Effects of gender and gonadectomy on growth and plasma cholesterol levels in pigs

Chong-Eon Lee^{1§}, Nam-Young Kim¹ and Kyu-Il Kim²

¹Jeju Substation, National Institute of Animal Science, RDA, San 175-6 Odeung-dong, Jeju 690-150, Republic of Korea

²Department of Animal Biotechnology, Cheju National University, 1 Ara-dong, Jeju 690-756, Republic of Korea

Received January 19, 2009, Revised February 23, 2009, Accepted Feburary 27, 2009

Abstract

We conducted two studies to determine the effect of gender, gonadectomy (GDX) on growth and plasma cholesterol levels in pigs. In experiment 1, five sham-operated and five GDX female Landrace pigs (26 kg) were allowed to have free access to water and feed up to market weight (approximately 100 kg). Body weight and feed consumption were recorded biweekly, and daily body weight gain, daily feed intake and feed efficiency (gain/feed) were calculated during the feeding period. In experiment 2, 10 male (26 kg) and 10 female (26 kg) Landrace pigs were used; five male and five female pigs were assigned to sham-operated or GDX. Pigs were allowed to have free access to water and a diet without added cholesterol (Table 1) until they were 6 months old (male 104 and female 98 kg) and thereafter they were fed a hypercholesterolemic diet (Table 1) containing 0.5% cholesterol and 0.1% cholate for 10 days. GDX of female pigs increased average daily gain (P<0.05), compared with their sham-operated counterparts during the growing-finishing period, but had no effect (P>0.05) on feed efficiency. Plasma cholesterol levels in pigs fed a hypercholesterolemic diet for 10 days were much higher (P<0.05) in females than in males (161 vs 104 mg/100 mL plasma), and were increased by GDX only in male pigs. HDL-cholesterol/LDL+VLDL-cholesterol ratio appeared to be higher in males than in females, and was not influenced by GDX in either sex. Results suggested that the lower growth rate of female pigs than their male counterparts is attributable to the ovarian activity, and the lower plasma cholesterol level in male than in female pigs fed a hypercholesterolemic diet is due to the testicular activity.

Key Words: Pigs, growth, gonadectomy, cholesterol, sex hormones

Introduction

There have been conflicting reports on the role of ovary (estrogen) and testis (testosterone) in growth and cholesterol metabolism. Our recent study (Lee *et al.*, 2008) demonstrated that a lower growth rate in female rats was due to estrogen, which induced secretion of the stress hormone cortisol. Similarly, other studies (Burgess & Handa, 1992; Lesniewska *et al.*, 1990) also showed gender differences in cortisol secretion, indicating that estrogen elevates basal levels of cortisol, corticosterone and adrenocorticotropin, and their responses to various stimuli in rats. Short-term estradiol treatment to young men increased cortisol and norepinephrine concentrations in the saliva following stress over a placebo control (Kirschbaum *et al.*, 1996).

In the Framingham study, the risk of cardiovascular disease was shown to be significantly increased in women who had taken estrogen (Wilson *et al.*, 1985). Manhem *et al.* (1996) reported that estrogen administration resulted in an enhanced cardiovascular response to mental stress in young menstruating women. In contrast, estrogen has long been known to exert cardioprotective effect (Barrett-Connor & Bush, 1991), although the precise mechanism underlying its benefits is unknown (Sudhir *et al.*, 1997). Antioxidant activity of estrogen has been suggested as

its role in cardioprotection (Walsh *et al.*, 1999), but Santanam *et al.* (1998) found no inhibitory action of estrogen on LDL oxidation at physiological concentrations.

Our previous studies (Lee et al., 1999; Lee et al., 2008) showed a much higher plasma cholesterol level in female than in male rats when they were fed a hypercholesterolemic diet (CD). Naito et al. (1995) proposed that the effect of sex hormones on lipid metabolism is not likely to account for the sex difference in cardiovascular disease. Furthermore, in the 1995 Ancel Keys lecture, Barrett-Connor (1997) indicated that plasma estrogen levels do not explain coronary heart disease (CHD) risk in either sex.

In the present study, we conducted two experiments to determine the effect of gender and gonadectomy on growth, feed efficiency and plasma cholesterol levels, using pigs as a model animal to confirm the findings of our previous study done with rats (Lee *et al.*, 2008).

Materials and Methods

Animals and diets

In experiment 1 conducted to assess the effect of gonadectomy

[§] Corresponding Author: Chong-Eon Lee, Tel. 82-64-754-5721, Fax. 82-64-754-5713, Email. leece00@rda.go.kr

Chong-Eon Lee et al. 39

Table 1. Composition of the experimental diet11

Ingredient	%
Corn	29.2
Wheat	38.0
Wheat bran	0.5
Soybean meal	23.5
Limestone	0.7
Salt	0.3
Monocalcium phosphate	1.4
Tallow	3.2
Molasses	3.0
Vitamin mix	0.1
Mineral mix	0.1
Total	100

^{1) 0,5%} cholesterol and 0,1% cholate were added at the cost of corn in experiment 2,

(GDX) on growth and feed efficiency in female pigs, five sham-operated intact and five GDX Landrace female pigs (26 kg) were allowed to have free access to water and feed until they reached market weight (approximately 100 kg). Body weight and feed consumption were recorded biweekly, and daily body weight gain, daily feed intake and feed efficiency (gain/feed) were calculated during the feeding period.

In experiment 2 conducted to determine the effect of gender and GDX on plasma cholesterol levels in pigs, 10 male (26 kg) and 10 female (26 kg) Landrace pigs were used; five male and five female pigs were assigned to sham-operated intact or GDX. Pigs were allowed to have free access to water and diet without added cholesterol until they were 6 months old (male 104 and female 98 kg) and thereafter a hypercholesterolemic diet (Table 1) containing 0.5% cholesterol and 0.1% cholate for 10 days. All animal management and sampling procedures were in accordance with the Guide for the Care and Use of Agricultural Animals in Agricultural Research and Teaching (Consortium, 1988).

Gonadectomy, plasma sample preparation and carcass measurement

Gonadectomy was carried out surgically by removing testicles or ovaries. Testicles were removed through a small incision in the tip of the scrotum after ligation of the spermatic cords. Ovariectomy was done through a dorsal paramedial incision at the level of the lower pole of the kidneys. The surgery was done under light anesthesia and the incision was closed with stitches. The stitches were removed 7 d after surgery. The control intact pigs were sham-operated leaving the organs intact. The surgery on pigs used in both experiments was done when they were 10 days old. At the end of feeding period, pigs were fasted for 16 hr and killed (Exp. 1), and from the pigs used in experiment 2, blood samples were collected into vacutainer tubes containing EDTA before and after a 10-day feeding period of the hypercholesterolemic diet. Plasma was prepared from the blood

samples by centrifugation, and stored at $-20\,^{\circ}\mathrm{C}$ for later analysis. For pigs used in experiment 1, the average value of measurements at 11^{th} thoracic and 1^{st} lumbar vertebra was used as the back fat thickness. Dressing percentage was calculated from the slaughter weight over the live weight.

Analyses of cholesterol

Total and high-density lipoprotein (HDL) cholesterol concentrations in plasma were determined using commercial assay kits (International Reagent Corp., Tokyo, Japan for the former, and WAKO Pure Chemical Ind., Osaka, Japan for the latter) according to the manufacturer's instruction. Low-density lipoprotein (LDL) plus very-low-density lipoprotein (VLDL) cholesterol concentration was calculated by subtracting HDL cholesterol from the total.

Statistical analysis

The student t-test was used to assess the effect of ovariectomy on pig performance in experiment 1. Data from experiment 2 were analyzed by the two-way analysis of variance (SAS, 1988). In the analysis of variance, the main sources of variation for all variables were gender and gonadectomy. When the F-value in the analysis of variance was significant, the Duncan's multiple range test was used to compare individual means.

Results

Overiectomy increased average daily gain (P<0.05), but had no effect (P>0.5) on feed efficiency during the growing-finishing period (Table 2). Back fat thickness and dressing percentage in ovariectomized pigs tended to be higher (P>0.05) than those found in sham-operated intact female pigs (Table 2). No differences in the plasma total or HDL cholesterol levels and HDL/LDL+VLDL ratio were found before feeding the hypercholesterolemic diet between sexs or between intact and GDX (Table 3). However, plasma cholesterol levels in pigs fed the hypercholesterolemic diet for 10 days were much higher in females than in males (161 vs 104 mg/100 mL plasma). GDX

Table 2. Effect of ovariectomy on daily weight gain and feed efficiency in female pigs¹⁾ - Exp. 1

ltem	Sham	OVX ²⁾	Р
Initial BW, kg	26.1 ± 3.6	26.4 ± 6.3	
Feed intake, kg/d	2.37 ± 0.11	2.58 ± 0.21	0.09
Weight gain, kg/d	0.88 ± 0.06	1.01 ± 0.09	0.04
Gain/feed	0.37 ± 0.04	0.39 ± 0.07	0.44
Carcass traits Back fat thickness ³⁾ , mm	20.4 ± 3.7	23.6 ± 4.4	0.25
Dressina percentage	71.2 ± 1.6	73.2 ± 1.4	0.07

 $^{^{11}}$ Values are means \pm standard deviation of 5 pigs,

Ovariectomized at 10 d of age

³⁾ The average measured at the 11th thoracic and 1st lumbar vertebra.

Table 3. Effect of gender and gonadectorny on plasma cholesterol levels in adult pigs fed a hypercholesterolemic diet for 10 d - Exp. 2

, ,				
	Male		Female	
Item	Sham	GDX ⁴	Sham	GDX⁴
TC ¹ , mg/100 mL				
Before feeding C	96.8 ± 7.9	105.0 ± 11.5	111.0 ± 11.2	107.0 ± 8.2
After feeding C ^c	104.2 ± 21.6 ^a	135.8 ± 22.6 ^{ab}	161.0 ± 28.3 ^b	160.4 ± 26.1 ^b
HDL-C2, mg/100 mL				
Before feeding C	44.4 ± 3.2	43.0 ± 4.8	44.0 ± 3.5	38.8 ± 6.4
After feeding C	44.5 ± 15.4	49.8 ± 8.1	38.2 ± 11.5	41.8 ± 4.6
LDL+VLDL-C ³ , mg/100 mL				
Before feeding C	52.4 ± 10.9	62.0 ± 10.1	67.0 ± 14.0	68.2 ± 11.8
After feeding C ^c	59.8 ± 26.3 ^b	86.0 ± 19.9 ^{ab}	122.8 ± 32.7°	118.6 ± 25.8 ^a
HDL/LDL+VLDL-C				
Before feeding C	0.89 ± 0.25	0.70 ± 0.12	0.68 ± 0.19	0.59 ± 0.22
After feeding C ^c	0.74 ± 0.17^{a}	0.57 ± 0.06^{ab}	0.34 ± 0.09 ^b	0.37 ± 0.05^{b}

Values are means ± standard deviation of 5 pigs.

increased plasma cholesterol levels in male but not in female pigs. HDL/LDL+VLDL ratio appeared to be higher (P<0.01) in male than in female pigs, and was not influenced by GDX in either sex.

Discussion

The present pig study data (Table 2) support our previous findings of the rat study (Lee *et al.*, 2008), in which depressed growth in female rats compared with male rats was mostly due to their ovarian activity. Estrogen secreted in the ovary was responsible for the growth depression because the injection of estradiol markedly depressed growth in GDX rats of either sex, while enhancing plasma cortisol levels. This gender difference was abolished by ovariectomy and was reinstated by estradiol administration (Le Mevel *et al.*, 1979). The pattern of cortisol secretion was also different between sexes in the nonhuman primate macaques, and estradiol implant in castrated male macaques elicited a female pattern of plasma cortisol levels (Norman *et al.*, 1992; Smith & Norman, 1987).

In healthy men, short-term treatment with estradiol led to enhanced hypothalamic-pituitary-adrenal (HPA) and sympathetic responsiveness to psychological stress, resulting in increased adrenocorticotropic hormone (ACTH), cortisol and norepinephrine concentrations in the saliva compared to the placebo (Kirschbaum et al., 1996). Similarly, Burgess and Handa (1992) showed that corticosterone and ACTH levels after a 5-second footshock stress with one mamp were much higher and maintained for a prolonged time in OVX rats when administered with estradiol. In addition, they found that estrogen treatment resulted in a loss of the

glucocorticoid receptor's ability of autoregulation.

Interestingly, estrogen has been reported to have both stimulatory and inhibitory effects on HPA functions, depending on the time after ovariectomy and different doses of estradiol (Luber *et al.*, 1991; Redei *et al.*, 1994). Cortisol, as well as catecholaminergic responses to stress, has also been known to vary with estrous cycle in women (Collins *et al.*, 1985) and in rats (Carey *et al.*, 1995).

Taken together, our data obtained from both rat (Lee *et al.* 2008) and the present pig studies, and others' (Le Mevel *et al.*, 1979; Phillips & Poolsaguan, 1978) clearly indicate that estrogen induces the secretion of hormones that increase with stress, in turn resulting in growth depression. Therefore, ovariectomy may be an economically viable practice to improve growth of female pigs.

Studies have shown marked gender difference in plasma cholesterol levels in Sprague Dawley rats (Lee *et al.*, 1999; Lee *et al.*, 2008) and in guinea pigs (Fernandez *et al.*, 1995), when animals were fed hypercholesterolemic diets. The present study also showed a gender difference in plasma cholesterol levels in pigs, and gonadectomy of male but not female increased plasma total cholesterol and LDL-C concentrations without decreasing HDL-C when pigs were fed the hypercholesterolemic diet (Table 3).

Testosterone replacement therapy in hypogonadal and elderly men showed a beneficial effect on cardiovascular system, decreasing total cholesterol and atherogenic fraction of LDL-C without significant alterations in HDL-C levels or its subfractions HDL₂-C and HDL₃-C (Zgliczynski *et al.*, 1996). Tchernof *et al.* (1997) also showed that increased testosterone levels were associated with reduced triacylglycerol and Apo B; total and LDL cholesterol concentrations; and increased HDL/total cholesterol and HDL₂-C / HDL₃-C ratios. Hypotestosteronemia was observed in Chinese male patients with coronary heart disease (CHD), and positive correlation of plasma testosterone level with plasma HDL-C was found but negative correlation with plasma Lp(a) level, suggesting that testosterone had a protective effect against atherosclerosis (Zhao & Li, 1998).

On the contrary, several studies have shown that testosterone has an adverse effect on cholesterol metabolism by altering lipoprotein profiles. Total testosterone concentrations and sex hormone-binding globulin (SHBG) were significantly associated with LDL size in men (Haffner et al., 1996). Testosterone, per se, when administered at sufficiently high doses and for long durations, can significantly raise serum levels of total cholesterol, triacylglycerol, LDL-C and Apo-B, meanwhile lower HDL-C levels in androgenized women, indicating that male predilection for cardiovascular disease may be due to adverse effects of high androgen levels on lipid and lipoprotein profiles (Goh et al., 1995). In a study done with men, Anderson et al. (1995) also showed that serum HDL-C levels were significantly depressed by intramuscular injection of testosterone enanthate (200 mg per week for 12 months), while plasma total and LDL cholesterol or triacylglycerol levels were not affected.

ab Means in the same row not sharing the same superscripts differ (P<0.01),

^c Gender effect (P<0,01)

¹⁾ Total cholesterol, mg/100 mL

²⁾ High density lipoprotein-cholesterol, mg/100 mL

³⁾ Low density + very low density lipoprotein-cholesterol, mg/100 mL

⁴⁾ Gonadectomized

Handa *et al.* (1997) reported that the plasma free testosterone level was associated with lower levels of HDL-C, but total estradiol was related to elevated levels of HDL-C in Japanese men in their early fifties, supporting that testosterone may be causally associated with atherosclerosis in men through altered lipoprotein metabolism. Male hamsters fed a hypercholesterolemic diet containing 0.05% cholesterol developed greater elevations in plasma total cholesterol and HDL-C levels, and a higher rate of early aortic atherosclerosis, compared with female hamsters (Wilson *et al.*, 1999). Further studies appear to be inevitable to clarify the involvement of testosterone in cholesterol metabolism, such as cholesterol synthesis (e.g., HMG-CoA reductase gene expression) and absorption, or cholesterol oxidation and excretion (e.g., cholesterol 7α-hydroxylase gene expression).

Estrogen's involvement in lipoprotein metabolism (e.g., an increased ratio of HDL to LDL cholesterol) has been implicated in its role in cardioprotection (Campos et al., 1997; Wagner et al., 1991). If estrogen plays any role at all in reducing cardiovascular disease in females, it may be through actions on other than controlling blood cholesterol levels. Naito et al. (1995) indicated that the effect of sex hormones on lipid metabolism is not likely to account for the sex difference in cardiovascular disease because: 1) men with premature myocardial infarction was shown to have increased estrogen levels (Phillips, 1976), and 2) men who had received high doses of estrogen showed an increased frequency of cardiovascular events (Coronary Drug Project Research Group, 1976; Veterans Administration Cooperative Urological Research Group, 1967).

In conclusion, ovarian activity depresses growth in female pigs and thus ovariectomy (or inhibition of estrogen secretion or its activity) can be an economically viable method to improve growth of female pigs. Testis (testosterone) is responsible for the lower plasma cholesterol level in male pigs than in female pigs when they were fed hypercholesterolemic diets.

Literature cited

- Anderson RA, Wallace EM & Wu FCW (1995). Effect of testosterone enanthate on serum lipoproteins in man. *Contraception* 52:115-119.

 Rayrett-Connor E (1997). Say differences in coronary heart disease.
- Barrett-Connor E (1997). Sex differences in coronary heart disease. Circulation 95:252-264.
- Barrett-Connor E & Bush TL (1991). Estrogen and coronary heart disease in women. J Am Med Assoc 265:1861-1867.
- Burgess LH & Handa RJ (1992). Chronic estrogen-induced alterations in adrenocorticotropin and corticosterone secretion, and glucocorticoid receptor-mediated functions in female rats. *Endocrinology* 131:1261-1269.
- Campos H, Walsh BW, Judge H & Sacks FM (1997). Effect of estrogen on very low density lipoprotein and low density lipoprotein subclass metabolism in postmenopausal women. J Clin Endocrinol Metab 82:3955-3963.
- Carey MP, Deterd CH, De Koning J, Helmerhorst F & De Kloet ER (1995). The influence of ovarian steroids on hypothalamic-pituitary-adrenal regulation in the female rat. *J Endocrinol* 144:

- 311-321.
- Collins A, Eneroth P & Landgren BM (1985). Psyconeuroendocrine stress responses and mood as related to the menstrual cycle. *Psychosom Med* 47:512-527.
- Consortium (1988). Guide for the Care and Use of Agricultural Animals in Agricultural Research and Teaching. Savoy, Illinois. 118A
- Coronary Drug Project Research Group (1976). The coronary drug project: initial findings leading to modifications of research protocol. J Am Med Assoc 214:1303-1313.
- Fernandez ML, Vergara-Jimenez M, Romero AL, Erickson SK & McNamara DJ (1995). Gender differences in response to dietary soluble fiber in guinea pigs: effects of pectin, guar gum, and psyllium. J Lipid Res 36:2191-2202.
- Goh HH, Loke DFM & Ratnam SS (1995). The impact of long-term testosterone replacement therapy on lipid and lipoprotein profiles in women. *Maturitas* 21:65-70.
- Haffner SM, Laakso M, Miettinen H, Mykkanen L, Karhapaa P & Rainwater DL (1996). Low levels of sex hormon-binding globulin and testosterone are associated with smaller, dencer low density lipoprotein in normoglycemic men. J Clin Endocrinol Metab 81:3697-3701.
- Handa K, Ishii H, Kono S, Shinchi K, Imanishi K, Mihara H & Tanaka K (1997). Behaviocular stress responses: modulation as a function of menstrual cycle phases. J Psychosom Res 28:475-483.
- Kirschbaum C, Schommer N, Federenko I, Gaab J, Neumann O, Oellers M, Rohleder N, Untiedt O, Hanker J, Pirke KM & Hellhammer DH (1996). Short-term estradiol treatment enhances pituitary-adrenal axis and sympathetic responses to psychosocial stress in healthy young men. *J Clin Endocrinol Metab* 81:3639-3443
- Lee CE, Kang JS & Kim KI (2008). Effects of gender, gonadectomy and sex hormones on growth and plasma cholesterol level in rats. *Ann Nutr Metab* 53:1-5.
- Lee CE, Koh DH, Chang YM & Kim KI (1999). Effects of sex and diets containing alfalfa, laver or pine-needle meals on the plasma cholesterol level in rats. *Nutritional Sciences* 2:46-50.
- Le Mevel JC, Abitbol S, Beraud G & Maniey J (1979). Temporal changes in plasma adrenocorticotropin concentration after repeated neurotropic stress in male and female rats. *Endocrinology* 105:812-827.
- Lesniewska B, Miskowiak B, Nowak M & Malendowicz LK (1990). Sex differences in adrenocortical structure and function. XXVII. The effect of ether stress on ACTH and corticosterone in intact, gonadectomized, and testosterone- or estradiol-replaced rats. Res Exp Med 190:95-103.
- Luber AH, Mobbs CV, Muramatsu M & Pfaff DW (1991). Estrogen receptor messenger RNA expression in rat hypothalamus as a function of genetic sex and estrogen dose. *Endocrinology* 129: 3180-3186.
- Manhem K, Hansson L, Milsom I, Pilhall M & Jern S (1996). Estrogen and progestagen modify the hemodynamic response to mental stress in young women. Acta Obstet Gynecol Scand 75:57-62.
- Naito M, Hayashi T & Iguchi A (1995). New approaches to the prevention of atherosclerosis. *Drugs* 50:440-453.
- Norman RL, Smith CJ, Pappas JD & Hall J (1992). Exposure to ovarian steroids elicits a female pattern of plasma cortisol levels in castrated male macaques. *Steroids* 57:37-43.
- Phillips GB (1976). Evidence for hyperestrogenemia as a risk factor

- for myocardial infarction in men. Lancet 2:14-18.
- Phillips JG & Poolsanguan W (1978). A method to study temporal changes in adrenal activity in relation to sexual status in the female laboratory rat. *Endocrinology* 77:283-291.
- Redei E, Li L, Halasz I, McGivern RF & Aird F (1994). Fast glucocorticoid feedback inhibition of ACTH secretion in the ovariectomized rat: Effect of chronic estrogen and progesterone. *Neuroendocrinology* 60:113-123.
- Santanam N, Shern-Brewer R, McClatchey R, Castellano PZ, Murphy AA, Voelkel S & Parthasarathy S (1998). Estradiol as an antioxidant: incompatible with its physiological concentrations and function. *J Lipid Res* 39:2111-2118.
- SAS (1988). SAS/STATR: User's Guide (Release 6.03). SAS Inst. Inc., Gary, North Carolina. USA
- Smith CJ & Norman RL (1987). Influence of the gonads on cortisol secretion in female rhesus macaques. *Endocrinology* 121:2192-2198.
- Sudhir K, Esler MD, Jennings GL & Komesaroff PA (1997). Estrogen supplementation decreases norepinephrine-induced vasoconstriction and total body norepinephrine spillover in perimenopausal women. *Hypertension* 30:1538-1543.
- Tchernof A, Labrie F, Bélanger A, Prud'homme D, Bouchard C, Tremblay A, Nadeau A & Després JP (1997). Relationship between endogenous steroid hormone, sex hormone-binding globulin and lipoprotein levels in men: contribution of visceral obesity, insulin levels and other metabolic variables. *Atherosclerosis* 133:235-244. Veterans Administration Cooperative Urological Research Group

- (1967). Treatment and survival of patients with cancer of the prostate. Surg Gynecol Obstet 124:1101-1017.
- Wagner JD, Clarkson TB, St Clair RW, Schwenke DC, Shively CA & Adams MR (1991). Estrogen and progesterone replacement therapy reduces low-density lipoprotein accumulation in the coronary arteries of surgically postmenopausal cynomonkeys. J Clin Invest 88:1995-2002.
- Walsh BA, Mullick AE, Walzem RL & Rutledge JC (1999). 17 beta-estradiol reduces tumor necrosis factor-alpha-mediated ldl accumulation in the artery wall. J Lipid Res 40:387-396.
- Wilson PWP, Garrison RJ & Castelli WP (1985). Postmenopausal estrogen use, cigarette smoking, and cardiovascular morbidity in wemen over 50: Framingham Study. New Engl J Med 313:1038-1043
- Wilson TA, Nicolosi RJ, Lawton CW & Babiak J (1999). Gender differences in response to a hypercholesterolemic diet in hamsters: effects on plasma lipoprotein cholesterol concentrations and aortic atherosclerosis. *Atherosclerosis* 146:83-91.
- Zgkiczynsky S, Ossowski M, Slowinska-Srzednicka J, Brzezinska A, Zgliczynski W, Soszynski P, Chotkowska E, Srzednicki M & Sadowski Z (1996). Effect of testosterone replacement therapy on lipids and lipoproteins in hypogonadal and elderly men. *Atherosclerosis* 121:35-43.
- Zhao SP & Li XP (1998). The association of low plasma testosterone level with coronary artery disease in Chinese men. Int J Cardiol 63:161-164.