

CHLAMYDIAL INFECTION, PLASMA PEROXIDATION AND OBESITY IN TUBAL INFERTILITY

A.C. Nsonwu-Anyanwu¹, M.A.Charles-Davies¹, A.A.Oni², V.O. Taiwo³ and F.A. Bello⁴

¹Department of Chemical Pathology, University College Hospital, Ibadan

²Department of Medical Microbiology & Parasitology, University College Hospital, Ibadan

³Department of Veterinary Pathology, University College Hospital, Ibadan

⁴Department of Obstetrics and Gynaecology, University of Ibadan, Nigeria

Correspondence:

Dr. Folasade A. Bello

Department of Obstetrics and Gynaecology

University of Ibadan

Ibadan, Nigeria

E-mail: dr.nikebello@yahoo.com

Tel: +2348037084505

ABSTRACT

Background: Genital tract infections and obesity are both sources of oxidative stress. Alterations in immune and antioxidant parameters may arise from this or from an indeterminate autoimmune mechanism.

Objective: This study aimed to investigate the association of Chlamydial infection, obesity and oxidative response with tubal infertility in Nigerian women.

Methods: It was a case-control study of 40 women with tubal infertility and 32 fertile women, respectively, recruited from the Infertility and Family Planning Clinics respectively, of the University College Hospital, Ibadan, Nigeria. Anthropometric indices were measured in each subject and endocervical swabs were taken to screen for current genital tract infection. Antioxidant, hormonal and immunologic analysis were performed on serum.

Results: None of the subjects had current genital tract infections. Chlamydia trachomatis IgG positivity was significantly higher in infertile than in fertile subjects [OR 4.33; 95%CI (0.078-0.681)]. No significant variations were observed in the anthropometric indices, antioxidant parameters and hormones between infertile and the fertile women. Body mass index correlated positively with oxidative stress in infertile subjects. Waist and hip circumferences correlated negatively with oestradiol in women with tubal infertility.

Conclusion: Chlamydial infection is associated with tubal factor infertility, however, obesity seems to increase oxidative stress and reduce fertility potential in women with tubal factor infertility.

Key words: Tubal infertility, obesity, oxidative stress, Chlamydia

INTRODUCTION

Human reproductive failure is as old as mankind. This public health problem involves all regions of the world and its prevalence worldwide varies from 10-15%¹; 10-32% in Africa² and 31.5% in Nigeria³. The various categories of human reproductive failure include: infertility, recurrent pregnancy loss (spontaneous miscarriages or abortions and preterm birth) and ectopic pregnancy. Causes include environmental and lifestyle factors, congenital malformations, endocrine disorders, immunologic abnormalities and sequelae of genital tract infections^{4,5}.

The association of infertility with genital tract infections (GTIs) or sexually transmitted infections has been demonstrated. GTIs associated with human

reproductive failure include: *Treponema pallidum*, *Neisseria gonorrhoeae*, *Chlamydia trachomatis*, *Trichomonas vaginalis* and *Schistosoma haematobium* infections among others. The spread of gonococcal and Chlamydial infection to upper female genital tract may cause pelvic inflammatory disease with severe tubal scarring leading to tubal infertility⁶. Genital tract infections evoke both cellular and humoral immune response leading to activation of polymorphonuclear leucocytes, macrophages and the release of cytokines. These immunologic factors produced in response to GTIs influence various aspects of reproduction including follicle development, ovulation, luteinisation, oocyte quality, fertilization, implantation, foetal development and pregnancy immunotolerance⁷⁻⁹.

Abnormal immunologic profile which in part is characterized by autoantibody abnormalities have been described in infertile women¹⁰. GTIs induce increased production of reactive oxygen species and nitric oxide leading to oxidative stress which has been implicated in the pathophysiology of tubal and peritoneal factor infertility, endometriosis, preeclampsia, hydatidiform mole, free radical-induced birth defects and abortions¹¹⁻¹⁶. Their pathological effects are exerted by various mechanisms including lipid peroxidation, peroxidative DNA damage, mitochondrial alterations and apoptosis¹⁷. The complex interplay between GTI and its induction of oxidative stress and pathologic immune response both systemic and locally and the deleterious effects of these responses on the male and female genital tract which include tissue damage, tubal scarring, pelvic inflammatory disease, chronic salpingitis, endometritis and distal tubal obstruction have been implicated as pathologic mechanisms of infertility.

Excessive weight and central distribution of body fat have been related to an increased risk of normogonadotrophic anovulation¹⁸. High body mass index (BMI) has been implicated in development of insulin resistance and hyperinsulinaemia. Elevated insulin levels have been reported to inhibit the conversion of testosterone to oestrogen. Lower levels of oestrogen will lead to anovulation and infertility¹⁹. Obesity appears to have a major impact on reproductive performance and can compromise it in a variety of ways; including menstrual disorders and anovulation. Weight reduction in obese patients reduces hyperandrogenism and hyperinsulinaemia; both of which factors influence the ovarian response to follicular stimulating hormone. Weight loss can re-establish ovulation in obese anovulatory patients or improve their response to ovulation induction. Clark *et al.*²⁰ in 1995 found that weight loss re-established ovulation in obese anovulatory patients or improved their response to ovulation induction: in a series of 67 anovulatory women, 90% resumed ovulation after weight loss and 78% conceived. The same study group confirmed these findings in a larger series²¹. Similar results were obtained in preliminary observations by Crosignani *et al.*, 1999²² and 2002²³. In another series, among the 27 out of 33 patients with irregular menstrual cycles who lost weight, 18 re-established regular cycles¹⁸. A total of 60% had ovulatory levels of plasma progesterone after weight loss. Neither menstrual cycle improvement, nor ovulatory values of progesterone nor pregnancies occurred in the eight patients who did not lose weight. Therefore, it is presumed that obesity may be responsible for the relative ovarian insensitivity to infertility treatment¹⁸.

This study aimed to investigate the association of Chlamydial infection, obesity and oxidative response with tubal infertility in Nigerian women.

MATERIALS AND METHODS

Study design

This case control study was conducted in the University College Hospital (UCH), Ibadan, Nigeria. It served as a pilot for an on-going larger work studying the association of oxidative stress and pathologic response to stress with infertility. The study population comprised of female patients of reproductive age attending the Infertility Clinic and a control group of age-matched fertile women who were new clients at the Family Planning Clinic. The study protocol was approved by UI/UCH Ethical Committee (Ref UI/EC/08/0083).

Hypothesis

The hypothesis was that tubal infertility is not associated with oxidative stress and obesity. The outcome measures were the anthropometric and oxidative stress parameters of the subjects.

Selection of Subjects

Forty consenting women with infertility of at least one year's duration (with tubal blockage identified by hysterosalpingography) were recruited from the infertility clinic, while 32 controls were recruited from the family planning clinic. The controls were women without previous infertility who had childbirth within the last two years, and who had not been on any form of contraception prior to recruitment. Exclusion criteria included women that were undergoing any form of contraceptive therapy, previous history of uterine surgery, malignancy, long term medication, chronic organ or systemic illness and those who did not give consent.

Anthropometric indices (height, weight, waist and hip circumferences) were taken to calculate the body mass index and waist-hip ratio (WHR) respectively.

Sample Collection

Ten millilitres of venous blood samples were collected aseptically from each subject on days 3-5 and 21-23 of a 28-30 day menstrual cycle, respectively. The samples were dispensed into universal containers. After clot retraction, the samples were centrifuged at 3000 rev/s for ten minutes after which serum was extracted and stored in small aliquots at -20°C. High vaginal swabs (HVS) and endocervical swabs (ECS) were also taken from all subjects of study using sterile swab sticks for detection of common sexually-transmitted infections.

Laboratory Methods

Endocrinological analysis: follicular stimulating hormone (FSH), luteinizing hormone (LH), progesterone (P4), estradiol (E2) and prolactin (PRL) were measured with enzyme immunoassay method (EIA) from Immunometrics Ltd., London, UK.

Antioxidant profile: total antioxidant potential (TAP) was estimated using the ferric reducing antioxidant power (FRAP) method of Benzie and Strain²⁴. Total plasma peroxides (TPP), as a biomarker of Oxidative Stress (OS), were estimated using the modified FOX 2 method of Harma *et al.*²⁵. Oxidative stress index (OSI) was calculated as a ratio of TPP/TAP.

Microbiological Analysis: screening for *Trichomonas vaginalis* was by microscopy while *Neisseria gonorrhoeae* was by gram staining, followed by culture if necessary. Immunologic analysis: endocervical swabs were tested for *Chlamydia trachomatis* antigen with a rapid screening test (DiaSpot™, Bresta Perkasa, Indonesia). Serum was screened for *Treponema pallidum* antibodies (IgG & IgM) by immunochromatographic method with Exact® syphilis diagnostic device, USA; and *Chlamydia trachomatis* antibodies (CT IgG) with ImmunoComb®, Organics Ltd., Yavne, Israel.

Statistical Analysis

Data was analyzed using the statistical package of social science (SPSS) software 15.0 version. For quantitative variables, paired student's t-test was used to test for mean differences. Pearson's correlation analysis was employed to determine associations between variables. For non-quantitative variables, χ^2 -square analysis was used for determination of associations between variables. Significant *p* was <0.05.

RESULTS

The screening tests for the afore-mentioned were all negative, suggesting that none of the patients had current genital tract infection. Chlamydial antibodies were further tested for; to identify women with previous infection which may have led to the tubal damage. Table 1 shows the comparison of Chlamydia

	Infertile (n=40)	Fertile (n=32)	Total
<i>CTIgG +ve</i>			
Subjects	20	6	26
%	50.0	18.3	
<i>CTIgG -ve</i>			
Subjects	20	26	46
%	50.0	81.7	
	40	32	72

Table 1: Comparison of Chlamydia trachomatis IgG (CTIgG) antibody positivity in Fertile and Infertile Women

OR=4.33, 95%CI (0.078-0.681)
²=7.847, df=1, P=0.005

CT IgG= Chlamydia trachomatis immunoglobulin G
 +ve = Positive
 -ve = Negative

trachomatis IgG (CT IgG) antibody positivity in fertile and infertile women. The fertile group of the study population were significantly less likely to have Chlamydia trachomatis antibodies than the infertile group [OR=4.33; 95%CI (0.078-0.681)].

Table 2 shows the comparison of anthropometric data, hormone profiles and antioxidant parameters among fertile and infertile women; luteinizing hormone was significantly higher in the infertile group.

The correlation of anthropometric indices with oxidative stress and hormonal parameters showed that BMI had a significant positive correlation with TPP (*p*<0.001) and OSI (*p*<0.001) in the infertile group but not in controls. In infertile women, waist circumference (*p*<0.006) and hip circumference (*p*<0.001), respectively, correlated negatively with oestrogen.

DISCUSSION

The total antioxidant potential (TAP), total plasma peroxides (TPP) and therefore, oxidative stress index (OSI) were not significantly different between infertile women and their fertile controls. This finding is in agreement with other authors, who reported no significant differences in TAP in infertile and fertile controls^{26,27}. Total plasma peroxides were positively associated with BMI; this indicated that increasing BMI may be a risk factor for increased oxidative stress and the associated deleterious effects on the general body system. Oxidative stress has been implicated in the pathogenesis of more than 100 disease conditions including infertility¹⁶. The place of obesity in anovulatory infertility has been discussed—however, the subjects of the current study were ovulating, as observed by their mid-luteal progesterone assays. This finding suggests that BMI may increase oxidative stress and its influence on infertility in the absence of anovulation.

The negative correlation observed between oestradiol and waist and hip circumferences imply that obesity in women may lead to a hormonal imbalance that may reduce their fertility. It has been reported that ovulating subfertile women with a BMI over 29 kg/m² have lower pregnancy rates compared with those with normal weight²⁸. Our study also points to this fact as indicated by the increased oxidative stress.

Index	Infertile	Fertile	t	p
	n = 40	n = 32		
Age (yrs)	32.3±4.0	33.5±4.8	-1.185	0.240
Weight (kg)	69.2±11.4	66.1±14.4	1.019	0.312
Height (m)	1.6 ± 0.1	1.6 ± 0.1	0.423	0.673
BMI (kg/m ²)	27.1±0.4	26.1±5.1	0.879	0.382
WC (cm)	84.9±8.3	84.4±12.4	0.179	0.858
HC (cm)	103.7±9.6	103.0±12.4	0.272	0.786
WHR	0.82±0.0	0.82±0.1	-0.027	0.979
TAP (μmol TE/L)	767.9±146.8	748.9±90.2	0.643	0.522
TPP (μmol H ₂ O ₂ /L)	6.3±7.8	4.4±1.4	1.337	0.185
OSI	0.9±0.9	0.6±0.2	1.569	0.121
FSH (IU/L)	13.4±11.8	10.1±4.7	1.462	0.148
LH (IU/L)	16.1±21.2	5.7±3.9	2.712	0.008
Prolactin (IU/L)	452.2±486.2	611±874.9	-0.976	0.332
P4 (p mol/L)*	22.6±19.8	-	3.264	-
E2 (p mol/L)	0.85±1.87	0.51±0.41	1.000	0.321

*see text for explanation'

BMI- body mass index, WC-waist circumference, HC-hip circumference, WHR- waist-hip ratio, TAP- total antioxidant potential, TPP- total plasma peroxides, OSI- oxidative stress index, FSH-follicular-stimulating hormone, LH-luteinizing hormone, P4-progesterone, E2-oestrogen

Table 2: Comparison of anthropometric data, hormone profiles and antioxidant parameters in Fertile and Infertile Women

We did not expect the fertility hormone profile to be significantly different between infertile and fertile women in this study; the profile was studied to ascertain that the cases did not have anovulatory infertility, which could be a confounder. The only isolated significant hormone was luteinizing hormone. The current study cannot fully explain the association; however, the mid-luteal progesterone suggested these women were

actually ovulating—therefore, the elevated LH may be of little consequence. The study design did not allow for testing mid-luteal progesterone in the controls. We required these fertile controls to be contraceptive- and hormone-naïve, so they were recruited at their first visit to family-planning clinic. A mid-luteal progesterone assay would have required a follow-up visit, but by this time, they would have been

commenced on a contraceptive method, so would not be suitable for inclusion any longer. The relatively high values of prolactin in the fertile women were probably because most of them were still nursing their infants at the time of recruitment. The prevalence of Chlamydia antibody positivity was found to be significantly higher in infertile women compared to their fertile controls. This finding is in agreement with the findings of other workers^{3,29-34}. Chlamydial infection evokes pathologic immune response and generation of reactive oxygen species which results in inflammatory response, apoptosis, tissue damage, scarring, fibrosis, hydrosalpinx, tubal occlusion leading to tubal infertility¹⁶.

The findings of this study may be limited by the small sample size; the association of obesity with tubal infertility may be validated in the larger study to follow. The study will include evaluation of cellular as well as humoral pathologic response found associated with tubal infertility, and will hopefully shed more light on the aetiopathogenesis of infertility.

CONCLUSION

Previous Chlamydial infection is significantly associated with tubal factor infertility. Obesity seems to reduce fertility potential in women with tubal factor infertility as well as increase oxidative stress. Larger studies are required to explore this.

REFERENCES

- World Health Organization. Recent Advances in medically assisted conception. Report of a WHO scientific group. Technical report series, No. 820. Geneva, World Health Organization 1992.
- Gerias AS**, Rushman H. Infertility in Africa. *Popul Sci* 1992; 12: 26-46.
- Sule JO**, Erigbali P, Eruom L. Prevalence of infertility in women in a southwestern Nigerian community. *Afr J Biomed Res* 2008; 11: 225-227.
- Radojic L**, Marjanovic S, Vicovac L, Kataranovski M. Anticardiolipin antibodies in women with unexplained infertility. *Physiol Res* 2004; 53: 91-96.
- Trounson A**. Development in infertility therapy-diagnosis of genetic disease in embryos. *Aust Fam Physician* 2005; 34(3): 123-125.
- Sparling PF**. Biology of *Neisseria gonorrhoeae*. In: Holmes KK, Mardh PA, Sparling PF *et al* (eds). Sexually transmitted diseases (3rd ed). New York, McGraw-Hill 1999: 433-449.
- Gleicher N**. Some thoughts on the reproductive autoimmune failure syndrome (RAFS) and Th-1 versus Th-2 immune responses. *Am J Reprod Immunol* 2002; 48: 252-254.
- Robertson SA**. Control of the immunological environment of the uterus. *Rev Reprod* 2000; 5: 164-174.
- Clark DA**. T cell in pregnancy: illusion and reality. *Am J Reprod Immunol* 1999; 41: 233-238.
- Vujisic S**, Zidovec S. Follicular immunology environment and the influence on In-vitro fertilization outcome. *Curr Womens Health Rev* 2005; 1: 49-80.
- Dong M**, Shi Y, Cheng Q, Hao M. Increased nitric oxide in peritoneal fluid from women with idiopathic infertility and endometriosis. *J Reprod Med* 2001, 46:887-891.
- Polak G**, Koziol-Montewka M, Gogacz M *et al*. Total antioxidant status of peritoneal fluid in infertile women. *Eur J Obstet Gynecol Reprod Biol* 2001, 94: 261-263.
- Van Langendonck A**, Casanas-Roux F, Donnez J. Oxidative stress and peritoneal endometriosis. *Fertil Steril* 2002; 77: 861-870.
- Harma M, Kocyigit A, Demir N. Role of plasma nitric oxide in complete hydatidiform mole. *Eur J Gynaecol Oncol* 2004, 25: 333-335.
- Loeken MR**. Free radicals and birth defects. *J Matern Fetal Neonatal Med* 2004, 15: 6-14.
- Agarwal A**, Gupta S, Sharma KR. Role of oxidative stress in female reproduction. *Rep Biol Endocrinol* 2005; 3: 28-49.
- Agarwal A**, Saleh RA, Bedaiwy MA. Role of reactive oxygen species in the pathophysiology of human reproduction. *Fertil Steril* 2003; 79: 829-843.
- Al-Hasani S**, Zohni K. The Overlooked Role of Obesity in Infertility. *J Fam Reprod Health* 2008; 2: 115-122.
- Sudha G**, Reddy KSN, Reddy KK. Association between body mass index and infertility: A cross sectional study. *Asia Pac J Soc Sci* 2009; 1: 73-81.
- Clark AM**, Ledger W, Galletly C *et al*. Weight loss results in significant improvement in pregnancy and ovulation rates in anovulatory obese women. *Hum Reprod* 1995; 10: 2705-2712.
- Clark AM**, Thornley B, Tomlinson L *et al*. Weight loss in obese infertile women results in improvement in reproductive outcome for all forms of fertility treatment. *Hum Reprod* 1999; 13: 1502-1505.
- Crosignani PG**, Piloni S, Gessati A *et al*. Resumption of fertility with diet in PCOS patients. *ASRM/CFAS Conjoint Annual Meeting*, September 25-30; Toronto, Ontario, Canada, *Fertil Steril* 1999; Suppl 72: S233.
- Crosignani PG**, Vegetti W, Colombo M, Ragni G. Resumption of fertility with diet in overweight women. *Reprod Biomed Online* 2002; 5: 60-64.

24. **Benzie IFF**, Strain JJ. The ferric reducing ability of plasma (FRAP) as a measure of "antioxidant power": the FRAP assay. *Anal Biochem* 1996; 239: 70-76.
25. **Harma M**, Harma M, Enel O. Increased oxidative stress in patients with hydatidiform mole. *Swiss Med Wkly* 2003; 133: 563-566.
26. **Veena BS**, Sharmila U, Satish KA, Pratap KN. Evaluation of oxidative stress, antioxidants and prolactin in infertile women. *Indian J Clin Biochem* 2008; 23(2): 186-190.
27. **Agarwal A**, Allamaneni SS. 2004 Role of free radicals in female reproductive diseases and assisted reproduction. *Reprod Biomed Online* 2004; 9: 338-347.
28. **van der Steeg JW**, Steures P, Eijkemans MJ *et al.* Obesity affects spontaneous pregnancy chances in subfertile, ovulatory women. *Hum Reprod* 2008; 23(2): 324-328.
29. **Otolorin EO**, Ojengbede O, Falase OA. Laparoscopic evaluation of the tuboperitoneal factor in infertile Nigerian women. *Int J Gynaecol Obstet* 1987; 25: 47-52.
30. **Ekwere PD**, Archibong EI, Bassey EE *et al.* Infertility among Nigerian couples as seen in Calabar. *Port Harcourt Med J* 2007; 2: 35-40.
31. **Olatunji AO**, Sule-Odu AO. The pattern of infertility cases at a University Hospital. *West Afr J Med* 2003; 22: 205-207.
32. **Omo-Aghoja LO**, Okonofua FE, Onemu SO *et al.* Association of Chlamydia trachomatis serology with tubal infertility in Nigerian women. *J Obstet Gynaecol Res* 2007; 33: 688-695.
33. **Ikechebelu I.I**, Adinma I.I, Ori E.F, Ikegwuonu S.O. High prevalence of male infertility in Southeastern Nigeria. *J Obstet Gynaecol* 2003; 23: 657-659.
34. **Okonofua F**, Menakaya U, Onemu SO *et al.* A case study of risk factors for male infertility in Nigeria. *Asian J Androl* 2005; 7: 351-361.