IGF-I groups, and maximum serum IGF-I concentrations were observed at 32 wks and 34 wks, respectively. The concentrations of serum IGF-I in the high group were higher than those in the low group during the experimental period. Especially, serum IGF-I concentrations were significantly different between high and low groups at 26, 30, 32, 34 and 36 wks (Figure 1).

Compared to the IGF-I profile, a much different profile of serum IGF-II expression was obtained in KNOC (Figure 2). Generally, IGF-II concentrations progressively decreased with age of KNOC, and maximal serum IGF-II concentrations were detected at 22 wk in both groups. However, any significant differences between two groups were not observed.

Comparison of growth performance

A possible effect of serum IGF-I concentration on body weight was investigated by comparing body weights and weight gain between the high and low IGF-I expression groups (Figures 3 and 4).

Body weight increased with age both the high and low groups, showing a higher body weight in the IGF-I high group than that in IGF-I low group. In particular, significant differences between two groups was found at 34 and 36 wks,

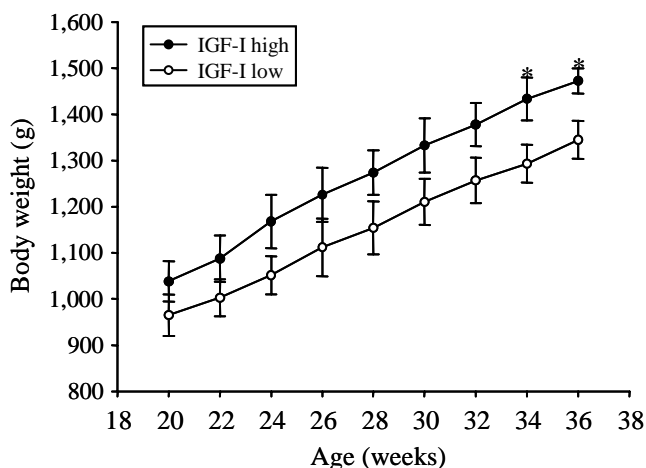


Figure 3. Comparison of body weight between high and low IGF-I expression groups. Values are the mean \pm SE. Means with different superscripts differ significantly ($p < 0.05$).

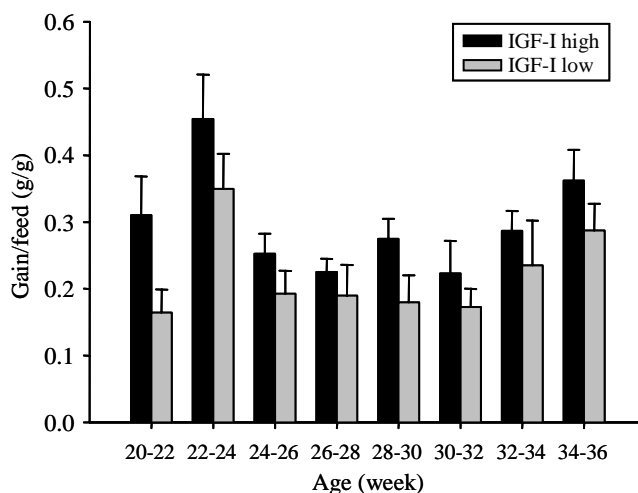


Figure 5. Comparison of feed efficiency between high and low IGF-I expression groups. Values are the mean \pm SE.

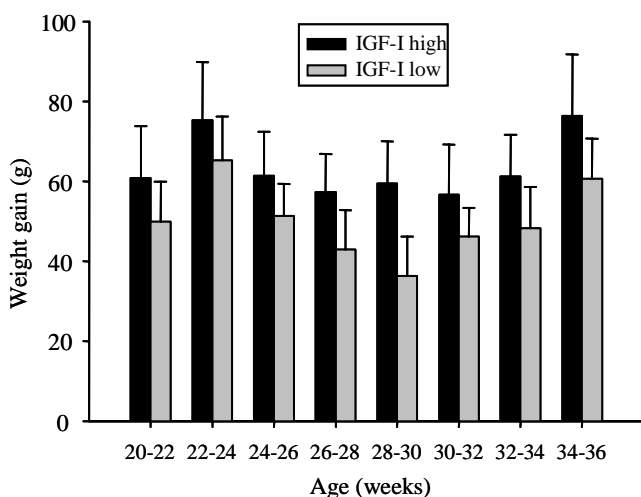


Figure 4. Comparison of weight gain between high and low IGF-I expression groups. Values are the mean \pm SE.

respectively ($p < 0.05$, Figure 3).

A comparison study on weight gain in both groups revealed no significant difference, but the serum IGF-I high group tended to show higher weight gain than the low group (Figure 4).

Comparison of feed efficiency

In order to investigate whether a link between IGF-I concentration and feed efficiency exists, two groups were compared with respect to serum IGF-I and feed efficiency.

Similarly to a weight gain pattern, a significant difference was not found and the IGF-I high group tended to show higher feed efficiency than the IGF-I low group (Figure 5).

Correlation coefficient of serum IGF-I/II with feed efficiency, weight gain, and body weight

Table 1 shows the correlation coefficients of serum IGF-

I with feed efficiency, weight gain, and body weight. The both body weight and feed efficiency generally showed positive correlation with serum IGF-I, particularly at 24 and 34 wks ($p < 0.05$). Moreover, a highly significant correlation between serum IGF-I and feed efficiency, including weight gain, was observed at 30 wks ($p < 0.01$). In addition, weight gain and body weight also showed positive correlation with serum IGF-I at 22, 24 and 32 wks, respectively ($p < 0.05$).

On the contrary, serum IGF-II did not show any significant correlation with body growth parameters except at 22 wk (Table 2).

DISCUSSION

Body growth of domestic animals is regulated by the coordinated actions of nutritional, genetic, environmental and endocrine factors (Etherton and Kensinger, 1984). Among the several endocrine factors, IGF-I in particular is generally known to be an important regulator of muscle development in a number of species (Bass et al., 1999; Yun et al., 2003a). For unclear reason in chickens, IGF-I is not always accompanied by enhanced body growth. For instance, administration of IGF-I has been reported to have no apparent effect on growth rate (McGuinness and Cogburn, 1991; Huybrichts et al., 1992; Czerwinski et al., 1998). Similarly, Mitchell and Burke (1995) reported no clear relationships between growth and plasma IGF-I concentrations. However, different results have also been claimed, reporting that IGF-I treatment consistently has been associated with an increased growth rate (Kocamis et al., 1998; Tomas et al., 1998; Colon and Kita, 2002). These indicate that the relationships between IGF-I and body growth is not apparent in poultry. Therefore, the present study analyzed the concentrations of IGF-I/II, feed efficiency, and body growth to clarify the roles of IGF-I in

Table 1. Correlation coefficients of serum IGF-I concentration with feed efficiency and body weight in KNOC

	IGF-I concentration								
	20 w	22 w	24 w	26 w	28 w	30 w	32 w	34 w	36 w
Feed efficiency	-	0.393	0.601**	-0.049	0.507*	0.624**	0.083	0.561*	0.383
Weight gain	-	0.428*	0.142	-0.034	0.473*	0.535**	0.099	0.227	0.306
Body weight	0.267	0.295	0.418*	0.136	0.242	0.101	0.463*	0.441*	0.256

* p<0.05, **<0.01.

Table 2. Correlation coefficients of serum IGF-II concentration with feed efficiency and body weight in KNOC

	IGF-II concentration								
	20 w	22 w	24 w	26 w	28 w	30 w	32 w	34 w	36 w
Feed efficiency	-	-0.550	0.398	0.258	0.181	.371	0.286	0.159	0.062
Weight gain	-	-0.174	0.360	0.318	0.082	.278	-0.038	0.044	-0.043
Body weight	-0.099	0.548*	0.293	0.427	0.204	.067	0.375	0.056	0.149

* p<0.05.

body growth of KNOC as a model and to furthermore investigate the possibility that serum IGF-I concentration could be used as a selection marker for body growth in chicken.

The concentrations of serum IGF-I in the IGF-I high group were higher than those in the low group, especially at 26, 30, 32, 34 and 36 wks (p<0.05, Figure 1). Similar profiles on feed efficiency (Figure 4), body weight, and weight gain (Table 1) were also obtained.

Although IGF-I treatment did not affect chicken growth rate and feed consumption (McGuinness and Cogburn, 1991; Czerwinski et al., 1998), the results in the present study are consistent with other reports that IGF-I stimulates posthatching muscle development in chickens (Colon and Kita, 2002; Guernec et al., 2003) and that IGF-I treatment promotes lean growth and feed utilization efficiency in broiler chickens (Kocamis et al., 1998; Tomas et al., 1998). Moreover, serum concentration of IGF-I in chickens selected for high growth rate was higher than that in chickens selected for low growth rate (Beccavin et al., 2001). These reports indicate that serum IGF-I controls the body growth, suggesting that they may stimulate chicken feed efficiency and body weight. In pigs, a positive correlation between serum IGF-I concentrations and body growth has been demonstrated (Yun et al., 2003a).

Contradictory arguments on the roles of IGF-I in weight gain were also published. Infusion of IGF-I and growth hormone did not produce differences in weight gain and feed intake (Huybrechts et al., 1992; Vasilatos-Younken et al., 1999) but, others (Kita et al., 2002; Tesseraud et al., 2003) along with the present study (Figure 4, Table 1) pointed toward a positive relationship between IGF-I expression and weight gain.

Spencer et al. (1996) and Tomas et al. (1998) have observed that administration of IGF-II did not affect feed efficiency and body growth in chickens, implying no involvement of IGF-II in feed efficiency and body growth. The present study also shows that serum IGF-II is not correlated with feed efficiency, weight gain, or body weight

(Table 2). Moreover, serum IGF-II concentration between the IGF-I high and low groups were not significantly different (Figure 2). However, serum IGF-II concentration at 22 wk was positively correlated to body weight (0.548, p<0.05, Table 2). Similarly, maximal concentration of serum IGF-II was also detected at 22 wk. Like these results, Beccavin et al. (2001) have reported that IGF-II mRNA level showed a similar trend with nutritional state. Moreover, Decuypere et al. (1993) suggested that the plasma IGF-II level was related with body weight in chickens, indicating that the serum IGF-II concentration may not directly affect feed efficiency but rather indirectly affect body weight in KNOC.

In conclusion, the body weight, weight gain, and feed efficiency in the serum IGF-I high group were higher than those in the serum IGF-I low group during the experimental period. Moreover, significantly positive correlations of serum IGF-I concentration with body weight, weight gain, and feed efficiency were observed, suggesting that IGF-I directly promotes not only feed efficiency but also weight gain, resulting in increased body growth of KNOC. Collectively, this study shows that serum IGF-I plays an important role in the improvement of body growth and suggests a possibility that serum IGF-I could be used as a selection marker for body growth in KNOC.

ACKNOWLEDGMENTS

This work was supported by grant No. R01-2002-000-00553-0 from the Basic Research Program of the Korea Science & Engineering Foundation (KOSEF) through the Korea University, and partially conducted by the Regional Animal Industry Research Center at Jinju National University.

REFERENCES

- Auchtung, T. L., E. E. Connor, S. M. Barao, L. W. Douglass and G. E. Dahl. 2001. Use of growth hormone response to growth

- hormone-releasing hormone to determine growth potential in beef heifers. *J. Anim. Sci.* 79:1566-1572.
- Bacon, W. L., K. E. Nestor, D. A. Emmerson, R. Vasilator-Younken and D. W. Long. 1993. Circulating IGF-I in plasma of growing male and female turkeys of medium and heavy weight lines. *Domest. Anim. Endocrinol.* 10:267-277.
- Ballard, F. J., R. J. Johnson, P. C. Owens, G. L. Francis, F. M. Upton, J. P. McMurtry and J. C. Wallace. 1990. Chicken insulin-like growth factor-I: amino acid sequence, radioimmunoassay, and plasma levels between strains and during growth. *Gen. Comp. Endocrinol.* 79:459-468.
- Bass, J., M. Oldham, R. Sharma and R. Kambaddur. 1999. Growth factors controlling muscle development. *Domest. Anim. Endocrinol.* 17:191-197.
- Beccavin, C., B. Chevalier, L. A. Cogburn, J. Simmon and M. J. Duclos. 2001. Insulin-like growth factors and body growth in chickens divergently selected for high and low growth rate. *J. Endocrinol.* 168:297-306.
- Buyse, J. and E. Decuyper. 1999. The role of the somatotrophic axis in the metabolism of the chicken. *Domest. Anim. Endocrinol.* 17:245-55.
- Colon, M. A. and K. Kita. 2002. Muscle protein synthesis rate is altered in response to a single injection of insulin-like growth factor-I in seven day-old Leghorn chicks. *Poult. Sci.* 81:1543-1547.
- Czerwinski, S. M., J. M. Cate, G. Francis, F. Tomas, D. M. Brocht and J. P. McMurtry. 1998. The effect of insulin-like growth factor-I (IGF-I) on protein turnover in the meat-type chicken (*Gallus domesticus*). *Comp. Biochem. Physiol. C. Pharmacol. Toxicol. Endocrinol.* 119:75-80.
- Decuyper, E., F. R. Leenstra, J. Buyse, L. M. Huybrechts, F. C. Buonomo and L. R. Berghman. Plasma levels of growth hormone and insulin-like growth factor-I and -II from 2 to 6 weeks of age in meat-type chickens selected for 6-week body weight or for feed conversion and reared under high or normal environmental temperature conditions. *Reprod. Nutr. Dev.* 33:361-372.
- Duclos, M. J., C. Beccavin and J. Simon. 1999. Genetic models for the study of insulin-like growth factor (IGF) and muscle development in birds compared to mammals. *Domest. Anim. Endocrinol.* 17:231-243.
- Etherton, T. D. and R. Kensinger. 1984. Endocrine regulation of fetal and postnatal meat animal growth. *J. Anim. Sci.* 59:511-528.
- Gonzales, E., J. Buyse, J. R. Sartori, M. M. Loddi and E. Decuyper. 1999. Metabolic disturbance in mail broilers of different strains. 2. Relationship between the thyroid and somatotrophic axes with growth rate and mortality. *Poult. Sci.* 78:516-521.
- Guernec, A., C. Berri, B. Chevalier, N. Wacrenier-Cere, E. Le Bihan-Duval and M. J. Duclos. 2003. Muscle development, insulin-like growth factor-I and myostatin mRNA levels in chickens selected for increased breast muscle yield. *Growth Horm. IGF. Res.* 13:8-18.
- Gultom, D., A. Songsang and U. Ter Meulen. 2001. The effect of chlorocholine chloride (CCC) inclusion in the diets of growing hens on growth rate, oestrogen levels and the onset of lay. *J. Anim. Physiol. Anim. Nutr. (Berl.)* 85:1-8.
- Humbel, R. E. 1990. Insulin-like growth factors I and II. *Eur. J. Biochem.* 190:445-462.
- Huybrechts, L. M., E. Decuyper, J. Buyse, E. R. Kuhn and M. Tixier-Biohard. 1992. Effect of recombinant human insulin-like growth factor-I on weight gain, fat content, and hormonal parameters in broiler chickens. *Poult. Sci.* 71:181-187.
- Jones, J. I. and D. R. Clemmons. 1995. Insulin-like growth factors and their binding proteins: biological actions. *Endocr. Rev.* 16:2-34.
- Kang, W. J., D. S. Seo and Y. Ko. 2003. Association among egg productivity, granulosa layer IGF-I and ovarian IGF-I in Korean Native Ogol Chicken. *Asian-Aust. J. Anim. Sci.* 16:325-330.
- Kcoamis, H., D. C. Kirkpatrick-Keller, H. Klandorf and J. Killefer. 1998. *In ovo* administration of recombinant human insulin-like growth factor-I alters postnatal growth and development of the broiler chicken. *Poult. Sci.* 77:1913-1919.
- Kita, K., S. Kato, M. Amanyaman, J. Okumura and H. Yokota. 2002. Dietary L-carnitine increases plasma insulin-like growth factor-I concentration in chicks fed a diet with adequate dietary protein level. *Br. Poult. Sci.* 43:117-121.
- Kocamis, H., D. C. McFarland and J. Killefer. 2001. Temporal expression of growth factor genes during myogenesis of satellite cells derived from the biceps femoris and pectoralis major muscles of the chicken. *J. Cell. Physiol.* 186:146-152.
- Lee, C. Y. and D. M. Henricks. 1990. Comparisons of various acidic treatments of bovine serum on insulin-like growth factor-I immunoreactivity and binding activity. *J. Endocrinol.* 127:139-148.
- Lee, C. Y., H. P. Lee, J. H. Jeong, K. H. Baik, S. K. Jin, J. H. Lee and S. H. Sohnt. 2002. Effects of restricted feeding, low-energy diet, and implantation of trenbolone acetate plus estradiol on growth, carcass traits, and circulating concentrations of insulin-like growth factor (IGF)-I and IGF-binding protein-3 in finishing barrows. *J. Anim. Sci.* 80:84-93.
- Liu, J. L. and D. LeRoith. 1999. Insulin-like growth factor I is essential for postnatal growth in response to growth hormone. *Endocrinology* 140:5178-5184.
- McGuinness, M. C. and L. A. Cogburn. 1990. Measurement of developmental changes in plasma insulin-like growth factor-I levels of broiler chicken by radioreceptor assay and radioimmunoassay. *Gen. Comp. Endocrinol.* 76:446-458.
- McGuinness, M. C. and L. A. Cogburn. 1991. Response of young broiler chicken to chronic injection of recombinant-derived human insulin-like growth factor-I. *Domest. Anim. Endocrinol.* 8:611-620.
- McMurtry, J. P., G. L. Francis and Z. Upton. 1997. Insulin-like growth factors in poultry. *Domest. Anim. Endocrinol.* 14:199-229.
- Mitchell, R. D. and W. H. Burke. 1995. Posthatching growth and pectoralis muscle development in broiler strain chickens, bantam chickens and the reciprocal crosses between them. *Growth Dev. Aging.* 59:149-161.
- Oudin, A., B. Chevalier, J. Simon and M. J. Duclos. 1998. Muscle insulin-like growth factor-I (IGF-I) receptors in chickens with high or low body weight: effects of age and muscle fibre type. *Growth Horm. IGF. Res.* 8:243-250.
- Pym, R. A., R. J. Johnson, D. B. Etse and P. Eason. 1991.

- Inheritance of plasma insulin-like growth factor-I and growth rate, food intake, food efficiency and abdominal fatness in chickens. *Br. Poult. Sci.* 32:285-293.
- Radecki, S. V., M. C. Capdevielle, F. C. Buonomo and C. G. Scanes. 1997. Ontogeny of Insulin-like Growth Factors (IGF-I and IGF-II) and IGF-Binding Proteins in the Chicken Following Hatching. *Gen. Comp. Endocrinol.* 107:109-117.
- Rosebrough, R. W., J. P. McMurtry and R. Vasilatos-Younken. 1991. Effect of pulsatile or continuous administration of pituitary-derived chicken growth hormone (p-cGH) on lipid metabolism in broiler pullets. *Comp. Biochem. Physiol. A.* 99:207-214.
- Rosselot, G., J. P. McMurtry, R. Vasilatos-Younken and S. Czerwinski. 1995. Effect of exogenous chicken growth hormone (cGH) administration on insulin-like growth factor-I (IGF-I) gene expression in domestic fowl. *Mol. Cell. Endocrinol.* 114:157-66.
- Rotwein, P. 1991. Structure, evolution, expression and regulation of insulin-like growth factors I and II. *Growth Factors* 5:3-18.
- Seo, D. S., J. S. Yun, W. J. Kang, G. J. Jeon, K. C. Hong and Y. Ko. 2001. Association of insulin-like growth factor-I (IGF-I) gene polymorphism with serum IGF-I concentration and body weight in Korean Native Ogol Chicken. *Asian-Aust. J. Anim. Sci.* 14:915-921.
- Simmen, F. A., L. Badinga, M. L. Green, I. Kwak, S. Song and R. C. Simmen. 1998. The porcine insulin-like growth factor system: at the interface of nutrition, growth and reproduction. *J. Nutr.* 128:315-320.
- Spencer, G. S. G., E. Decuyper, J. Buyse and M. Zeman. 1996. Effect of recombinant human insulin-like growth factor-II on weight gain and body composition of broiler chickens. *Poult. Sci.* 75:388-392.
- Tanaka, M., Y. Hayashida, K. Sakaguchi, T. Ohkubo, M. Wakita, S. Hoshino and K. Nakashima. 1996. Growth hormone-independent expression of insulin-like growth factor I messenger ribonucleic acid in extrahepatic tissues of the chicken. *Endocrinology* 137:30-34.
- Tesseraud, S., R. A. Pym, E. Le Bihan-Duval and M. J. Duclos. 2003. Response of broilers selected on carcass quality to dietary protein supply: live performance, muscle development, and circulating insulin-like growth factors (IGF-I and -II). *Poult. Sci.* 82:1011-1016.
- Tomas, F. M., R. A. Pym, J. P. McMurtry and G. L. Francis. 1998. Insulin-like growth factor (IGF)-I but not IGF-II promotes lean growth and feed efficiency in broiler chickens. *Gen. Comp. Endocrinol.* 110:262-275.
- Vasilatos-Younken, R., X. H. Wang, Y. Zhou, J. R. Day, J. P. McMurtry, R. W. Rosebrough, E. Decuyper, N. Buys, V. Darras, J. L. Beard and F. Tomas. 1999. New insights into the mechanism and actions of growth hormone (GH) in poultry. *Domest. Anim. Edocrinol.* 17:181-190.
- Yoon, J., M. S. Rhee, D. S. Seo, B. C. Kim and Y. Ko. 2001. Monitoring of blood cytokines by PIT-1 genotypes in day 150 male pigs. *Asian-Aust. J. Anim. Sci.* 14:1659-1664.
- Yun, J. S., D. S. Seo, M. S. Rhee, S. Oh, B. C. Kim and Y. Ko. 2003a. Relationship of concentrations of endocrine factors at antemortem and postmortem periods to carcass weight and backfat thickness in pigs. *Asian-Aust. J. Anim. Sci.* 16:335-341.
- Yun, J. S., W. J. Kang, D. S. Seo, C. Y. Lee, S. Oh and Y. Ko. 2003b. Relationships of circulating of insuline-like growth factor (IGF)-I and -II to egg production and growth rate in the Korean Native Ogol Chicken. *Asian-Aust. J. Anim. Sci.* 16:481-488.