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## **Environmental Influences in Normal Weight Women with History of Recurrent Pregnancy Loss**

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### **Authors' contributions**

*This work was carried out in collaboration between all authors. Author OBA participated in conception and design of the study, managed the literature searches, did the laboratory and statistical analyses of the study, interpreted data, and prepared the first draft of the study. Author MAC participated in the conception and design of the study, literature searches and interpretation of data, critical revision of article for important intellectual content and wrote the final version to be published. Author MAO participated in the design of the study, enrolment of participants in the study and revision of article for important intellectual content. All authors read and approved the final manuscript.*

**Research Article**

**Received 6<sup>th</sup> March 2012**  
**Accepted 7<sup>th</sup> March 2013**  
**Published 12<sup>th</sup> July 2013**

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### **ABSTRACT**

**Aim:** The study investigated the possible relationship of luteal phase progesterone, toxic heavy metals and nutrients in normal-weight women with history of recurrent pregnancy loss for adequate management.

**Study Design:** Cross sectional study.

**Place and Duration of Study:** The Obstetrics and Gynaecology Clinic, University College Hospital, Ibadan, Olabisi Onabanjo University Teaching Hospital, Shagamu and State Hospital, Ijebu-Ode; the University of Ibadan and environs between April and September, 2009.

**Methodology:** 90 apparently healthy women with normal weight aged 18-45years with

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regular and ovulatory cycles of 26-30 days were enrolled. They were 60 women with history of recurrent pregnancy loss (cases) age-matched with 30 women without history of recurrent pregnancy loss (controls). Demographic and anthropometric measurements were done by standard methods. Progesterone was determined by enzyme immunoassay (Immunometrics UK Ltd). Total cholesterol, triglyceride and high density lipoprotein were performed by enzymatic methods (Randox laboratories, USA) while low density lipoprotein was calculated using Friedwald's formula. Zinc, selenium, chromium, manganese, iron, magnesium, copper, lead, and cadmium were estimated by atomic absorption spectrophotometry while Vitamin E was measured by high performance liquid chromatography. Statistical analysis was done using SPSS version 16.0.

**Results:** Results showed significantly higher levels of triglycerides, cadmium and lead, and significantly lower levels of progesterone, iron, copper, magnesium, chromium, selenium and vitamin E in cases compared with controls ( $p < 0.013$ ).

**Conclusion:** Oxidative stress mechanisms in normal-weight women with history of recurrent pregnancy loss were implicated. Healthcare policies should focus on pollution reduction and increase awareness on healthy diet for optimal periconceptional micronutrient requirements.

*Keywords: Progesterone; metabolic syndrome; micronutrients; endocrine disruptors; recurrent pregnancy loss.*

## 1. INTRODUCTION

Early pregnancy loss is one of the most common disorders of human pregnancy and a public health concern with risk factors that have not been elucidated [1,2,3]. Recurrent pregnancy loss (RPL), defined as 3 or more consecutive pregnancy losses before 20 weeks of gestation affects 0.5%–3% of women in the reproductive age group. 50%–60% of these pregnancies are idiopathic and highly frustrating for both patients and physicians [4].

Advanced maternal age (>35yrs), obesity, increased number of previous pregnancy, and luteal phase insufficiency with reduced progesterone levels are known risk factors in the aetiology of RPL with less clearly defined mechanisms [5,6,7,8,9]. Recent reports implicate environmental and lifestyle factors in RPL [2,10].

Cadmium (Cd) and lead (Pb) are toxic heavy metals [10,11] associated with several modern industrial processes and obtained through the food chain [10,12]. Accumulation of heavy metals like Cd and Pb in African cat fish (*Clarias gariepinus*) from Ogun River near six major industries in the region of this present study has been observed [13]. Cd and Pb disrupt endocrine function and are recognized risk factors for hormonal changes, impaired placental development, congenital malformations, fetal growth restriction and spontaneous abortion [10]. The negative impact of heavy metals on pituitary and ovarian functions has been reported [14]. Correlations of heavy metal excretion with different hormones- progesterone, oestradiol, prolactin, thyroid stimulating hormone have been reported [14].

Disruption of metal ion homeostasis leads to increased formation of potentially destructive reactive oxygen species (ROS), which overwhelms body antioxidant protection. The resultant oxidative stress (OS) subsequently induces DNA damage, lipid peroxidation, protein modification and eventual cellular damage [11,15]. Increased free radical activity has been shown in the pathogenesis of recurrent abortion [16]. Systemic and placental oxidative stress have been implicated in the pathophysiology of RPL. Depletion of antioxidant

defences caused by increased oxidative stress has been demonstrated in RPL [2,4,16]. Diet, particularly its constituent antioxidants, and oxidative stress may influence the timing and maintenance of a viable pregnancy [2]. Thus micronutrient deficiencies might affect fertility, embryogenesis and placentation and their prophylactic use in preventing these adverse pregnancy outcomes have been advocated [17]. This study was designed to understand the roles of heavy metals and nutrients and their specific influences on progesterone in normal weight women with RPL to elucidate underlying mechanisms for both prevention and informed management.

## **2. MATERIALS AND METHODS**

### **2.1 Subjects**

A total of 90 apparently healthy women with normal body mass index (BMI) of 18.5-24.9 kg/m<sup>2</sup> [18] with regular and ovulatory cycles of 26-30 days aged 18-45 years were recruited after informed consent for this prospective cross sectional study using stratified random sampling method. These were 60 non-pregnant women with history of recurrent abortion and at least 3 pregnancy losses before 20 weeks of gestation (cases) age matched with 30 non-pregnant women without history of recurrent abortion with at least one child (controls). All cases had normal values of luteinising hormone, and normal transvaginal/ abdominal scan done recently without fibroid, polycystic ovarian syndrome or abdominal mass. None of the women was on lipid lowering or hormonal drugs. Individuals with cardiovascular diseases, diabetes and renal failure and those who did not give consent were excluded from the study. The cases were selected at the Obstetrics and Gynaecology Department of the University College Hospital, Ibadan (UCH), Olabisi Onabanjo University Teaching Hospital, Shagamu and State Hospital, Ijebu-Ode while controls were selected from University of Ibadan and environs. Ethical approval was obtained from the University of Ibadan/UCH Ethical Board.

### **2.2 Collection of Blood**

10ml of fasting venous blood sample was aseptically taken from the antecubital fossa vein with minimal stasis on day 21 of the menstrual cycle of all women. 5ml was dispensed into ethylene diamine tetra-acetic acid (EDTA) tubes for the estimation of lipids while the remaining 5ml were dispensed into plain tubes and kept for 1-2 hours to clot for the estimation of micronutrients, heavy metals and progesterone. The samples were then centrifuged at 500g for 5mins to obtain plasma and serum respectively, and stored at -20°C until analysis.

### **2.3 Demographic Characteristics**

Semi structured pre-test questionnaire was completed by each subject in order to obtain demographic data. The demographic characteristics tested were age and age at menarche (AM), length of menstrual cycle (LMC), number of previous pregnancies (NPP) and number of live birth(s) (NLB).

### **2.4 Anthropometric Characteristics**

The anthropometric parameters measured were weight, height, body mass index (BMI), waist circumference (WC), hip circumference (HC), and waist hip ratio (WHR) according to methods described by Umoh et al. [19].

## **2.5 Biochemical Characteristics**

The biochemical variables tested were progesterone, lipids [triglyceride (TG); high density lipoprotein (HDL), total cholesterol (TC) and low density lipoprotein-cholesterol (LDL)], micronutrients [zinc (Zn), iron (Fe), copper(Cu), manganese (Mn), magnesium(Mg), chromium (Cr) and selenium (Se) and Vitamin E] and heavy metals [lead (Pb) and cadmium (Cd)]. Progesterone was estimated by enzyme immunoassay method [20] (Immunometrics UK. Ltd.). Total cholesterol was estimated by enzymatic method [21]. TG was analysed by method described by Trinder [22]. HDL was estimated by precipitating low density lipoprotein cholesterol, very low density lipoprotein cholesterol and chylomicron fractions using phosphotungstic acid in the presence of magnesium ions. The HDL cholesterol which remained in the supernatant was estimated using the total cholesterol method. Low density lipoprotein cholesterol was calculated using formula of Friedwald et al. [23]. All analyses of lipid fractions were done by commercially produced kits (Randox Laboratories, USA). Micronutrients(Zn, Fe, Cu, Mn, Mg, Cr, Se) and heavy metals (Pb and Cd) were estimated by Atomic Absorption Spectrophotometer (Beck 200) based on the direct method described by Kaneko [24] while vitamin E was estimated by high performance liquid chromatography (Waters, USA).

## **2.6 Statistical Analysis**

Statistical package for social scientist (SPSS) version 16.0 was employed for the analyses of data. Student's t –test was used for the comparison of quantitative variables. Pearson's correlation coefficient was used for the relationship between quantitative variables.  $p < 0.05$  was regarded as significant.

## **3. RESULTS AND DISCUSSION**

### **3.1 Lipids, Progesterone, Demographic and Anthropometric Characteristics**

Table 1 shows the comparison of lipids, progesterone, demographic and anthropometric characteristics between women with recurrent abortion (cases) and controls. Of all parameters tested, only NPP and TG were significantly higher while progesterone and NLB were significantly lower in cases compared with controls ( $p < 0.0001$ ). The steroid hormone, progesterone is critical in maintenance of pregnancy [25] and luteal phase insufficiency with low progesterone production is an established endocrine cause of RPL [6,26]. Other risk factors are advanced maternal age ( $>35$  yrs) and obesity [9]. In this present study of non-obese women (aged  $<35$  yrs with cases age-matched with controls), our findings confirm earlier reports [6,9,26] and suggest that TG, NPP, NLB and progesterone may be independent of obesity and age as risk factors of RPL.

**Table 1. Comparison of Demographic, Anthropometric, Lipids and Progesterone between Women with Recurrent Abortion (cases) and controls**

Parameters	Cases n = 60	Controls n = 30	p-value
Demographic Parameters			
Age (years)	34.37±0.55	33.97±4.31	0.750
AM (years)	13.92±0.19	17.23±3.34	0.165
LMC (days)	28.45±0.40	28.77±1.48	0.596
NLB	01.43±0.12	02.67±0.14	0.000**
NPP	04.67±0.13	02.67±0.13	0.000**
Anthropometric Parameters			
Height (m)	01.70±0.02	01.68±0.02	0.542
Weight (kg)	64.22±0.66	64.70±0.71	0.650
BMI (kg/m <sup>2</sup> )	22.74±0.43	22.87±0.54	0.860
WC (cm)	16.93±0.26	16.93±0.31	1.000
HC (cm)	47.80±0.47	46.73±0.50	0.163
WHR	0.353±0.003	0.359±0.004	0.219
Lipids			
TC (mg/dl)	139.15±5.00	153.50±3.80	0.061
TG (mg/dl)	93.15±3.41	90.67±5.38	0.025*
HDL (mg/dl)	45.30±0.64	46.40±0.89	0.318
LDL (mg/dl)	75.55±4.94	88.97±4.28	0.082
Progesterone (nmol/L)	04.84±0.61	18.35±3.09	0.000**

\*= significant, n=no. of subjects, T-test=student's t-test, p = probability, cases= Non-pregnant women with history of recurrent abortion, controls= Non-pregnant women without history of recurrent abortion, BMI= body mass index, WHR= waist hip ratio, LMC=length of menstrual cycle, AM= age at menarche, NLB= number of live births, NPP= number of previous pregnancy HC= hip circumference, WC= waist circumference, TC=total cholesterol, TG=triglyceride, HDL=high density lipoprotein, LDL=low density lipoprotein

Cholesterol is the precursor of all steroid hormones and impairment of lipid metabolism has long been implicated in the luteal phase deficiency of progesterone resulting in RPL [27]. It is hypothesised that cholesterol insufficiency impedes the initiation of sex hormones resulting in foetal loss [28]. Our data showed no significant differences in cholesterol containing lipid fractions-TC, LDL and HDL ( $p>0.318$ ) between cases and controls. However, the observed higher TG levels in the cases compared with controls in this study, contrary to earlier findings of Oladimeji et al. [29] in our geographical sub-region, implicates current lifestyle factors. Metabolic syndrome characterised by increased levels of triglycerides, abdominal obesity and insulin resistance is now prevalent in Nigeria [30]. This has been attributed to changes by Nigerians to western diet and lifestyle [31,32] and might contribute to RPL. Obesity and/or insulin resistance have been associated with an increased risk of pregnancy loss [26].

### 3.2 Micronutrients and Heavy Metals

Table 2 shows the comparison of micronutrients and heavy metals between women with recurrent abortion (cases) and controls. Cd and Pb were significantly higher in cases compared with controls ( $p<0.013$ ). Cd and Pb are known to accumulate in fish [13]. Consumption of fish as well as continuous and prolonged exposure to Cd and Pb have been linked to adverse pregnancy outcomes [10]. The overall effect of toxic metals on any cell or

tissue has been attributed to a synergy of several mechanisms with one mechanism predominating in a specific cell type [12]. Chemical pollutants have been shown to interact additively to diet and lifestyle to modulate endocrine function. As endocrine disruptors, Cd and Pb target the homeostasis of sex steroids through multiple diverse pathways [10,12].

**Table 2. Comparison of micronutrients and heavy metals between women with recurrent abortion (cases) and controls**

<b>Micronutrients</b>	<b>Cases n= 60</b>	<b>Controls n= 30</b>	<b>p- value</b>
Zinc (µmol/L)	13.45±0.80	12.24±0.43	0.308
Selenium (µmol/L)	08.46±0.21	11.95±0.36	0.000**
Manganese(µmol/L)	09.49±0.18	11.04±0.36	0.000**
Chromium (µmol/L)	11.14±0.21	12.21±0.41	0.012**
Magnesium (µmol/L)	00.75±0.04	01.04±0.03	0.000**
Iron (µmol/L)	11.42±0.67	18.36±0.77	0.000**
Vitamin E (mg/dl)	03.48±0.24	06.89±0.30	0.000**
Copper (µmol/L)	11.71±0.44	15.39±0.54	0.000**
<b>Toxic trace metals</b>			
Cadmium(µmol/L)	03.58±0.17	02.45±0.13	0.000**
Lead(µmol/L)	02.93±0.08	02.05±0.33	0.001**

\* = significant, n = number of subjects, p = probability, Cases= Non-pregnant women with history of recurrent abortion, Controls= Non pregnant women without recurrent abortion, T- test= student's t- test.

Oxidative stress has been implicated as one of the main underlying mechanisms in the pathogenesis of spontaneous abortion [2,4].

Increased OS seems a common end-point to several endocrine disruptors [10]. Loss of antioxidant defences, high levels of biochemical markers of ROS-induced membrane damage such as lipid peroxidation products and oxidant/antioxidant imbalance are associated with pregnancy loss. Oxidative damage to macromolecules and DNA and ROS-induced signal transduction for various genes might result in recurrent pregnancy loss [2,4,12].

Female OS is postulated as a likely mediator of conception and early pregnancy loss might be placenta related [1,2]. The placenta is exposed to high oxygen metabolism [2,4]. Placental OS, with associated necrosis and apoptosis of the trophoblastic epithelium of the placental villous tree has been suggested. Development of the placento–decidual interface is severely impaired leading to early and widespread onset of maternal blood flow and major oxidative degeneration, which influences the timing of early pregnancy loss [1].

The mechanism of metal-induced formation of free radicals is tightly influenced by the action of cellular antioxidants [11]. The antioxidant status is known to be a physiological barrier against free radical attack in certain body compartments. Retardation of trophoblastic outgrowth and development, placental necrosis and suppression of steroid biosynthesis, and altered handling of nutrient metals by the placenta all contribute to implantation delay and possible early pregnancy loss [12]. Placental oxidative stress ultimately depletes antioxidant defences [2,4]. We observed significantly lower levels of micronutrients (Fe, Cu, Mn, Mg, Cr, Se, Vit.E) in cases compared with controls (p<0.013) Table 2. Iron deficiency contributes to one of the largest prevalence of micronutrient deficiencies among pregnant women [33]. Nutrients interact with endocrine disruptors to mitigate their effects. Vitamin E with

antioxidant activities appear to moderate the effect of endocrine disruptors and other toxicants [10]. The ability of Cd and Pb to imitate nutritive metals that are divalent cations-Fe, Zn, Mg and Mn in many biological pathways and processes as well as interact with membrane transporters involved in their uptake, is a possible explanation for their toxicity. Divalent nutrients have demonstrably protected embryonic and adult tissue from toxic metal induced damage. Se probably blocks Cd uptake by embryonic cells [12]. Zn, an essential component of numerous proteins involved in the defence against oxidative stress, has an impact on the immune system and possesses neuroprotective properties. Its depletion is known to impair DNA repair mechanisms and enhance DNA damage [11]. The similar Zn levels in both cases in our study (Table 2), suggest synergistic action of multinutrients to prevent adverse pregnancy outcomes [33].

### **3.3 Relationship of Lipids, Demographic and Anthropometric Characteristics with Progesterone**

Rapid non-genomic events might critically determine the functional response of progesterone in the female reproductive system in a cell-type- and environment-specific manner [25]. We observed significantly positive correlations of only age and weight with progesterone in cases only ( $p < 0.026$ ). No correlation was observed between all parameters tested and progesterone in controls ( $p > 0.05$ ). Our observations suggest that increasing age and weight may predict progesterone levels in non-obese women with RPL aged below 35 years. The lack of correlation of progesterone with Cd and Pb, micronutrients and any of the lipid fractions suggest that their importance in RPL may be due to factors other than progesterone. Cadmium has recently been recognized as an estrogenic endocrine disruptor. Cd and Pb might target sex steroids other than progesterone, possibly acting on the pathways mediated by oestrogen receptors [10]. We did not estimate levels of other hormones like oestrogen, implicated in RPL, which is the limitation of this study.

Pregnancy represents a state of increased metabolic requirements and micronutrient malnutrition among women of reproductive age and pregnant women is much more common in developing countries than in industrialized countries [33,34]. Deficient micronutrients in the non-pregnant state and/or increased micronutrient requirements during repeated pregnancies of our cases might have further exacerbated the pre-existing maternal deficiency [33]. Recommendations for prenatal micronutrient supplements are absent in most developing countries, contrary to some developed countries where such supplements are routinely recommended [34]. Micronutrient supplementation in pregnant women particularly in developing countries is controversial [33,34]. Our study show the impact of the environment in addition to diet and lifestyle and progesterone in women with RPL. However, these factors may be independent causes of RPL particularly in normal weight women of less than 35 yrs with RPL. Healthcare strategies show focus on reduction of environmental pollution and increase awareness on healthy diet for optimal periconceptional micronutrient requirements [10,17]. Tightly monitored clinical trials on the safety, efficacy and effective delivery of maternal micronutrient supplementation in women with RPL [34] before and during pregnancy are recommended.

## **4. CONCLUSION**

We observed in our study that reduced progesterone levels, number of previous pregnancies, number of live births, levels of micronutrients (Iron, Copper, Manganese, Magnesium, Chromium, Selenium, Vitamin E) and increased levels of triglycerides, toxic

metals (cadmium, lead) are independent risk factors for recurrent pregnancy loss in normal weight women below 35 years. Of all factors, only age and weight correlated with progesterone. Our findings implicate oxidative stress mechanisms. Inadequate micronutrients in our cases might be further exacerbated in the next pregnancy due to increased micronutrient demands thus forming a vicious cycle of repeated/recurrent pregnancies losses. Healthcare strategies and research should therefore focus on reduction of environmental pollution and increase awareness on healthy diet for optimal micronutrient requirements in women with recurrent pregnancy loss.

## **CONSENT**

We declare that informed consent was obtained from all the participants in the study after detailed explanation of the study.

## **ETHICAL APPROVAL**

We hereby declare that the study protocol was examined and approved by the University of Ibadan/University College Hospital, Ibadan ethics committee and have therefore been performed in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki.

## **ACKNOWLEDGEMENTS**

This research was self funded.

## **COMPETING INTERESTS**

Authors declare that no competing interests exist.

## **REFERENCES**

1. Jauniaux E, Poston L, Burton GJ. Placental-related diseases of pregnancy: involvement of oxidative stress and implications in human evolution. *Hum Reprod Update.* 2006;12:747-755.
2. Ruder EH, Hartman TJ, Blumberg J, Goldman MB. Oxidative stress and antioxidants: exposure and impact on female fertility. *Hum Reprod Update.* 2008;14:345-357.
3. Caserta D, Maranghi L, Mantovani A, Marci R, Maranghi F, Moscarini M. Impact of endocrine disruptor chemicals in gynaecology. *Hum Reprod Update.* 2008;14:59-72.
4. Gupta S, Agarwal A, Banerjee J, Alvarez JG. The Role of Oxidative Stress in Spontaneous Abortion and Recurrent Pregnancy Loss: A Systematic Review. *Obstetrical and Gynecological Survey.* 2007;62(5):335-347.
5. Lashen H, Fear K, Sturdee DW. Obesity is associated with increased risk of first trimester and recurrent miscarriage: matched case-control study. *Hum Reprod.* 2004;19:1644-6.
6. Romero R. Prevention of spontaneous preterm birth: the role of sonographic cervical length in identifying patients who may benefit from progesterone treatment. *Ultrasound Obstet Gynecol.* 2007;30:675-686.



7. Metwally M, Ong KJ, Ledger WL, Li TC. Does high body mass index increase the risk of miscarriage after spontaneous and assisted conception? A meta-analysis of the evidence. *Fertil Steril.* 2008;90:714-26.
8. Metwally M, Saravelos SH, Ledger WL, Li TC. Body mass index and risk of miscarriage in women with recurrent miscarriage. *Fertil Steril.* 2010;94:290-5.
9. Kroon B, Harrison K, Martin N, Wong B, Yazdani A. Miscarriage karyotype and its relationship with maternal body mass index, age, and mode of conception. *Fertil Steril.* 2011;95:1827-1829.
10. Caserta D, Mantovani A, Marci R, Fazi A, Ciardo F, La Rocca C, Maranghi F, Moscarini M. Environment and women's reproductive health. *Hum Reprod Update.* 2011;17:418-433.
11. Jomova K, Valko M. Advances in metal-induced oxidative stress and human disease. *Toxicology.* 2011;283:65–87.
12. Thompson J, Bannigan J. Cadmium: Toxic effects on the reproductive system and the embryo. *Reproductive Toxicology.* 2008;25:304–315.
13. Farombi EO, Adelowo OA, Ajimoko YR. Biomarkers of oxidative stress and heavy metal levels as indicators of environmental pollution in African cat fish (*Clarias gariepinus*) from Nigeria Ogun River. *Int J Environ Res Public Health.* 2007;4:158-65.
14. Gerhard I, Waibel S, Daniel V, Runnebaum B. Impact of heavy metals on hormonal and immunological factors in women with repeated miscarriages. *Hum. Reprod Update.* 1998;4:301-309.
15. Tremellen K. Oxidative stress and male infertility—a clinical perspective *Hum. Reprod Update.* 2008;14:243-258.
16. Vural P, Akgu C, Yildirim A, Canbaza M. Antioxidant defence in recurrent abortion. *Clinica Chimica Acta.* 2000;295:169–177.
17. Cetin I, Berti C, Calabrese S. Role of micronutrients in the periconceptual period. *Hum Reprod Update.* 2010;16:80-95.
18. World Health Organization. Obesity: preventing and managing the global epidemic. Report of a WHO consultation on obesity. Geneva: WHO; 1998
19. Umoh U, Charles-Davies M A, Adeleye J. Serum testosterone and lipids in relation to sexual dysfunction in males with metabolic syndrome and type2 diabetes mellitus. *International Journal of Medicine and Medical Sciences.* 2010;2:402-412.
20. Pugeat MM, Dum JF, Nisula BC. Transport of steroid Hormones interactions of 70 Drugs with testosterone binding globulin and corticosteroid binding globulin in human plasma. *J. Chn and metals.* 1981;53:69-75.
21. Alain CC, Poon LS, Chan CSG, Richmand W, Fu PC. Enzymatic detemination of total serum cholesterol. *Clin. Chem.* 1974;20:470.
22. Trinder P. Determination of glucose in blood using glucose oxidase with an alternative oxygen acceptor. *Ann Clin Biochem.* 1969;6:24–27.
23. Friedwald WT, Levy RI, Fredrickson, D.S. Estimation of the concentration of low density lipoprotein cholesterol in plasma, without use of the preparative ultracentrifuge. *Clin Chem.* 1972;18:499-502.
24. Kaneko JJ. *Clinical biochemistry of animals*, Fourth ed. New York:Academic Press Inc; 1999.
25. Gellersen B, Fernandes MS, Brosens JJ. Non-genomic progesterone actions in female reproduction. *Hum Reprod Update.* 2009;15:119-138.
26. Dhont M. Recurrent miscarriage. *Current Women's Health Reports.* 2003;3:361–366.
27. Coste J, Jabspera N, Fernandez H. Risk factors for spontaneous abortion: a case control in France. *Hum Reprod.* 1991;6:1332-1337.

28. Shekeres-Bartho J, Shekeres GY, Debre P, Autra B, Chaouat G. Reactivity of lymphocyte to a progesterone receptor-specific monoclonal antibody. *Cell Immunol.* 1990;125:273-281.
29. Oladimeji OS, Magbagbeola OA, Peter SS, Adewole TA, Akinwande IA. Serum Vitamin E, Cholesterol and Triglyceride Levels of Nigerian Women with Unexplained Infertility and Recurrent Miscarriage *Nig Jnl Health & Biomedical Sciences.* 2002;1:71-75.
30. Charles-Davies MA, Arinola OG, Fasanmade AA, Olaniyi JA, Oyewole OE, Owolabi MO, et al. Indices of Metabolic Syndrome in 534 Traders. *Journal of the US-China Medical Science.* 2012;9:91-100.
31. Utinwa G.O. Prevalence of obesity and Predisposition to metabolic syndrome amongst school based adolescents in Botswana and Nigeria. *The African Symposium-an online Journal of African Education Research Network.* 2009;9:48–52.
32. Adegoke OA, Adedoyin RA, Balogun MO, Adebayo RA, Bisiriyu LA, Salawu AA. Prevalence of Metabolic Syndrome in a Rural Community in Nigeria. *Metabolic Syndrome and Related Disorders.* 2010;8:59–62.
33. Haider BA, Mohammad YY, Bhutta ZA. Effect of multiple micronutrient supplementation during pregnancy on maternal and birth outcomes. *BMC Public Health.* 2011;11(Suppl 3):S19.
34. Kawai K, Spiegelman D, Shankar AH, Fawzi WW. Maternal multiple micronutrient supplementation and pregnancy outcomes in developing countries:meta analysis and meta-regression. *Bull World Health Organ.* 2011;89:402–411B.

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