Clinicopathological Features of Ovarian Tumours - A Prospective Observational Study

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Abstract: Information from developing countries regarding clinicopathological features for ovarian tumours is lacking. Influence of menarche, menopause, nulliparity, mean age of presentation and type of tumour needs to be identified. This will help develop an analysis for clinicopathological features of ovarian tumour. This was a prospective observational study conducted from 1 January 2014 to 31 August 2015 at KLE’s Dr. Prabhakar Kore Hospital and Medical Research Centre, Belgaum which included 119 patients satisfying the inclusion criteria. Incidence of ovarian tumour and clinicopathological features of ovarian tumours was studied. Percentage distribution of clinical and pathological features of ovarian tumours was studied. The incidence of ovarian tumours at KLE’s Dr. Prabhakar Kore Hospital and Medical Research Centre from 1 January 2014 to 31 August 2015 was found to be 6.9% of all gynaecological admissions. In a study of 119 women 92.43% of women presented with pain per abdomen, 83% of women were multiparous. Only 17.64% of women were post menopausal. Amongst 101 patients whose HPR was available, 14 patients had malignant lesions on HPR. The commonest benign lesion was serous cystadenoma followed by simple cyst and the commonest malignant lesion being papillary adenocarcinoma. The sensitivity and specificity of CA 125 in detecting malignant lesions among 70 patients was 70% and 85% respectively. The sensitivity and specificity of RMI in comparison to HPR in 68 patients was found to be 66.6% and 94.64% respectively. Thus, it is concluded that on morphological grounds, tumours originating from surface epithelium are the commonest variant and various modalities will help in early detection of malignant lesions of ovary thereby, reducing the mortality rates.

Keywords: Ovarian Tumour, Benign ovarian tumour, malignant ovarian tumour, Borderline ovarian tumour, Sensitivity, Specificity

INTRODUCTION

Ovarian tumours frequently present as adnexal masses and are frequent reasons for referral to Gynaecologist. Information from developing countries regarding clinicopathological features for ovarian tumours is lacking. The influence of mean age of presentation, parity, menopause, type of tumour needs to be studied [53]. This will help develop a analysis for clinicopathological features of ovarian tumour. This encouraged us to conduct the present study at KLE’s Dr. Prabhakar Kore Hospital and Medical Research Centre, Belgaum.

Ovarian tumours are common form of neoplasia in women. Ovarian tumours constitute 3% to 4% of all Gynaecological admission [59]. Ovarian tumours account for about 30% of female genital cancers. Asian countries have rate of 2 to 6 new cases per 1, 00,000 women per year.² Benign ovarian cysts are the commonest constituting about 90% of ovarian tumours. Gynecologists receive the major load due to ovarian lesion not only because of anatomical location but also since these tumours may remain unnoticed for long period of time[57]. Amongst benign tumours, 60% of them are epithelial in origin. Among benign epithelial tumour, serous cystadenoma are most common (30%), occurring most commonly in reproductive age group [60]. They are bilateral in 10% of cases. Benign or mature cystic teratoma is the most common germ cell tumour, filled with thick sebaceous material. They account for 40% of all ovarian tumours. Mostly benign ovarian tumours are asymptomatic. If symptomatic present with dull aching pain may be acute severe pain in torsion, rupture, haemorrhage, infection [3]. They may present with menstrual disturbances in hormone secreting tumour like granulosa cell tumour.

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Borderline ovarian tumours or ovarian epithelial tumours of low malignant potential were first described by Taylor in 1929 [3, 60]. Histologically, these are intermediate between truly benign neoplasms and those with invasive characteristics. They constitute 10-15% of all epithelial tumours; prevalence being 2.5 per 10,000 women [61]. Ovarian cancer is the leading cause of death in women with female genital cancers in developing countries. A women’s lifetime risk has been estimated to be about 1 in 55, which represents an increase from the 1970 [1]. Ovarian cancer is the fifth most common cause of cancer death in women. It is the third most common Gynaecological malignancy among women in western world, hence is the most lethal. Epithelial ovarian cancer is the eight most common cancer in women, and uterine (corpus and endometrial) is fourth. The ovaries are the ninth most common site of cancer in women. So, to know the incidence of ovarian tumour at KLE’s Dr. Prabhakar Kore Hospital and Medical Research Centre and to study the clinicopathological features of ovarian tumours, the present study was planned.

MATERIAL AND METHOD
Information was gathered from patients with ovarian tumours during interview regarding clinicopathological features of ovarian tumours which included demographic features, menstrual and reproductive history, clinical features, and pathological features. Investigations like USG, tumour markers, CT/ MRI were performed. After the enrolment demographic data, reproductive, obstetric history was obtained. These findings were recorded on a pre designed proforma.

Early menarche is defined as age <11 years at the onset. Late menopause is defined as >51 years of age.

All patients admitted with ovarian tumour at KLE’s Dr. Prabhakar Kore Charitable hospital and MRC, Belgaum from 1st January 2014 to 31st August 2015 which fulfilled the inclusion criteria i.e. all patients with our ovarian tumour attending OPD and admitted with the same at KLE’s Dr. Prabhakar Kore Hospital and Medical Research Centre, Belgaum. All patients given neoadjuvant chemotherapy were also included. Exclusion criteria included ovarian metastasis from any other malignancy and Recurrence of ovarian tumour. Women fulfilling the selection criteria were explained about the nature of study and a written informed consent was obtained prior to enrolment. The ethical clearance was obtained from the Institutional ethics committee, Jawaharlal Nehru Medical College, Belgaum.  The percentage distribution of clinical features and pathologic features of ovarian tumours was found. Categorical outcomes were summarized as rates and numerical outcomes as mean.

RESULTS
A total of 119 cases were studied from 1 January 2014 to 31 August 2015. The data obtained was coded and entered into master chart. The incidence of ovarian tumours was 6.9% of all Gynaecological admissions at KLE’s Dr. Prabhakar Kore Hospital and Medical Research Centre, Belgaum.

In the present study of 119 women, 18 were not operated and in the remaining 101 the histopathological reports were available.
In the present study 25.21% (30) of the women were in age group of 41 to 50 yrs and 24.36% (29) of the women were in 21 to 30 yrs, 22.68% (27) of women were in 30-40 years, 12.6% (15) women in >60 years age group. The mean age of the study population was 40.60 yrs.

In the study of 119 women 92.43% (110) of women presents with complaint of pain per abdomen and vaginal bleeding present in 29.41% (35) of women. Amongst signs 86.55% (103) of women had bulky uterus while ascites was present in 8.40% (10). In the present study 83.19% of women were multiparous while 10.08% were primiparous while 6.72% of women were multigravida. In the present study 82.35 % of women were pre menopausal while only 17.64% of women were post menopausal.
In this study of the total 70 women who had CA 125 value, 77.14% (54) of women had serum CA 125 levels of <35 IU/ml while 22.85% (16) of women had serum CA 125 levels >=35 IU/ml.

In this study, RMI Score was available in 68 women amongst which RMI Score was found to be < 200 in 83.82% (57) of women and in 16.17% (11) of women it was >200. In the present study the histopathological reports showed benign lesions in 86.13% (87) of women while in 13.86% (14) of the women malignant lesions were noted. In the present study, commonest malignant lesion was found to be serous papillary adenocarcinoma 50% (07) followed by endometrioid carcinoma and granulosa cell tumour 14.28% (02) each. In the present study, the commonest benign lesion was found to be serous cystadenoma 35.63% (31) followed by simple cyst and mucinous cystadenoma 17.24% (15) each out of the total 87 patients who had histopathological report showing benign lesions.

<table>
<thead>
<tr>
<th>Malignant</th>
<th>Number(n=14)</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Granulosa tumour</td>
<td>02</td>
<td>14.28</td>
</tr>
<tr>
<td>Serous cystadenocarcinoma</td>
<td>01</td>
<td>07.14</td>
</tr>
<tr>
<td>Endometrioid carcinoma</td>
<td>02</td>
<td>14.28</td>
</tr>
<tr>
<td>Papillary adenocarcinoma</td>
<td>06</td>
<td>42.85</td>
</tr>
<tr>
<td>Mucinous papillary cystadenocarcinoma</td>
<td>01</td>
<td>07.14</td>
</tr>
<tr>
<td>Dysgerminoma</td>
<td>01</td>
<td>07.14</td>
</tr>
<tr>
<td>Yolk sac tumour</td>
<td>01</td>
<td>07.14</td>
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</tbody>
</table>
Table 2: Benign ovarian tumour

<table>
<thead>
<tr>
<th>Tumour Type</th>
<th>Number</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mucinous cystadenoma</td>
<td>12</td>
<td>13.79</td>
</tr>
<tr>
<td>Papillary cystadenoma mucinous</td>
<td>03</td>
<td>03.44</td>
</tr>
<tr>
<td>Serous cyst</td>
<td>13</td>
<td>14.94</td>
</tr>
<tr>
<td>Simple cyst</td>
<td>15</td>
<td>17.24</td>
</tr>
<tr>
<td>Serous cystadenoma</td>
<td>16</td>
<td>18.39</td>
</tr>
<tr>
<td>Papillary serous cystadenoma</td>
<td>02</td>
<td>02.29</td>
</tr>
<tr>
<td>Fibroma</td>
<td>01</td>
<td>01.14</td>
</tr>
<tr>
<td>Haemorrhagic cyst</td>
<td>11</td>
<td>12.64</td>
</tr>
<tr>
<td>Corpus luteal cyst</td>
<td>03</td>
<td>03.44</td>
</tr>
<tr>
<td>Para ovarian cyst</td>
<td>02</td>
<td>02.29</td>
</tr>
<tr>
<td>Benign cystic lesion</td>
<td>04</td>
<td>04.59</td>
</tr>
<tr>
<td>Follicular cyst</td>
<td>02</td>
<td>02.29</td>
</tr>
<tr>
<td>Benign cystic teratoma</td>
<td>03</td>
<td>03.44</td>
</tr>
<tr>
<td>Total</td>
<td>87</td>
<td>100.00</td>
</tr>
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</table>

Table 3: Accuracy of CA 125 (IU/ml) in comparison to histopathology

<table>
<thead>
<tr>
<th></th>
<th>CA 125 &gt;=35</th>
<th>CA 125 &lt;35</th>
<th>Total(n=70)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Malignant</td>
<td>07</td>
<td>03</td>
<td>10</td>
</tr>
<tr>
<td>Benign</td>
<td>09</td>
<td>51</td>
<td>60</td>
</tr>
<tr>
<td>Total</td>
<td>16</td>
<td>54</td>
<td>70</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>Sensitivity (%)</th>
<th>Specificity (%)</th>
<th>PPV (%)</th>
<th>NPV (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>70</td>
<td>85</td>
<td>43.75</td>
<td>94.44</td>
<td></td>
</tr>
</tbody>
</table>

P<0.001
Table-4: Accuracy of RMI index in comparison to histopathology

<table>
<thead>
<tr>
<th>RMI index</th>
<th>Malignant</th>
<th>Benign</th>
<th>Total (n=68)</th>
</tr>
</thead>
<tbody>
<tr>
<td>&gt;=200</td>
<td>08</td>
<td>03</td>
<td>11</td>
</tr>
<tr>
<td>&lt;200</td>
<td>04</td>
<td>53</td>
<td>57</td>
</tr>
<tr>
<td>Total</td>
<td>12</td>
<td>56</td>
<td>68</td>
</tr>
</tbody>
</table>

P<0.001

<table>
<thead>
<tr>
<th>Sensitivity (%)</th>
<th>Specificity (%)</th>
<th>PPV (%)</th>
<th>NPW (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>66.6</td>
<td>94.64</td>
<td>72.72</td>
<td>92.98</td>
</tr>
</tbody>
</table>

Table-5: Comparison of ovarian crescent sign with histopathological report

<table>
<thead>
<tr>
<th>Ovarian crescent sign</th>
<th>Malignant</th>
<th>Benign</th>
<th>Total (n=68)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Malignant</td>
<td>01</td>
<td>39</td>
<td>40</td>
</tr>
<tr>
<td>Benign</td>
<td>08</td>
<td>20</td>
<td>28</td>
</tr>
</tbody>
</table>

P=0.002

In the present study amongst the total 21 women who had menopausal status only 2 women had shown malignant lesions on histopathological examination, while 19 of the women had benign lesions on histopathological examination. In this study, 70 women who had CA 125 levels were compared with the histopathological reports, amongst10 women who had malignant lesion on histopathological examination 07 women had CA 125 >= 35IU/ml while 03 women had CA 125 < 35IU/ml. Amongst 60 women who had benign lesions on histopathological examination CA 125 >= 35 IU/ml was found in 09 women only while rest 51 women had CA 125 < 35 IU/ml. In the present study 68 women in whom RMI index was calculated were compared with histopathological report, it was found that out of the 12 malignant lesions on histopathology, 8 had RMI Score>=200 while 4 women had RMI <200. Amongst 56 women who had benign lesions on histopathological report, 03 women had RMI >= 200 while 53 women had RMI < 200. The sensitivity and specificity of RMI in predicting malignant lesions as compared to histopathological report was 66.6% and 94.64% respectively.

In the study it was found that amongst 68 women, in whom ovarian crescent sign was studied 1 women had presence of ovarian crescent sign was found to have malignant lesion on HPR. Amongst 09 women who had malignant lesion on HPR, ovarian crescent sign was absent in 08 women while it was present in 01 woman only. This is in agreement with the literature which states that ovarian crescent sign is usually absent in malignant lesion. Amongst 59 women who had benign lesion on histopathological report, ovarian crescent sign was present in 39 women and was absent in 20 women.

DISCUSSION

A pelvic mass is one of the most frequent indications for referral to Gynaecologists. Diagnosis of ovarian tumours can be difficult due to variety of pathological conditions that can affect the ovaries and present with similar clinical manifestations. Knowledge of morphology and age specific characteristics can help refine the diagnosis. Our hospital is a tertiary care hospital where patients are referred from the adjoining and far flung areas. As it is a charitable hospital, a variety of Gynaecological diseases including malignancies are frequently seen. Thus, the present study was aimed to know the incidence and to study the clinicopathological features of ovarian tumours at KLE’s Dr. Prabhakar Kore Hospital and Medical Research centre, Belgaum.

In this study of 119 women, the commonest age group was 41 to 50 years (25.21%) followed by 21 to 30 years (24.36%). The mean age was found to be 40.60 years. These results were in agreement with the findings in literature stating that, the ovarian tumours can occur at any age but their peak incidence is in the reproductive age group [3, 15]. However, it was interesting to note very low frequency of early menarche, late menopause, nulliparity and advanced age at first child birth. Most of the women were multiparous and most of them had lactated in the present study.

In the present study of 119 patients, 92.43% women presented with pain abdomen, 29.41% presented with vaginal bleeding while only 12.60% had urinary complaints. Amongst signs 86.55% of women had bulky uterus while only 8.40% of women had ascites. With regard to obstetric history, most (83.19%) of the women reported were multiparous while only 6.72% were multigravida. In the present study 83.25% of women were premenopausal while 17.64% women were postmenopausal. The serum CA 125 levels were


<35 IU/ml in 77.14% while in 22.85% of women had serum CA 125 >35 IU/ml. According to a study it was found that CA 125 cannot adequately be characterized as a screening test due to the presence of overall low incidence of ovarian cancer in general population and the risk of false positive result [60, 63].

In the present study, the commonest benign lesion was found to be serous cystadenoma 18.39% (16) followed by simple cyst 17.24% (15), followed by serous cyst 14.94% (13) out of the total 87 patients who had histopathological report showing benign lesions. The commonest malignant lesion was found to be papillary adenocarcinoma 28.57% (04), followed by endometrioid carcinoma and granulosa tumour 14.28% (02) each. In the present study, RMI was found to be <200 in 83.82% of women and in 16.17% of women it was >200. The sensitivity and specificity of RMI index in detecting malignant lesions is 66.6% and 94.64% respectively. The data available from this study can help us in recognizing the pattern of ovarian tumours prevalent. Whether malignant tumours arise de novo or the benign tumour transforms into malignant is the subject of ongoing research and debate. Therefore, based on the results of this study it is evident that early diagnosis is crucial to help in decreasing morbidity and mortality among these patients.

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