

**Topic: Correlational analysis of Interleukin 6,
Adiponectin and Lipid indices in women
with Uterine Fibroids.**

ABSTRACT

Background: Uterine fibroids are the most common benign, monoclonal tumours affecting women of reproductive age.

Aim: This study investigated the association between interleukin-6 (IL-6), adiponectin (ADP) and lipid indices in women with uterine fibroids.

Subjects and methods: Sixty (60) participants were recruited for this study. They comprised thirty (30) subjects who had ultrasonographic evidence of uterine fibroids and thirty (30) subjects who had no ultrasonographic evidence of uterine fibroids. Subjects were recruited from the Gynaecology out-patient clinic of LAUTECH Teaching Hospital Ogbomoso, Nigeria. Anthropometric measurements were performed using standard method. About 5mL of venous blood was collected from each study participant and was dispensed into a plain bottle. Serum was obtained after clotting and centrifugation and was aliquoted into a small vial which was stored at -20°C until time of analysis. Interleukin-6 (IL-6), adiponectin (ADP), total cholesterol (TC), triglycerides (TG) and high density lipoprotein (HDL) were determined using enzyme linked immunosorbent assay (ELISA) and colorimetric method as appropriate. Low density lipoprotein (LDL) was estimated using Friedwald's equation. Data analysis was done using Student's t-test for comparison of variables and Pearson's correlation was used to determine the relationship between variables. *P*-value less than 0.05 was considered significant.

Results: Women with uterine fibroids had significantly elevated body mass index (BMI), waist circumference (WC) and waist hip ratio (WHR) when compared with the controls (*P*<0.05). The mean values of TC, TG, LDL and IL-6 were significantly elevated in women with uterine fibroids when compared with the control subjects (*P*<0.05). Women with uterine fibroids had significantly reduced levels of ADP when compared with controls (*P*<0.05) and there was no significant difference in the levels of HDL when both case and control subjects were compared (*P*>0.05). Interleukin-6 had a significant positive correlation with BMI (*R*=0.362, *P*=0.03), WC (*R*=0.456, *P*=0.02), WHR (*R*=0.374, *P*=0.04), TC (*R*=0.735, *P*=0.00), TG (*R*=0.429, *P*=0.01), and LDL (*R*=0.606, *P*=0.04). Interleukin-6 also had a significant negative correlation with HDL (*R*= -0.590, *P*=0.03) and ADP (*R*= -0.527, *P*=0.02).

Conclusion: This study demonstrated a significant direct relationship between IL-6 and dyslipidemia in women with uterine fibroids. Also, we observed a noteworthy inverse relationship between IL-6 and

35 adiponectin in women with uterine leiomyomas, thus emphasizing the low grade chronic inflammatory
36 state associated with uterine leiomyomas.

37 *Keywords: Uterine leiomyoma, interleukin-6, adiponectin, adipocytes, lipid profile.*

38

39

40 **1. INTRODUCTION**

41 Uterine fibroids (leiomyomas) are benign, monoclonal tumors originating from the myometrium and the
42 most common pelvic tumors of the female reproductive system (1). The symptoms of uterine fibroids
43 include menstrual disturbances such as menorrhagia, dysmenorrhea, intermenstrual bleeding,
44 dyspareunia and noncyclic pelvic pain (2) and pressure symptoms such as a sensation of bloatedness,
45 increased urinary frequency, and bowel disturbances (3). It has also been associated with impaired
46 reproductive outcomes, leading to subfertility, abortion, preterm labor and delivery, and cesarean delivery
47 (3, 4). The estimated prevalence of uterine fibroids in Southwest Nigeria is 7.0% (5).

48 The occurrence of fibroids has been linked with the hormones: estrogens, and progesterone. These
49 hormones are also responsible for fibroid growth. Some studies have reported the up-regulation of
50 progesterone receptor-A (PR-A) and PR-B, oestrogen receptor-alpha (ER- α) and -beta (ER- β) as well as
51 messenger RNA (mRNA) and protein expression in myoma tissue compared with normal myometrium (6,
52 7). Leiomyoma growth is influenced by progesterone interaction with some growth factors; it upregulates
53 the endothelial growth factor (8) and transforming growth factor-beta 3 (bimodal action) expression (9).
54 On one hand, progesterone seems to down-regulate interleukin growth factor-1 expression through PR-B
55 while PRA appears to inhibit this function (8). Other factors influencing the development of myomas
56 include: age, race, obesity, lifestyle such as smoking, alcohol and other drugs which can affect the uterine
57 wall (10). It is therefore hypothesized that oestrogens exert growth stimulatory effects on leiomyomas
58 intermediated by cytokines, growth factors, or apoptotic factors (11). The immune system plays an
59 important role in the aetiology of leiomyomas. This includes the chronic inflammation associated with
60 cytokine release by the immune system as well as the undifferentiated cells (5, 12, 13).

61 In obesity, adipocytes undergo molecular and cellular alterations which includes hypertrophy; with
62 resultant effect on systemic metabolism, thus leading to insulin resistance, metabolic disorders and
63 inflammatory responses due to the release of pro-inflammatory factors from the adipocytes (14). Also,
64 adipocytes secrete adiponectin and previous studies have demonstrated that adiponectin has insulin-
65 sensitizing and anti-inflammatory effects (15, 16).

66

67 Since obesity has been implicated as a predisposing factor for leiomyomas, there is likelihood of
68 dyslipidaemia. Also, studies have reported a link between tumor necrosis factor-alpha (TNF- α), a pro

69 inflammatory cytokine, and the growth of uterine fibroid (17, 18). This study is therefore focused on
70 determining adiponectin, Interleukin-6 and lipid indices in order to obtain valuable information on the
71 correlation between them in women with uterine fibroids.

72

73 **2. MATERIAL AND METHODS**

74

75 **2.1 Subjects**

76 A total of sixty (60) participants were recruited for this study. They comprised thirty (30) subjects who had
77 ultrasonographic evidence of uterine fibroids and thirty (30) subjects who had no ultrasonographic
78 evidence of uterine fibroids. Subjects were recruited from the Gynaecology out-patient clinic of LAUTECH
79 Teaching Hospital Ogbomoso, Nigeria. Subjects were infertile women referred for ultrasound examination
80 and based upon this examination, they were stratified into women with fibroid (case) and women without
81 fibroid (control). All subjects had normal menstrual cycle and they were not on any hormonal medication.
82 All subjects consented to participate in the study through writing.

83 A short structured questionnaire was administered to each study participant to obtain information on
84 smoking habits, medications and established diseases. Persons diagnosed with dyslipidemia and other
85 metabolic conditions, record of alcoholism, smoking, usage of medications that affect lipid status,
86 pregnant women and those who refuse to give consent were excluded from this study.

87 **2.2 Anthropometric measurement**

88 Body weight, height, body mass index (BMI), waist circumference (WC) and waist to hip ratio (WHR) were
89 obtained from the participants by standard methods as described by Charles-Davies *et al.* (19).

90 **2.3 Sample Collection and Biochemical Analyses**

91 Blood sample (5mL) was collected and dispensed into a plain bottle. Serum was obtained after clotting
92 and centrifugation and was aliquoted into a small vial which was stored at -20°C until time of analysis.
93 Serum Total Cholesterol (TC) and Triglyceride (TG) were determined using standard enzymatic method
94 while high density lipoprotein-cholesterol (HDL-C) was determined by a two-step procedure as previously
95 described by Adediji *et al.* (20). LDL cholesterol was estimated using Friedewald's equation; [Total
96 cholesterol – HDL cholesterol – (Triglyceride/ 5) mg/dL].

97 Serum Adiponectin and IL-6 were determined using ELISA kits manufactured by Elabscience
98 Biotechnology Co. Ltd (Wuhan, China) with the procedure as previously carried out by Ashraf *et al.* (21).

99 **2.3.1 Principle of ELISA**

100 In this assay, the antigen – adiponectin or IL-6 present in each sample reacts with the corresponding
101 antibody adsorbed to the surface of solid-phase polystyrene microtitre wells. On removal of unbound

102 proteins by washing, the antibodies conjugated with horseradish peroxidase (HRP) form complexes with
103 the previously bound antigen following the addition of a chromogenic substrate, 3, 3', 5, 5'-
104 tetramethylbenzidine (TMB). The absorbance at 450 nm is a measure of the concentration of the 'antigen'
105 in the test sample.

106 107 **2.3.2 ELISA Procedure**

108 A gradient of standard concentrations were prepared from the concentrated standard through serial
109 dilution to cover the expected assay range. One hundred microliters of the standards and sera were
110 pipetted into microwells already coated with specific antibodies and incubated at 37°C for 90 min.
111 Following incubation, the wells were aspirated of their contents without washing and 100µl of biotinylated
112 detection antibody was added to each well and incubated for 60 minutes at 37°C, after which each well
113 was completely filled with appropriate wash solution. The plate was washed three times.

114 One hundred microliters of appropriately diluted enzyme-antibody conjugate was pipetted into each well
115 and the plate was incubated at 37°C for 30 minutes. After incubation, another process of washing was
116 performed as described above and 90µl of TMB substrate solution was added to each well. This was
117 followed by incubation for 15 minutes at 37°C after which 50µl of stop solution was added to each well.
118 The absorbance (at 450 nm) was determined using a microplate reader.

119 120 **2.4 Statistical analysis**

121 Data analysis was done using SPSS version 20 (IBM, Armonk, NY, USA). All values were expressed as
122 mean±standard deviation for test and control groups. Comparison of variables was done using Student's
123 t-test and Pearson's correlation was used to determine the relationship between variables. *P* less than
124 0.05 was considered to be statistically significant.

125 126 **3. RESULTS**

127 The anthropometric and biochemical parameters of the study participants are shown in table 1. The mean
128 values of BMI, WC, WHR, TC, TG, LDL and IL-6 were elevated in women with uterine fibroids when
129 compared with the control subjects. The mean values of ADP were significantly reduced in women with
130 uterine fibroids when compared with the controls and there was statistical significant difference in the
131 levels of HDL when both test and control subjects were compared.

132
133 Table 2 shows the correlation IL-6, anthropometric and other biochemical parameters in women with
134 uterine fibroids. IL-6 had a significant positive correlation with BMI ($R=0.362$, $P=0.03$), WC ($R=0.456$,
135 $P=0.02$), WHR ($R=0.374$, $P=0.04$), TC ($R=0.735$, $P=0.00$), TG ($R=0.429$, $P=0.01$), LDL ($R=0.606$,
136 $P=0.04$). Also, IL-6 had a significant negative correlation with HDL ($R= -0.590$, $P=0.03$) and ADP
137 ($R= -0.527$, $P=0.02$) in women with uterine fibroids.

138
139

140 **Table 1: Anthropometric and biochemical parameters in study participants.**

141

Parameters	Subjects (n=30)	Control (n=30)	P-value
BMI (kgm ⁻²)	28.6 ±2.76	24.5 ±4.68	0.04*
WC (cm)	86.3 ±11.3	77.5 ±8.7	0.02*
WHR	1.15±0.08	1.04±0.13	0.00*
TC (mg/dL)	193.1±24.2	146.3±38.1	0.00*
TG (mg/dL)	180.6±29.9	131.6±45.6	0.00*
HDL (mg/dL)	42.5±5.43	43.3±3.38	0.56
LDL (mg/dL)	114.5±19.5	76.5±31.0	0.00*
IL-6 (pg/mL)	352.6±39.7	233.5±43.6	0.00*
ADP (µg/L)	121.9±33.3	189.2±81.8	0.00*

142 *Statistically significant at P<0.05. Results are expressed as mean±standard deviation. BMI-Body mass
 143 index; WC- Waist circumference; WHR- Waist to hip ratio; TC- Total cholesterol; TG- Triglycerides; HDL-
 144 High density lipoprotein; LDL- Low density lipoprotein; IL-6- Interleukin-6; ADP- Adiponectin.

145

146

147 **Table 2: Correlation between IL-6, anthropometric and other biochemical parameters in women**
 148 **with uterine fibroids**

149

Parameters	R	P- value
BMI [†]	0.362	0.03*
WC [†]	0.456	0.02*
WHR [†]	0.374	0.04*
TC [†]	0.735	0.00*
TG [†]	0.429	0.01*
HDL [†]	-0.590	0.03*
LDL [†]	0.606	0.04*
ADP [†]	-0.527	0.02*

150

151 *Statistically significant at P<0.05. BMI[†]= Correlation between IL-6 and BMI; WC[†]= Correlation between
 152 IL-6 and WC; WHR[†]= Correlation between IL-6 and WHR; TC[†]= Correlation between IL-6 and TC;
 153 TG[†]= Correlation between IL-6 and TG; HDL[†]= Correlation between IL-6 and HDL; LDL[†]= Correlation
 154 between IL-6 and LDL; ADP[†]= Correlation between IL-6 and ADP.

155

156

157

158

159 4. DISCUSSION

160
161 Uterine fibroids continue to be the most common gynecological benign tumours in women of reproductive
162 age and there are multiple risk factors which have been implicated in the development of uterine
163 leiomyoma.

164
165 In this study, body size and central obesity were measured by BMI, WC and WHR and we observed
166 increased BMI, WC and WHR in women with uterine fibroids when compared with controls. These
167 findings are consistent with previous studies that reported a significant positive association between
168 central obesity and incidence of uterine fibroids (22-24). This observation could be attributed to increased
169 adiposity which accentuates the aromatization of androgens to estrogens and the consequent
170 hyperestrogenic state is responsible for the growth of uterine fibroids in obese women (25-27).

171
172 Furthermore, the progressive adipocyte hypertrophy accentuates low grade chronic inflammation which is
173 thought to play a major role in the progression of uterine fibroids. Adipocytes secrete proinflammatory
174 cytokines like IL-6 and they also secrete anti-inflammatory mediators like adiponectin. In our study, we
175 observed a significant increase in the levels of IL-6 and a significant reduction in the levels of adiponectin,
176 when women with uterine fibroids were compared with the controls. This observation is further supported
177 by our observed significant inverse relationship between IL-6 and adiponectin in women with uterine
178 fibroids. The observed elevated IL-6 could be attributed to excessive secretion of IL-6 by peritoneal
179 macrophages which are abundantly present in enlarged adipocytes and are actively involved in their
180 activities (28-30). IL-6 and other proinflammatory factors produced due to excessive fat accumulation
181 have a direct effect on the myometrium and this consequently enhances the growth of uterine
182 leiomyomas (4, 30).

183
184 Moreover, the observed low serum adiponectin could also be attributed to increased adiposity which
185 promotes excessive secretion of IL-6 and this in turn, stymies the production of adiponectin in adipocytes
186 (21, 31). Previous studies have reported that low adiponectin levels has a direct relationship with
187 increased insulin levels and insulin is a potent regulator of sex hormone binding globulin (SHBG) (31, 32).
188 Thus, increased insulin level attenuates hepatic production of SHBG, leading to an increase in the
189 circulating levels of androgens which further enhances the growth of uterine leiomyomas in obese women
190 (24, 33).

191
192 In this study, we also observed significant increase in the levels of TC, TG and LDL when the case
193 subjects were compared with controls however, there was no statistical significant difference in the levels
194 of HDL-C. We also observed a significant positive correlation between IL-6, TG, TC and LDL while there
195 was significant inverse relationship between IL-6 and HDL-C in women with uterine fibroids. These
196 observations are consistent with previous studies (23, 24, 27) and could be attributed to the influence of

197 estrogen on TG accumulation and the ability of IL-6 to accentuate lipid abnormalities through its
198 pronounced inhibitory effect on the activities of adipocyte lipoprotein lipase. (34-36).

199

200 **5. CONCLUSION**

201 This study revealed a significant direct relationship between IL-6 and dyslipidemia in women with uterine
202 fibroids. Also, we observed a noteworthy inverse relationship between IL-6 and adiponectin in women
203 with uterine leiomyomas, thus emphasizing the low grade chronic inflammatory state associated with
204 uterine leiomyomas.

205 The major limitation of the study was sample size, therefore a cross sectional study on a larger population
206 is desirable in confirming our findings.

207

208 **COMPETING INTERESTS**

209 Authors have declared that no competing interests exist.

210

211

212 **REFERENCES**

213

2141) Reis F., Bloise E. and Ortiga-Carvalho T. Hormones and Pathogenesis of Uterine fibroid. Journal Best
215 Practice & Research Clinical Obstetrics and Gynaecology. 2015; 34:13-24.

2162) Eltokhi HM, Modi MN, Weston M, Armstrong AY, Stewart EA. The health disparities of uterine fibroid
217 tumors for African American women: a public health issue. Am J Obstet Gynecol. 2014; 210: 194-199.

2183) Andrea Ciavattini, Jacopo DiGiuseppe, Piergiorgio Stortoni, Nina Montik, Stefano Giannubilo, Pietro Litta,
219 Md. Sorifullslam, Andrea Tranquilli, Fernando Reis, and Pasquapina Ciarmela. Uterine Fibroids:
220 Pathogenesis and Interactions with Endometrium and Endomyometrial Junction. Obs&Gyne Int. 2013;
221 2013:1-11.

2224) Protic O, Toti P, Islam S, Occhini R, Giannubilo R, Catherino H, Cinti S, Petraglia F, Ciavattini A.,
223 Castellucci M, et al. "Possible Involvement of Inflammatory/Reparative Processes in the Development of
224 Uterine Fibroids". Cell Tissue Res. 2016; 364: 415–427.

2255) Ukwenya V, Maduemezia N, Afolayan O, Alese O and Thomas W. (2015). "Prevalence of Uterine Fibroid
226 in a South-Western Nigerian Population: A sonographic study. Journal of Exp&Clin.Anat. 2015; 14(1):24-
227 29.

2286) Ying Z and Weiyuan Z. Dual actions of progesterone on uterine leiomyoma correlate with the ratio of
229 progesterone receptor A: B. Gynecological Endocrinology. 2009; 25(8):520–523.

- 2307) Ishikawa H, Ishi K, Serna V. Progesterone is Essential for Maintenance and Growth of Uterine
231 Leiomyoma". *Endocrinology*. 2010; 151(24): 33-42.
- 2328) Yin P, Lin Z, Cheng Y et al. Progesterone receptor regulates Bcl-2 gene expression through direct binding
233 to its promoter region in uterine leiomyoma cells. *Journal of Clinical Endocrinology and Metabolism*. 2007;
234 92 (11):4459–4466.
- 2359) Arici A and Sozen I. Transforming growth factor- β 3 is expressed at high levels in leiomyoma where it
236 stimulates fibronectin expression and cell proliferation. *Fertility and Sterility*. 2000; 73(5):1006–1011.
- 237 10) Radmila Sparic, Ljiljana Mirkovic, Antonio Malvasi and Andrea Tinelli. Epidemiology of Uterine Myomas:
238 A Review. *International Journal of Fertility and Sterility*. 2016; 9(4):424-435.
- 239 11) Grings A.O, Lora V, Ferreira D.G, Brum I.S, Corleta H.V.E and Capp E. "Protein Expression of Estrogen
240 Receptors α and β and Aromatase in Myometrium and Uterine Leiomyoma. Gynecologic and Obstetric
241 Investigation. 2012; 73(2):113-117.
- 242 12) Wegienka G. "Are uterine leiomyoma a consequence of a chronically inflammatory immune system?"
243 Medical Hypotheses. 2012; 79(2):226–231.
- 244 13) Monia Orciani, Miriam Caffarini, Alessandra Biagini, Guendalina Lucarini, Giovanni Delli Carpini,
245 Antonella Berretta, Roberto Di Primio and Andrea Ciavattini. Chronic Inflammation May Enhance
246 Leiomyoma Development by the Involvement of Progenitor Cells. *Stem Cells International*; 2018; 2018:
247 1-13. Article ID 1716246.
- 248 14) Greenberg AS, Obin MS. Obesity and the role of adipose tissue in inflammation and metabolism. *Am J*
249 *Clin Nutr*. 2006; 83(Suppl):461S-465S.
- 250 15) Tong, H. V., Luu, N. K., Son, H. A., Hoan, N. V., Hung, T. T., Velavan, T. P., & Toan, N. L. (2017).
251 Adiponectin and pro-inflammatory cytokines are modulated in Vietnamese patients with type 2 diabetes
252 mellitus. *Journal of diabetes investigation*, 8(3), 295-305.
- 253 16) Ochiai H, Shirasawa T, Nishimura R, Nanri H, Ohtsu T, Hoshino H, et al. Abdominal obesity and serum
254 adiponectin complexes among population-based elementary school children in Japan: A cross-sectional
255 study. *BMC Pediatr*. 2014; 14:81.

- 256 17) Michal Ciebiera, Marta Włodarczyk, Magdalena Zgliczynska, Krzysztof Łukaszuk, Błażej Meczekalski,
257 Christopher Kobierzycki, Tomasz Łozinski and Grzegorz Jakiel. The Role of Tumor Necrosis Factor α in
258 the Biology of Uterine Fibroids and the Related Symptoms. *Int. J. Mol. Sci.* 2018; 19:1-26.
- 259 18) Geagea AG, Mallat S, Matar CF, Zerbe R, Filfili E, Francis M, Haidar H, Jurjus A. Adiponectin and
260 inflammation in health and disease: An update. *Open Medicine Journal.* 2018; 5 20 – 32
- 261 19) Charles-Davies MA, Arinola OG, Fasanmade AA, Olaniyi JA, Oyewole OO, Owolabi MO, et al. Indices of
262 metabolic syndrome in 534 apparently healthy traders in a local market in Ibadan, Nigeria. *Journal of*
263 *US-China Medical Science.* 2012; 9(2):91-100.
- 264 20) Adediji IO, Taiwo TD, Adepoju PO, Adelakun AA, Akinleye WA, Afolabi JO. Cortisol and Metabolic
265 Syndrome Components in Obese and Overweight Young Adults of a Nigerian Private University.
266 *AJMAH.* 2018; 11(2): 1-7.
- 267 21) Hamid Ashraf, Bashir Ahmad Laway, Dil Afroze, Arshad Iqbal Wani. Evaluation of Proinflammatory
268 Cytokines in Obese vs Non-obese Patients with Metabolic Syndrome. *Ind.J.Endo&Metab.* 2018;
269 22(6):751-756.
- 270 22) Luoto R, Kaprio J, Rutanen EM, Taipale P, Perola M, Koskenvuo M. Heritability and risk factors of uterine
271 fibroids the Finnish twin Cohort study. *Maturitas.* 2000; 37(1): 1526.
- 272 23) Takeda T, Sakata M, Isobe A, Miyake A, Nishimoto F, Ota Y, Kamiura S, Kimura T. Relationship
273 between metabolic syndrome and fibroids: a case-control study. *Gynecol Obstet Invest,* 66: 2008, 14-17.
- 274 24) Young Jin Tak, Sang Yeoup Lee, Sun Ki Park, Yun Jin Kim, Jeong Gyu Lee, Dong Wook Jeong, Seung
275 Chul Kim, In Joo Kim, Yu Hyun Yi. Association between uterine leiomyomas and metabolic syndrome in
276 parous perimenopausal women: A case-control study. *Medicine.* 2016; 95:46(e5325).
- 277 25) Okolo S. Incidence, aetiology and epidemiology of uterine fibroids. *Best Pract Res Clin Obstet Gynaecol.*
278 2008; 22(4): 571-588.
- 279 26) Wise LA, Laughlin-Tommaso SK. Uterine leiomyomata. In: Goldman MB, Troisi R, Rexrode KM, editors.
280 *Women and Health.* San Diego: Academic Press; 2013; 285-306.
- 281 27) He Y, Zeng Q, Li X, Liu B, Wang P. The association between subclinical atherosclerosis and uterine
282 fibroids. *PLoS One.* 2013; 8(2): e57089.

- 283 28) Fantuzzi, G. Adipose tissue, adipokines, and inflammation. *J. Allergy Clin. Immunol.* 2005; 115: 911–
284 919.
- 285 29) Essam El-Din R Othman, Daniela Homung, Hosam T. Salem, Essam A. Khalifa, Tarek H. El-Metwally,
286 Ayam Al-Hendy. Serum cytokines as biomarkers for non-surgical prediction of endometriosis.
287 *Eur.J.Obst&Gyn&ReproBiol.* 2007; 137 (2008):240-246.
- 288 30) Protic O, Islam M.S, Greco S, Giannubilo S.R, Lamanna P, Petraglia F, Ciavattini A, Castellucci M, Hinz
289 B, Ciarmela P. Activin A in inflammation, tissue repair and fibrosis: Possible role as inflammatory and
290 fibrotic mediator of uterine fibroid development and growth. *Semin. Reprod. Med.* 2017; 35; 499–509.
- 291 31) Noriyuki Ouchi, Koji Ohashi, Rei Shibata and Toyoaki Murohara. Adipocytokines and Obesity-linked
292 disorders. *Nagoya J. Med. Sci.*2012; 74:19-30.
- 293 32) Stefan N, Vojarova B, Funahashi T, Matsuzawa Y, Weyer C, Lindsay RS, Youngren JF, Havel PJ,
294 Pratley RE, Bogardus C, Tataranni PA: Plasma adiponectin concentration is associated with skeletal
295 muscle insulin receptor tyrosine phosphorylation, and low plasma concentration precedes a decrease in
296 whole-body insulin sensitivity in humans. *Diabetes.* 2002; 50:1884–1888.
- 297 33) Kong, S., Hou, J., Xia, M., Yang, Y., Xu, A., & Tang. Association of Hyperglycemia, Hyperlipemia with
298 the Risk of Uterine Leiomyomata: A Case-Control Study. *Cancer Cell Research.* 2014; 2(1):37-41.
- 299 34) Jose-Manuel Fernandez-Real, Montserrat Broch, Joan Vendrell, Cristobal Richart, and Wifredo Ricart.
300 Interleukin-6 Gene Polymorphism and Lipid Abnormalities in Healthy Subjects. *J.Clin.Endo&Metab.*2000;
301 85(3):1334-1339.
- 302 35) Khovidhunkit W, Kim M.S, Menon R.A, Shigenaga J.K, Moser A.H, Feingold K.R, Grunfeld C. Effects of
303 infection and inflammation on lipid and lipoprotein metabolism: mechanism and consequences to the
304 host. *J. Lipid Res.*2004; 45:1169-1196.
- 305 36) Shen M, Shi H. Sex hormones and their receptors regulate energy homeostasis. *Int. J. Endocrinol.* 2015;
306 2015:29478.