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Original Research Article

Ten-year review of the clinical presentation and treatment outcome of Asherman's syndrome in university of Maiduguri teaching hospital, Borno State, Nigeria

Ado Danazumi Geidam^{1*}, Adamu Malgwi²

¹Department of Obstetrics and Gynaecology, College of Medical Sciences, University of Maiduguri, Borno State, Nigeria,

²Department of Obstetrics and Gynaecology, University of Maiduguri Teaching Hospital, Maiduguri, Borno State, Nigeria

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***Correspondence:**

Dr. Ado Danazumi Geidam,

E-mail: adogeidam@yahoo.com

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ABSTRACT

Background: Asherman's syndrome (AS) is an important gynaecological disorder characterized by a menstrual abnormality (hypomenorrhea/amenorrhea) and infertility that is usually caused by activities that traumatize the endometrium. Objective of current study was to determine the prevalence, clinical presentation and treatment outcome of Asherman's syndrome in our hospital.

Methods: A retrospective review of patients with Asherman's syndrome managed at the University of Maiduguri Teaching hospital over 10 years (January 2008 to December 2017). Data about the patients were obtained and analyzed using Microsoft office excel 2007 and SPSS version 25 (IBM SPSS, Statistics) to generate descriptive statistics (frequencies, ratio and percentages) which were presented as tables.

Results: The prevalence of AS is 5.3%. Most of the patients were in the 25-34 age group (69.1%) and the majority (94.5%) were married. Dilatation and curettage (D & C) constituted the commonest risk factor for Asherman's syndrome (21/55, 38.2%) and infertility alone (29, 52.7%), and hypomenorrhea with infertility 10 (18.2%) were the commonest presentation. Twenty-five percent of the patients (14/55) achieved normal menses and 11 (20%) achieved pregnancy following treatment.

Conclusions: AS is not uncommon in our environment. D&C is the commonest risk factor and the patients usually present with infertility and hypomenorrhea.

Keywords: Asherman's syndrome, Achieve pregnancy, D and C, Infertility, UMTH

INTRODUCTION

Adhesion within the endometrial cavity is known as intrauterine adhesion or uterine synechiae. When it is associated with hypomenorrhea or amenorrhea, it is called Asherman's syndrome (AS).^{1,2} The true incidence of AS is unknown, as is dependent on the population studied and classification system used.^{3,4} However it is

steadily increasing particularly where dilation and curettage is still the most common method of pregnancy termination.^{5,6} An incidence of 1.6%, 29.2%, 4.2% has been reported in Abuja, Lagos, Uyo, respectively.^{3,4,6}

The destruction of the basal is endometrium by trauma prevent endometrial thickening in response to ovarian steroids, leading to a spectrum of avascular scarring

which ranges from filmy adhesion, dense bands or complete obliteration.¹⁻⁵ Vigorous curettage of the uterine cavity following a pregnancy complicated by infection is the commonest cause of Asherman's syndrome,⁶ and earlier studies have shown that intrauterine adhesion was found in 25% of women who had dilation and curettage in the postpartum period, in 30.9% following missed abortion and 40% in those that had dilatation and curettage for retained products of conception after miscarriage or retained placenta. The risk also increases with the number of dilatation and curettage performed as 16% is recorded after the first dilatation and 32% after three or more.^{3, 4} Other documented risk factors include evacuation of hydatidiform mole, caesarean section, myomectomy, metroplasty, diagnostic curettage in a non-gravid uterus, B-lynch, uterine artery embolisation, hysteroscopic surgery, insertion of intrauterine device complicated by infection, pelvic radiation, endometrial ablation, tuberculous endometritis, schistosomiasis endometritis and manual removal of placenta.¹⁻⁸

The clinical presentation of patients with Asherman's syndrome include menstrual abnormalities presenting as hypomenorrhea or secondary amenorrhea, others are; lower abdominal pain, dysmenorrhea, subfertility, recurrent miscarriage, fetal malposition and adherent placenta. Diagnosis requires a high index of suspicion with hysteroscopy being the gold standard, where direct visualization under magnification of the uterine cavity is possible. Transvaginal ultrasonography/3D USS, saline infusion sonography, hysterosalpingography and Magnetic resonance Imaging are other methods of diagnosis.¹⁻⁹ The diagnostic and clinical presentation has been used by different authorities to classify Asherman's syndrome based on severity. The commonly used are classification by The American fertility society who classifies it into minimal/mild, moderate and severe, Hamou et al classify it into isthmic, marginal, central and severe. Other classifications were by Nasr et al and the former European society for hysteroscopy.¹⁰ Treatment is aimed at restoring the endometrial cavity size, shape and prevention of adhesion reformation leading to normal function and possibly increasing the chances of pregnancy in patients that present with subfertility. These are achievable by direct vision-hysteroscopic adhesiolysis (gold standard) or through blind adhesiolysis. This procedure can be primary when first attempt at adhesiolysis is carried out or secondary when a repeat adhesiolysis is done.⁸⁻¹⁰ Other methods include; pressure lavage under ultrasound guidance and hysterotomy with transfundal adhesiolysis. Endometrial cavity splinting is done using a paediatric Foley's catheter, intrauterine device, stent or tubing. Endometrial regeneration can be improved by cyclical estrogen/progesterone therapy and sildenafil.¹⁰ Recently also, the use of amnion graft and stem cells from bone marrow and autologous menstrual blood are in small experimental women studies with remarkable results.^{11,12} However the degree of success depends on the severity of adhesion.¹

This study is aimed at determining the prevalence, clinical presentation and treatment outcome of Asherman's syndrome in our hospital.

METHODS

This was a retrospective analysis of patient's with Asherman's syndrome managed at the University of Maiduguri teaching hospital over 10 years (January 2008 to December 2017). The cases were identified using Gynaecology ward and theatre records and their case notes retrieved from the medical record department. A proforma design for the study was used to extract data from the case notes. The information extracted included demographic characteristics, the number of miscarriages, presenting complaint, predisposing factor, diagnostic tool used, management offered and outcome of the treatment. Data obtained was analyzed using Microsoft office excel 2007 and SPSS version 25 (IBM SPSS, Statistics) to generate descriptive statistics (frequencies, ratio and percentages) which were presented as tables.

RESULTS

During the study period, 81 patients were managed for Asherman's syndrome with 55 case files available giving a retrieval rate of 67.90%. There were 1521 gynaecological surgeries during the period of the study giving a prevalence of AS of 5.3% of gynaecological operations. The sociodemographic data of the patients is shown in (Table 1). Most of the patients were in the 25-34 age group (69.1%), followed by the 35-44 age group (18.2%). Most of the patients were married (94.5%) and a majority (34.5%) had tertiary education. Dilatation and curettage constituted the commonest risk factor for Asherman's syndrome (21/55, 38.2%), followed by myomectomy alone (11/55, 20.0%), puerperal infection 10.9% (6/55) and caesarean section 7.3% (3/55) (Table 2). Majority of the patients presented with infertility alone (29, 52.7%), hypomenorrhea and infertility 10 (18.2%) and recurrent miscarriage 5 (9.1%) (Table 3). The treatment outcome of the patients was as shown in (Table 4). Twenty-five percent of the patients (14/55) achieved normal menses and 11 (20%) achieved pregnancy. About 22 patients (40%) were lost to follow up within 6 months of treatment although 18 patients (32.7%) were followed up for one year or more (Tables 4-5).

DISCUSSION

This study revealed a prevalence of Asherman's syndrome of 5.3% with the age group 25-34 years being the commonest age group and D and C the commonest risk factor. The majority of the patients presented with infertility and 20% achieved pregnancy following treatment. The prevalence of Asherman's syndrome of 5.3% found in this study is similar to that found in Uyo, Southwestern Nigeria, higher than the 1.6% reported

from Abuja and 1.4% reported from Kano but lower than 29.2% reported from Lagos Southern Nigeria.^{3,4,6,13}

Table 1: Sociodemographic characteristics of the study group.

Parameters	N	%
Age (years)		
15 - 24	7	12.7
25 - 34	38	69.1
35 - 44	10	18.2
Total	55	100.0
Parity		
0	30	54.5
1-4	24	43.6
5 and above	1	1.8
Total	55	100.0
Marital status		
Single	3	5.5
Married	52	94.5
Total	55	100.0
Educational level		
Primary school	5	9.1
Secondary school	18	32.7
Tertiary education	19	34.5
No formal education	13	23.6
Total	55	100.0
Miscarriages		
0	20	36.4
1	16	29.1
2	11	20.0
3	8	14.5
Total	55	100.0

Table 2: Risk factors of the patients in the study.

Risk factors	N	%
Dilatation and curettage	21	38.2
Myomectomy	11	20.0
Puerperal infection	6	10.9
Non identified	4	7.3
Post caesarean section infection	4	7.3
Caesarean section	4	7.3
Post abortal sepsis	3	5.5
PID	2	3.6
Total	55	100

This varying prevalence maybe because of the variations in the populations studied and the methods of diagnosis employed. For example, hysteroscopy which is more accurate and the gold standard for diagnosis of Asherman's syndrome was used in making the diagnosis in the Lagos study that reported a high prevalence while HSG was the main tool used for making the diagnosis in our study and the study from Kano.

Similar to the reports of other studies majority of the patients in our study were in the age group 25-34 years.^{3,6,9}

Table 3: Presenting complaints of the patients.

Presenting complaints	N	%
Infertility	29	52.7
Hypomenorrhea and infertility	10	18.2
Recurrent miscarriages	5	9.1
Hypomenorrhea, infertility and cyclical pelvic pain	3	5.5
infertility and cyclical pelvic pain	3	5.5
Amenorrhea and infertility	2	3.6
Hypomenorrhea	2	3.6
Amenorrhea	1	1.8
Total	55	100

Table 4: Treatment outcome of the study population.

Outcomes	N	%
Achieved normal menses	14	25.5
Achieve pregnancy	11	20.0
Achieve normal menses and relief in cyclical pain	3	5.4
No change in condition	5	9.1
Lost to follow up	22	40.0
Total	55	100.0

Table 5: Duration of follow up.

Outcomes (months)	N	%
0-3	12	21.8
4-5	21	38.2
6-9	4	7.3
1 year and above	18	32.7
Total	55	100.0

This is the most reproductively active age group and the majority of the patients (94.5%) were married. They are therefore more likely to be pregnant and develop a pregnancy complication like abortions that can result in uterine evacuations which is a common cause of AS. The commonest risk factor for the development of AS in this study was D&C. This was similar to the reports of other studies.^{3,6,7,9} Endometrial injury induces ischemia and inflammatory reactions that lead to hypoxic cellular modifications and release of active substances leading to fibrosis and D&C is traumatic to the endometrium.^{14,15} The other risk factors identified including myomectomy and caesarean sections may have operated through the same trauma to the endometrium that can occur during these procedures. Although similar to the report of Charles et al infectious complications (post abortal and puerperal infections) were found to be risk factors of AS, the role of infection in the pathogenesis of AS remains unclear and Polishuk et al reported that the development

of endometritis after caesarean delivery is not associated with increased incidence of AS.^{6,16}

Patients with AS usually present with menstrual problems (hypomenorrhea or amenorrhea) and/or infertility, and similar to other studies the commonest presentation of the patients in our study was menstrual problems and/or infertility.^{6,7,17} Intrauterine adhesions caused reduction of the endometrial surface and therefore bleeding area leading to menstrual problems and AS is associated with endometrial trophic changes and unresponsiveness that can result in infertility. The treatment of the AS in our centre is done following a carefully developed departmental protocol that consists of D and C, insertions of size 14G Foley's catheter for 10 days, endometrial restoration with estrogen and post-operative antibiotics, although hysteroscopic resection has started in the last 2 years of the study period. Following treatment, 20% of our patients achieve pregnancy. This was higher than the 10.2% pregnancy rate reported by Opadiran et al from Abuja, Nigeria but lower than the 29% reported by Njoku et al, 27.3% reported by Shiktra et al and the 26.6% reported by Bhandari et al.^{6,7,18} The slightly lower pregnancy rate in our study compared to the other studies in our environment may be because 40% of our patients were lost to follow up within six of treatment although 37.2% of our patients were followed up for a year or more. Pregnancy rates ranging between 56 to 76% has been reported in patients with AS treated with hysteroscopic adhesiolysis.^{19,20} Hysteroscopy has become the gold standard for the treatment of AS, as it is safer and more effective compared to D and C. With a laparoscope the adhesions can be viewed directly and the magnification provided by the instrument allows accurate division of the adhesions to restore normal uterine calibre and free tubal Ostia in comparison to blind D and C.

CONCLUSION

AS is not uncommon in our environment and it is usually associated with activities that result in endometrial injury. The commonest presentation is a menstrual abnormality and/or infertility which can be ameliorated with intrauterine adhesiolysis.

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REFERENCES

1. Amenorrhea, In; Hoffman, Schoffer, Bradshaw, Schorge, Halvorson, Cunningham(ed). Williams Gynaecology, 2nd ed. United States of America: McGraw-Hill companies Inc; 2012:16;440-57.
2. Alex S, Wendy YC, Alan HD. In; Decherney AH, Nathan L, Laufer N, Ashley RS eds. Current diagnosis and treatment, Obstetric and gynaecology. 11th ed. United States of America: McGraw-Hill companies Inc; 2013:54;889-99.
3. Opadiran RO, Isah AD, Agida ET, Adewole N. Outcome of intrauterine adhesion management at a Nigerian tertiary hospital; a five year review. *Int J Reprod Contracept Obstet Gynecol.* 2018;7:375-9.
4. Ajayi A B, Afolabi B M, Ajayi V, Biobaba O, et al. Risk factors for intrauterine adhesions in a black African population. *Nigerians. Gynecol Obstet.* 2017; 7:436.
5. Chanrit P, Tawiwat P, Opas S, Usanee S. Intrauterine adhesions: causes and treatment outcomes among Thai women. *J med Assoc Thai.* 2016;99(10):1067-72.
6. Charles N, Cahethan E, Edu E, Boniface A, John E. Risk factors and management outcome of intrauterine adhesion in a constrained socioeconomic environment: a 10 year review in the university of Calabar teaching hospital, Calabar. *EJPM.* 2017;4(3): 25-30.
7. Kwari SD, Idrisa A. Intrauterine adhesions at the university of Maiduguri teaching hospital, Maiduguri, Nigeria. A 3 year review. *Trop J Obstet Gynecol.* 2011;28(10):50-1.
8. Saroja CSM, Nankani A, El-Hamamy E. Uterine compression sutures, an update: review of efficacy, safety and complications of B-Lynch suture and other uterine compression techniques for postpartum haemorrhage. *Arch Gynecol Obstet.* 2010;281(4):581-8.
9. Utuk N, Abasiattai A M, Asuquo O. Intrauterine adhesion in the university of Uyo teaching hospital, south-south, Nigeria. A 10 year review. *Trop J Obstet Gynecol.* 2019;6:39-43
10. Dreisler E, Kjer JJ. In Asherman's syndrome; current perspectives on diagnosis and management. *Int J Women's Health.* 2019;11:191-8.
11. Neeta S, Sujata M, Sona D. Autologous stem cell transplantation in refractory Asherman's syndrome: A novel cell based therapy. *J Hum Reprod Sci.* 2014; 7(2):93-8.
12. Changjiang L, Jia W. The study on the safety and efficacy of amnion graft for preventing the recurrence of moderate to severe intrauterine adhesions. *Genes Dis.* 2020;7(2):266-71.
13. Gaya SA, Adamu IS, Yakasai IA. Review of intrauterine adhesiolysis at the Aminu Kano Teaching Hospital, Kano, Nigeria. *Ann Afr Med.* 2012;11(2): 65-9.
14. Yu D, Wong YM, Cheong Y, Xia E, Li TC. Asherman syndrome-one century later. *Fertil Steril.* 2008;89:759-79.
15. Chen Y, Chang Y, Yao S. Role of angiogenesis in endometrial repair of patients with severe intrauterine adhesion. *Int J Clin Exp Pathol.* 2013;15:1343-50.
16. Polishuk WZ, Anteby SO, Weinstein D: Puerperal endometritis and intrauterine adhesions. *Int Surg.* 1975;60:418-20.
17. Abiodun OM, Balogun OR, Fawole AA. Aetiology, clinical features and treatment outcome of intrauterine

- adhesion in Ilorin, Central Nigeria. *West Afr J Med.* 2007;26:298-301.
18. Bhandari Sh, Bhavne P, Ganguly I, Baxi A, Agarwal P. Reproductive Outcome of Patients with Asherman's Syndrome: A SAIMS Experience. *J Reprod Infertil.* 2015;16(4):229-35.
19. Valle RF, Sciarra JJ. Intrauterine adhesions: hysteroscopic diagnosis, classification, treatment, and reproductive outcome. *Am J Obstet Gynecol.* 1988; 158(6):1459-70.
20. Yamamoto N, Takeuchi R, Izuchi D. Hysteroscopic adhesiolysis for patients with Asherman's syndrome: menstrual and fertility outcomes. *Reprod Med Biol.* 2013;12(4):159-66.

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