IJPSR (2011), Vol. 2, Issue 4





INTERNATIONAL JOURNAL PHARMACEUTICAL SCIENCES RESEARCH



Received on 02 December, 2010; received in revised form 22 January, 2011; accepted 24 March, 2011

HISTOPATHOLOGICAL CHANGES ASSOCIATED WITH ECTOPIC TUBAL PREGNANCY

N. Dahiya*, S. Singh, R. Kalra, R. Sen and S. Kumar

Department of Pathology, Pt. BDS PGIMS, Rohtak, Haryana, India

Keywords:

Ectopic gestation, Salpingitis isthmica nodosa, Pelvic inflammatory disease, Salpingitis

Correspondence to Author:

Dr. Narender Dahiya

727/23 DLF Colony, Rohtak, Haryana, India

Aim: The aim of this prospective study was to know the variable histopathological changes in the fallopian tube after having ectopic gestation and its correlation with predisposing factors.

ABSTRACT

Materials and Methods: 100 fallopian tubes having ectopic gestation for study group and

25 fallopian tubes received after sterilization operation for control group submitted in department of pathology between Jan. 2007- March 2009 were included. The conventional H&E stained microsections were prepared from specimen for extensive study. The sections were examined histopathologically for various pathological lesions by standard criteria.

Results: Acute salpingitis was found in 18% of patients. Chronic salpingitis and follicular salpingitis was seen in 39% of patients, while chronic salpingitis with salpingitis isthmica nodosa was found in only 8% of patients. Vessels were sclerotic in 8% of cases. Calcification was found in 7% cases. No case of tuberculosis and endometriosis was observed in our study group. Conclusion: Pelvic inflammatory disease, Salpingitis isthmica nodosa, Acute and Chronic salpingitis are predisposing factors for ectopic tubal gestation.

INTRODUCTION: Fallopian tube is one of the commonest sites for ectopic gestation. Pelvic inflammatory disease is one of the important predisposing factors for ectopic gestation. It requires careful diagnosis and prompt management to decrease maternal morbidity and mortality. Ectopic pregnancy accounts for 1.2% of all pregnancies and it is a serious cause of maternal morbidity and mortality ¹. Still the true incidence of ectopic gestation is difficult to determine accurately as it varies greatly according to race, socio economic factors and underlying predisposing risk factors ².

Ectopic gestation (fig. 1) occurs most frequently in the fallopian tubes accounting for approximate 95% of the cases 3. In case of diseased fallopian tube or other predisposing risk factor the fertilized ovum is structured in the tube itself leading to ectopic gestations ⁴. PID is considered a significant predisposing factor. On epidemiological grounds (correlations between time trends and the geographical distribution of hospital admissions for salpingitis (fig. 2) and those for ectopic pregnancy), there is a poor association between pelvic inflammatory disease and ectopic pregnancy 5. Keeping in view the high rate of tubal pregnancy and highly variable incidence of pathological changes in the tube, this study was conducted to evaluate the association between ectopic risk factors and histopathological pregnancy, findings.

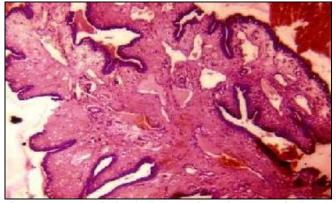


FIG. 1: TUBAL ECTOPIC GESTATION- OEDEMATOUS AND CONGESTED PLICAE (H & E- 40X)

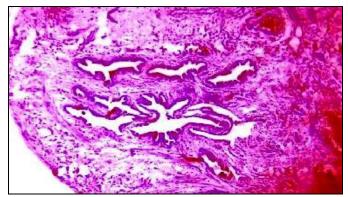


FIG. 2: SALPINGITIS ISTHMICA NODOSA- DIVERTICULAE OF TUBAL EPITHELIUM, THE THICKENED WALL SEPARATED BY BROAD BANDS OF SMOOTH MUSCLES (H & E- 40X)

The aim of the project was to study the histopathological changes in the fallopian tube associated with ectopic tubal pregnancy and to observe for predisposing factors, if any.

MATERIAL AND METHODS: In this prospective study, 100 fallopian tubes having ectopic gestation for study group and 25 fallopian tubes received after sterilization operation for control group were included. All these fallopian tubes (diseased and normal) were submitted in department of pathology for confirmation of disease of organ.

RESULTS: The maximum number of patients was in the age group of 25-29 years. Seventy nine percent of patients had ectopic gestation in ampullary region of fallopian tube and 16% of patients in fimbrial end of the fallopian tube in our study. Forty seven percent of patients had used an intrauterine contraceptive device (IUCD) at one or other time. Twenty seven present were having IUCD- in- situ at the time of presentation in the hospital, 53% of patients had never used any type of intrauterine contraceptive device.

TABLE 1: RELATION OF IUCD USAGE AND TUBAL ECTOPIC PREGNANCY

Usage	No. of patients	Percent
Current usage	27	27
Past usage	20	20
Not used	53	53

Acute salpingitis was found in 18% of patients. Chronic salpingitis and follicular salpingitis was seen in 39% of patients, while chronic salpingitis with SIN was found in only 8% of patients. Vessels were sclerotic in 8% of cases. No case of tuberculosis and endometriosis was observed in our study group.

TABLE 2: PATHOLOGIC FINDINGS OF THE PATIENTS WITH ECTOPIC PREGNANCY

Associated abnormalities	No. of patients	Percent
Acute salpingitis	18	18
Chronic salpingitis +follicular salpingitis	39	39
Chronic salpingitis + SIN	8	8
SIN	6	6
Calcification	7	7
Sclerotic vessel	8	8
Tuberculosis	0	0
Endometriosis	0	0

In, out of total is 56 contralateral tubes were examined of which, 13 patients (23.21%) had chronic salpingitis, three (5.35%) patients had SIN and two (3.5%) had acute salpingitis. Thirty eight (67.85%) patients were unremarkable. Twenty five fallopian tubes from control group examined did not show any significant pathology.

TABLE 3: PATHOLOGICAL FINDINGS IN CONTRALATERAL FALLOPIAN TUBE (WEIGHT INVOLVED WITH ECTOPIC PREGNANCY) (N=56)

Normal	38	67.85%
Acute salpingitis	2	3.5%
Chronic salpingitis	13	23.21%
SIN	3	5.35%

DISCUSSION: Ectopic pregnancy continues to be a significant cause of morbidity and mortality in women of reproductive age group throughout the world. Ruptured tubal ectopic pregnancy is also implicated in maternal death during the first trimester of pregnancy ⁶⁻¹¹.

There has been significant rise in incidence of ectopic pregnancy from 3-4 per 1000 to almost 16 per 1000 pregnancies ¹². This rise has been attributed to the residual tubal pathology after salpingitis to the current wide spread use of intrauterine contraceptive device (IUCD) and to the improved methods of diagnosis and reporting ².

Fallopian tube is common site for ectopic pregnancy with the highest frequency seen in ampullary portion ¹³ (79% in our study), while 16% of patients had fimbrial involvement consistent with findings of other studies ^{6, 14}. A high incidence of fimbrial ectopic pregnancy (16%) in our study was reason for presence of hemoperitoneum in our patients consistent with other studies ¹⁵.

78% patients had rupture tubal ectopic tube while in 22% cases intact tube was present. In the present study 22 patients had histopathologically intact tubal wall and all these women reported between 6.1-7 weeks of gestation with symptoms of chronic ectopic gestation. Tubal rupture may occur when size of pregnancy become too large to be accommodated by narrow lumen of fallopian tube. However various studies had reported the period of gestation in ruptured or unruptured tubal ectopic pregnancy significantly indifferent.

The risk among intra uterine contraceptive device users is 2.94-4.5 times that in non-users. In this study it was found that 47% of patients were having current or past history of usage of intra uterine contraceptive device. However in contrast only 3.9% had in situ contraceptive device and 5.6% were using oral contraceptives in an eight year review of ectopic pregnancy by Helvacioglu *et al.*, ¹⁶.In the present study 35% patients were found to have a history of pelvic inflammatory disease and 19% admitted to previous abdominal surgery. Out of these 19 % patients, (84.19%) has history of some tubal surgery, tuboplasty and salpingectomy etc. The figures of present study are consistent with studies conducted by other authors ¹⁶.

Of the total 100 cases of tubal ectopic gestation, 86% showed severe associated abnormality other than the effects of developing ectopic pregnancy in the form of tubal dilation, hemorrhage, necrosis, presence of trophoblastic tissue and villi. Eighteen cases showed acute salpingitis by virtue of presence of acute inflammatory cells. Kutluay *et al.*, reported acute salpingitis in 34.8% cases, but related and attributed this directly to ovum plantation ¹⁷. However, if this is so, the incidence would have been higher than observed.

The next common pathological finding associated was chronic salpingitis (39%) which was characterized by presence of lymphoplasmacytic infiltrates in the lamina propria and by distortion of tube due to fibrosis.

Follicular salpingitis was diagnosed when mucosal rugae had lost their normal structure and were represented by thickened mucosal folds adherent to one another, forming partitioned cavities of variable sizes ¹⁸. Some authors have reported a very high incidence of chronic salpingitis of 89% and 88% in ectopic pregnancy ¹⁹⁻²¹. Our incidence of 39% is likely due to advent of antibiotics, better management and lower incidence of pelvic inflammatory diseases in our population.

Salpingitis Isthmica Nodosa is other well described pathological lesson of unknown etiology, which is defined as the microscopic presence of tubal epithelium, well in myosalpinx or beneath the tubal serosa in a complete cross section of fallopian tube. Features of SIN were reduced in 8% cases and six cases had associated inflammation. In incidence of 45.9% and 36.4% of salpingitis isthmica nodosa has been reported by other studies ^{14, 19}, where as 12% by Kutluacy *et al.*, ¹⁷.

In our study presence of chronic salpingitis with SIN in six cases (6/8) indicate association between Salpingitis Isthmica Nodosa and other infectious lesions. Seven cases in this study showed

calcification in the wall of tubes which may be squeal of some healed lesion such as tuberculosis. However multiple sections may sometimes be required for diagnosis of active disease. Endometriosis was not found in any one of our cases in the same tube or in contralateral tube.

ISSN: 0975-8232

CONCLUSION: Salpingitis Isthmica Nodosa, acute as well as chronic salpingitis is associated with high incidence of ectopic gestation. This inflammatory disease may be secondary to postpartum or postabortal or intrauterine contraceptive device usage. With increased knowledge and experience early recognition of signs and symptoms of pelvic inflammation and adequate antibiotic treatment in acute phase of disease can prevent ectopic gestation by presenting certain predisposing factors.

REFERENCES:

- Karaer A, Filiz AA, Batioglu S. Risk factors for ectopic pregnancy: A case-control study. Aus NZ J Obstet Gynaecol 2000; 46: 521-7.
- Stabile I, Grudzinskas JG. Ectopic pregnancy; A review of incidence, etiology and diagnostic aspects. Obstet Gynaecol Surv 1990; 45: 335-47.
- Niles JH, Clark JFJ. Pathogenesis of tubal pregnancy. Am J Obstet Gynecol 1969: 105: 1230-4.
- 4. Anderson MC. The fallopian tube. In: Symmers WC, editor. Systemic Pathology, Female Reproductive System. 3rd ed. London: Churchil Livingstone; 1991; 6: 241-61.
- 5. Beral V. An epidemiological study of recent trends in ectopic pregnancy. Br J Obstet Gynaecol 1975; 82: 775-9.
- Roussos D, Panidis D, Matalliotakis I, Mavromatidis G, Neonaki M, Mamopoulos M, et al. Factors that may predispose to rupture of tubal ectopic pregnancy. Euro J Obstet Gynecol 2000; 89: 15-7.
- Dwarakanath LS, Mascarenhas L, Penketh RJA. Persistent ectopic pregnancy following conservative surgery for tubal pregnancy. Br J Obstet Gynaecol 1996; 103: 1021-4.
- 8. Karande VC, Flood JT, Heard NI. Analysis of ectopic pregnancies resulting from in vitro fertilization and embryo transfer. Hum Reprod 1991; 6: 446-9.
- Peek C, Graham FM. Ectopic pregnancy in a non-patent fallopian tube following transfer of embryos to the contralateral tube. Hum Reprod 1992; 7: 136-40.
- 10. Coste J, Job-Spira N, Aublet-Cuvelier B. Incidence of ectopic pregnancy. First results of a population-based register in France. Hum Reprod 1994; 9: 742-5.
- 11. Rouso D, Klearchou N, Rouso I. Combined intrauterine and abdominal pregnancy. J Obstet Gynaecol 1997; 17: 62-7.

- 12. Gupta V, Goel G, Gupta R, Bansal S, Chaturvedi J. Conventional surgical management of ectopic pregnancy in remote areas. Obstet Gynecol 2007; 57: 142-4.
- 13. Kumtepe Y, Borekci B, Polat P, Cetinkaya K, Kadanali S. The rarest form of ectopic pregnancy; Intramural ectopic pregnancy and medical treatment. Turkish German Gynaecol. 2007; 8: 416-9.
- 14. Homm RJ, Holtz G, Garvin AJ. Isthmic ectopic pregnancy and salpingitis isthmica nodosa. The American fertility society. 1987; 48: 756-60.
- 15. Saxon D, Falcone T, Mascha EJ, Marino T, Yao M, Tulandi T. A study of ruptured tubal ectopic pregnancy. Obstet Gynecol 1997; 90: 46-9.
- 16. Helvacioglu A, Long EM. Yang SL. Ectopic pregnancy. Obstet Gynecol. 1979; 22: 87-92.

17. Kutluay L, Vicdan K, Turan C, Batioglu S, Oguz S, Gokmen O. Tubal histopathology in ectopic pregnancies. Euro J Obstet Gyncecol 1994; 57: 91-4.

ISSN: 0975-8232

- 18. Majumdar B, Henderson PH, Semple PA. Salpingitis isthmica nodosa; a high risk factor for tubal pregnancy. Pathol Gynecol Obstet 1983; 62: 73-8.
- Dubuisson JB, Aubriot FX, Cardone V, Vacher-Lavenu MC. Tubal causes of ectopic pregnancy. Fertil Steril 1986; 46: 970-
- 20. Green LK, Kott ML. Histopathologic findings in ectopic tubal pregnancy. Int J Obstet Gynaecol Pathol 1989; 8: 255-6.
- 21. Persaud Y. Etiology of tubal ectopic pregnancy. Obstet Gynaecol 1970; 36: 257-63.
