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Prevalence; Characteristics and Management of Endometriosis Amongst Infertile Women: A One Year Retrospective Study

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ABSTRACT

Background: Endometriosis appears to affect every aspect of a women's reproductive system resulting in infertility and spontaneous pregnancy loss. This study aims to find out the prevalence & clinical characteristics of endometriosis amongst infertile women.

Settings and Design: A Hospital based retrospective study over a period of one year.

Materials and Methods: It is a retrospective study conducted in the gynaecology department in Institute of Kidney Diseases & Research Centre; Ahmedabad from April 2012 to March 2013 amongst women with a primary complaint of infertility (Primary/ Secondary). A total of 372 patients underwent diagnostic hysterolaparoscopy and of these 180 patients who had laparoscopic evidence of endometriosis was included in the study. All of these patients and their findings were analysed with respect to the clinical signs and symptoms. The outcome after appropriate management was analysed in subsequent follow up.

Statistical analysis: All collected data was entered into the SPSS version 20. Categorical data are expressed in frequency or percentage. Chi-Square test and Fisher-Exact test has been performed to carry out p-value for categorical data. P-value <0.05 shows statistically significant difference.

Results: The frequency of endometriosis among women with infertility subjected to diagnostic hysterolaparoscopy was found to be 48.38%. Statistical significant association with severity of disease was associated with symptoms like dysmenorrhea, chronic pelvic pain, restricted uterine mobility and adnexal tenderness. (p < 0.01) Ultrasound finding of endometrioma with ground glas appearance also had statistical significant association with staging of disease (p < 0.01).

Conclusion: Endometriosis amongst infertile women is increasingly being detected due to greater use of laparoscopy in evaluation of infertility. Though most signs do not correlate with severity of disease however the presence of restricted uterine mobility, adnexal tenderness & chronic pelvic pain should always raise the suspicion of endometriosis. Laparoscopy remains the gold standard for diagnosing and staging endometriosis.

Keywords: Infertility, Laparoscopy, Uterine cavity

INTRODUCTION

Endometriosis is a gynaecological enigma since it is difficult to diagnose and treat. It is defined as a chronic and recurrent disease characterized by the presence and proliferation of functional endometrial glands and stroma outside the uterine cavity. It is responsible for varied and disabling symptoms and also has adverse effects on reproductive potential. The precise mechanism by which endometriosis causes infertility needs to be evaluated. Endometriosis appears to affect every part of a woman's reproductive system. It is postulated that women with endometriosis have increased amount of peritoneal fluid associated with increased peritoneal concentrations of prostaglandins, proteases and cytokines including inflammatory cytokines such as IL 1, IL 6 and TNF alpha, and angiogenic cytokines like IL 8 and VEGF [1]. These alterations have an adverse effect on oocyte, sperm, embryo and fallopian tube function [2].

Endometriosis is one of the most common conditions encountered in gynaecological practice. Classic studies have suggested that 25 to 50% of infertile women have endometriosis and 30-50% of women with endometriosis are infertile [3]. The true prevalence of endometriosis is difficult to quantify as very wide ranges have been reported in literature [4,5]. Endometriosis is found in 45% - 82% of women with chronic pelvic pain and in 2.1%-78% of infertile women [6,7]. Nevertheless, its prevalence depends on patient profile and diagnostic tools utilized. However, its prevalence is 6-21 times higher in infertile as opposed to fertile women [8,9].

The role of Ultrasound in the clinical diagnosis of endometriosis is of limited value as it lacks resolution for visualizing adhesions and superficial peritoneal/ovarian implants. Hence laparoscopy is the mainstay in the diagnosis as it provides a visual proof of the minute endometriosis lesions and helps in staging of the disease. In this study, we aim to find out the prevalence of endometriosis amongst infertile women, the demographic & clinical characteristics associated with endometriosis.

MATERIALS AND METHODS

This is a retrospective study conducted at the department of Obstetrics and Gynaecology, Institute of Kidney disease and Research centre, Ahmedabad, Gujarat for the period from April 2012 to March 2013. This study aimed to determine the demographic, clinical and laparoscopic characterization of infertile women with endometriosis in addition to type of treatment administered.

Inclusion Criteria: All patients of primary or secondary infertility subjected to diagnostic hystero laparoscopy and chromopertubation test who were diagnosed to have endometriosis were included in the study.

Exclusion Criteria: Women with PID, adhesions due to previous surgeries or infections were excluded.

All patients included in the study were analysed with respect to the following characteristics.

Clinical Characteristics: Points noted were age, type of

infertility, duration of infertility, menstrual cycle - frequency and flow, association of symptoms like dysmenorrhoea, dyspareunia, chronic pelvic pain, urinary symptoms & their correlation to stage of endometriosis.

- Physical Examination: Findings were analysed with respect to presence of abdominal/adnexal masses, mobility of uterus and presence of adnexal tenderness.
- USG FINDING: Particularly to note endometriomas and probe tenderness.
- Laparoscopic Findings: Endometriotic lesions which were noted varied from dark blue, powder-burn black, red, white, yellow, brown or non-pigmented lesions. The size, depth and location of these lesions were noted to grade the severity of endometriosis. This laparoscopic staging was based on the revised AFS scoring [10] which categorized the finding into 4 stages.

Stage I: (Minimal) involved a few endometrial implants, most often in the cul de sac.

STAGE II: (Mild) comprised of endometrial implants affecting one or both ovaries.

STAGE III: (Moderate) involved moderate levels of endometriosis with implants in several reproductive areas & in one or both ovaries.

Stage IV: (Severe) involved wide spread endometriosis implants through the pelvic area.

STATISTICAL ANALYSIS

All collected data was entered into the SPSS version 20. Categorical data are expressed in frequency or percentage. Chi Square test and Fisher Exact test has been performed to carry out P-value for categorical data. p-value <0.05 shows statistically significant difference.

RESULTS

A total of 372 patients with infertility were subjected to diagnostic hysterolaparoscopy & chromopertubation test during the period from April 2012 to March 2013. Of these, 180 (48.38%) patients had laparoscopic evidence of endometriosis. One hundred thirty five patients (75.0%) had primary infertility and forty five patients (25.0%) had secondary infertility. The mean age of patients was 29± 4.3 years (Range: 19-40 years). Amongst the 180 patients studied, apart from infertility, the commonest complaints were dysmenorrhea (42.22%) followed by menstrual irregularity (17.77%), menorrhagia (12.2%), dyspareunia (9.4%) and chronic pelvic pain (4.41%). However, more than 50% of cases were asymptomatic. There was a statistical significant association between adnexal tenderness and restricted uterine mobility with staging of the disease (p< 0.01). Abnormal USG findings were seen in 15% of cases. Presence of cysts/

endometrioma with ground glass appearance was seen in 12.7% of cases. This particular finding was found to be clinically significant as sonograpically detected endometriomas were confirmed laparoscopically. All 11 (100%) cases of stage IV endometriosis in our series had endometriomas in one or both ovaries. Based on Revised AFS score (1985); STAGE 1 endometriosis was seen in 119 patients (66.1%); STAGE II endometriosis in 39 patients (21.66%); STAGE III endometriosis in 11 patients (6.11%); and STAGE IV endometriosis in 11 patients (6.1%). [Table/Fig-1] shows the association of clinical signs and symptoms with stage of disease. [Table/Fig-2] shows the association of laparoscopy& USG findings with stage of disease. In our study, there was a definite correlation of USG and lap evidence of endometriosis with stage of disease. Although most patients of endometriosis in our study were asymptomatic, the presence of dysmenorrhea, dyspareunia and chronic pelvic pain are clinically significant and should always anticipate the presence of subtle endometriosis in these patients. All patients of minimal and mild endometriosis were treated by fulguration /cauterization followed by three doses of GnRH agonist. Moderate and severe endometriosis were treated accordingly depending on laparoscopic findings i.e. adhesiolysis, endometrioma cyst wall excision. This was followed by three doses of leupragon (3.75mg) at an interval of 28 days.

DISCUSSION

Endometriosis affects 2.5-3.3 percent of women of reproductive age [11] and is diagnosed in 20-68 percent of the women studied for infertility [12]. According to Tsuzi et al., worldwide endometriosis has been found up to 63% [13]. The true incidence of endometriosis is difficult to establish since endoscopy or laparotomy is required for a definitive diagnosis and the disease may still undoubtedly exist in patients who are asymptomatic. It is generally believed, the disease is relatively less common in India, Pakistan, Iran, countries of Middle East and black Africa [14]. Our study demonstrates a very high incidence of 48.33%. Therefore, we feel that the clinical impression of low incidence in Asian and black women may have been due to limited medical and diagnostic facilities available to these women. In our study, every alternative patient had evidence of endometriosis. The commonest sites of endometriosis were the uterosacral ligament (65 cases), Pelvic wall (56cases) Pouch of douglas (40 cases), ovarian fossa (34 cases), and endometrioma (23 cases). The other rare sites were uterovesical fold of peritoneum, bowel, round ligament, rectovaginal septum and diaphraghm. One interesting point which we noted was that left side was more involved than the right though it was not found to the statistically significant. The mean age of patients was 29± 4.3 years. This is in comparison to other studies which quote low prevalence of endometriosis in either extremes of age & high prevalence in women of reproductive age [15]. Seventy five percent of cases were cases of primary infertility & 25% of case was of secondary infertility. This finding is similar to other descriptive studies [16].

Clinical signs and symptoms	Stage 1 (of 119)		Stage 2 (of 39)		Stage 3 (of 11)		Stage 4 (of 11)		p-value
	N (%)	OR	N (%)	OR	N (%)	OR	N (%)	OR	,
Menstrual Irregularity	25 (21.00%)	1.00	4 (10.25%)	0.43	1 (9.09%)	0.38	2 (18.18%)	0.84	0.40 (NS)
Heavy Menstrual flow	12 (10.08%)	1.00	5 (12.82%)	1.31	2 (18.18%)	1.98	3 (27.27%)	3.34	0.36 (NS)
Scant Menstrual flow	6 (5.04%)	1.00	3 (7.69%)	1.57	2 (18.18%)	4.19	1 (9.09%)	1.97	0.39 (NS)
Dysmenorrhea	40 (33.61%)	1.00	23 (58.97%)	2.83	6 (54.54%)	2.37	7 (63.63)	3.45	0.01 *
Dyspareunia	5 (4.20%)	1.00	4 (10.25%)	2.61	3 (27.27%)	8.55	5 (45.45%)	19	<0.01 *
Chronic Pelvic pain	1 (0.84%)	1.00	1 (2.56%)	3.11	1 (9.09%)	11.8	5 (45.45%)	98.33	<0.01 *
Abdominal mass	0 (0.00%)	1.00	0 (0.00%)	N/A	0 (0.00%)	N/A	0 (0.00%)	N/A	-
Tenderness	4 (3.36%)	1.00	4 (10.25%)	3.29	6 (54.54%)	34.5	11 (100%)	N/A	<0.01 *
Adnexal mass	1 (0.84%)	1.00	0 (0.00%)	N/A	2 (18.18%)	26.22	6 (54.54%)	141.6	<0.01 *
Restricted Mobility	1 (0.84%)	1.00	3 (7.69%)	9.83	4 (36.36%)	67.42	10 (90.90%)	1180	<0.01 *

[Table/Fig-1]: Association of clinical presentations of endometriosis with staging (OR represents Odds Ratios. p <0.05 considered to be statistically significant difference Here, NS represents Non-Significant difference between these groups.)

Laparoscopic findings	Stage 1 (of 119)		Stage 2 (of 39)		Stage 3 (of 11)		Stage 4 (of 11)		p-value
	N (%)	OR	N (%)	OR	N (%)	OR	N (%)	OR	
Blocked tubes on laparoscopy	16 (13.44%)	1.00	10 (25.64%)	2.22	5 (45.45%)	5.36	9 (81.81%)	28.97	<0.01 *
Endometrioma on Ultrasonography	0 (0.00%)	1.00	7 (17.94%)	N/A	5 (45.45%)	N/A	11 (100%)	N/A	<0.01 *
Endometrioma on laparoscopy	0 (0.00%)	1.00	2 (5.12%)	N/A	3 (27.27%)	N/A	11 (100%)	N/A	<0.01 *

[Table/Fig-2]: Association of ultrasonographic & laparoscopic findings with staging of endometriosis OR represents Odds Ratios. p < 0.05 considered to be statistically significant difference Here, NS represents Non-Significant difference between these groups.)

Commonest symptoms encountered in our study were dysmenorrhea (42.2%); irregular cycles (17.7%); menorrhagia (12%); dyspareunia (9.4%) and chronic pelvic pain (4.4%). However, other rare complaints which were noted were urinary complaints like dysurea and difficulty in defaecation in one case of stage 4 endometriosis with frozen pelvis. Statistical significant association of symptoms and signs like dyspareunia, chronic pelvic pain, restricted uterine mobility and adnexal tenderness with staging of disease was noted. However majority of the cases (57.8%) were asymptomatic. We feel that, though none of clinical signs are decisive of endometriosis, positive clinical findings like tenderness on clinical examination, fixation or relatively decreased mobility of uterus or a fixed retroverted uterus or a pelvic mass should always raise the index of suspicion towards endometriosis. The final diagnosis is always by laparoscopy, the gold standard in diagnosis of endometriosis, preferably with histological confirmation [17].

Transvaginal ultrasound lacks adequate resolution for visualizing adhesions and superficial peritoneal/ovarian implants. But when the presence of endometrioma with a typical ground glass appearance is identified, it usually indicates that moderate to severe endomeriosis is present. In our study, there was significant correlation between presence of cysts/endometriomas with ground glass appearance & severity of disease. Out of the 23 diagnosed cases of endometrioma by transvaginal USG, 17 cases were confirmed to have stage III and IV endometriosis on laparoscopy according to R-AFS score (1985).

Majority of the cases of our study had STAGE I endometriosis i.e. 110 cases (66.1%). Most of these patients were asymptomatic and this suggests an early presentation. STAGE II endometriosis was seen in 21% of case and STAGEIII & IV in 6.1% of cases each. All cases of stage IV endometriosis in our study (11 cases) had bilateral endometrioma & frozen pelvis with bowel adhesions. Tubal block was seen in almost 81.81% of case of stage IV endometriosis. Thus there were strong association of laparoscopic findings of endometrioma and blocked tubes. Since our study is a retrospective study, one of the important drawbacks is certain data could not be extrapolated like BMI. BMI & endometriosis by European & Western Studies suggested a positive association [18,19].

All patients in stage I/II received medical management of 3 doses of leupragon (3.75mg) along with fulguration by monopolarcautery of endometriotic spots. Stage III/IV endometriosis underwent adhesiolysis & endometrioma cyst wall excision (based on individual findings) followed by 3 doses of leupragon at 28 days interval. Thus this study highlights the higher prevalence of endometriosis

in our population particularly in asymptomatic infertile females. Laparoscopy is the gold standard for diagnosing endometriosis as recommended by ESHRE guidelines [20].

CONCLUSION

Endometriosis in infertile females is not uncommon & is increasingly being detected because of greater use of diagnostic modalities like laparoscopy in evaluation of infertility. Though most females are asymptomatic, dysmenorrhoea, chronic pelvic pain, restricted uterine mobility & adnexal tenderness raises the suspicion of endometriosis. Ultrasound evidence of endometrioma has strong correlation to severity of disease. But it is of limited value for diagnosing and determining extent of endometriosis. Laparoscopy remains the gold standard for diagnosing and staging endometriosis.

REFERENCES

- Bedaiwy MA, Falcone T, Sharma RK, Goldberg JM, Attaran M, Nelson DR, et al. Prediction of endometriosis with serum and peritoneal fluid markers:a prospective controlled trial. Hum Reprod. 2002;17:426-31
- Lebovic DI, Mueller MD, Taylor RN. Immunobiology of endometriosis. Fertil Steril.
- 2001;75:1-10. Missmer SA, Hankinson SE, Spiegelman D, Barbieri RL, Marshall LM, Hunter DJ. Incidence of laparoscopically confirmed endometriosis by demographic, anthropometric, and lifestyle factors. *Am J Epidemiol*. 2004;160:784-96. Stilley JA, Birt JA, sharpe-timms KL. Cellular and molecular basis for endometriosis associated infertility. *Cell tissue Res*. 2012;349(3):849-62. Practice Committee of American Society for Reproductive medicine. Endometriosis
- and infertility: a committee opinion. Fertil Steril. 2012;98(3):591-98.
- Mahmood TA, Templetion A. Prevalance and generis of endometriosis. Hum Reprod. 1991;6:544-49.
- Meuleman D, D'Hooghe T. High prevalance of endometriosis in infertile women with normal ovulation and normospermie patients. Fertil steril. 2009:68-74. Rawson JM. Prevalance of endometriosis in asymptomatilc women. J Reprod Med.
- 1991:36:513-15.
- Louis GMB, Hedigar ML, Peterson CM, Croughan M, Sundaram R, Stanford J, et al Incidence of endometriosis by study population and diagnostic method: the ENDO study. Fertil steril. 2011;96:360-65.
- [10] American Society for Reproductive Medicine. Revised American Society for Reproductive Medicine classification of endometriosis: 1996. Fertil Steril. 1997;67:817-21.
- [11] Houston DE, Nouer KL, Melton LJ III, Selwyn BI, Hardy RJ. Incidence of pelvic endometriosis in Rochestar, Minnessota, 1970-1979. AMJ Epedemiol. 1987;125:959-69.
 [12] Matorras R, Rodrigez F, Pijoan JI, et al. Epidemiology of endometriosis in infertile
- women Fertil Steril 1995:63:34-38 Tsuji I, Ami K, Miyazaki A, Hujinami N, Hoshiai H. Benefit of diagnostic laparoscopy for
- patients with unexplained infertility and normal hysterosalping graphy findings. *Tohoku J Exp Med*. 2009;219:39-42.
- Ranney B. Endometriosis: IV Hereditary Tendency. ObstetGynaecol. 1971;37:734 Farquhar CM. Extracts from the "Clinical Evidence" Endometriosis BMJ. 2000;320:1449-
- [16] Gao X, Outley J, Bottenam M, Spalding J, Simon JA, Pashos CL. Economic burden of endometriosis. Fertil Steril. 2006;86:1561-72.
- Albert LH, Izabella K, Pamela S. Invasive and noninvasive methods for diagnosis of endometriosis. *Clinobstet Gynecol.* 2010;53(2):413-19. Ferrero S, Anserini P, Remorgida V, Ragni N. Body mass index in endometriosis. *Eur J obstet gynaecol Reprod Biol.* 2005;121:94-98.
- Hediger ML, Hartnett HJ, Louis GM. Association of endometriosis with body size &figure. Fertile steril. 2005;84:1366-74.
- Kennedy S, Berggquist A, Chapron C, D'Hooghe T, Dunselman G, Greb R, et al. ESHRE guidelines for the diagnosis and treatment of endometriosis. *Hum Reprod*. 2005;20:2698-704.

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