Evaluation of risk of malignancy index in adnexal masses at a tertiary hospital: a prospective study

Ritanjali Behera, Paramita Pradhan*, Bharati Misra

Department of Obstetrics and Gynecology, M. K. C. G. Medical College, Berhampur, Odisha, India

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*Correspondence:
Dr. Paramita Pradhan,
E-mail: drparamitapradhan@gmail.com

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ABSTRACT

Background: The discrimination between benign and malignant adnexal masses is important in deciding clinical management and optimal surgical planning. The aim of the study was to evaluate the effectiveness of risk of malignancy index (RMI) to identify cases with high potential of ovarian malignancy at a tertiary hospital.

Methods: This prospective study was conducted over a period of two years from September 2017 to August 2019 at obstetrics and gynecology department of M. K. C. G. Medical College and Hospital, Berhampur. A total case of 130 patients with adnexal masses who underwent surgical treatment were included as histopathological report was taken as gold standard to calculate accuracy of RMI.

Results: Of the total masses, 85 (65.4%) were benign and 45 (34.6%) were malignant. The mean age of patients was 41.03±14 years. The best cut off value for the RMI-3 was 225 with highest area under the ROC curve 87%, sensitivity of 75.55%, specificity of 98.82%, PPV of 97.14%, NPV of 88.42% and an accuracy of 90.76%.

Conclusions: The present study demonstrated that RMI was a reliable method in detecting malignant ovarian tumors. The RMI is a simple and practically applicable tool in preoperative discrimination between benign and malignant adnexal masses in non-specialized gynecologic departments, particularly in developing countries.

Keywords: Adnexal masses, CA-125, Ovarian tumors, Risk of malignancy index

INTRODUCTION

Adnexal masses are a common cause for admission of patients to gynaecology clinics, and one of the most common reasons for referral to gynaecologic oncology departments for possibility of uterine or ovarian malignancies. The most prevalent type adnexal masses is ovarian masses, which vary from benign cysts to malignant ovarian cancers. Ovarian cancer ranks third after cervical and uterine cancer among gynecological malignancies. It also has worst prognosis and the highest mortality rate. Ovarian cancer constitutes 3rd most common cancer and contributes to about 6% of total cancer cases among the Indian women. The mean AAR (age adjusted rate) was observed to be 5.3 per 100,000 populations. About 50% of the total cases occurs between 45-65 years of age. The age specific incidence rates increase sharply with every ten years rise after the age of 35 years. The factors associated with an increased risk include older age, race (white), nulliparity, and family history of ovarian, endometrial, or breast cancer.

Ovarian cancer is often asymptomatic or present with a variety of vague symptoms like pelvic or abdominal pain, bloating, poor appetite, urinary urgency. So, the high mortality rate of ovarian cancer is due to asymptomatic and indolent growth of the tumor, delayed onset of symptoms, and lack of proper screening that results in its diagnosis in the advanced stages.

The goal of evaluation of adnexal masses is to discriminate between benign and malignant which helps...
in deciding clinical management and surgical planning. Early identification of ovarian carcinomas and referral to a gynaeco-oncologist can facilitate accurate staging of the disease and optimal cyto-reductive treatment, enhancing patient survival. Several diagnostic methods for adnexal masses have been reported, such as abdominal and transvaginal ultrasonography, three-dimensional ultrasound, color doppler ultrasonography and tumor markers. However, none of these methods used individually has shown significantly better performance in detecting malignant tumors from clinically restricted ovarian masses. Treatment efficiency in patients with ovarian cancers could be increased by standardization of preoperative evaluation. A formula-based scoring system known as risk of malignancy index (RMI) was introduced by Jacobs et al. in 1990, which was termed RMI. It is a product of ultrasound findings (U), the menopausal status (M) and serum CA-125 levels. (RMI = U × M × CA-125).

RMI was modified in 1996 by Tingulstad et al, as RMI 2 and again in 1999 known as RMI 3.8 The difference between the three indices lies in the different scoring of ultrasound findings and menopausal status. The three versions of RMI have been confirmed retrospectively and prospectively in different clinical studies, where a cut off value of 200 for RMI revealed the best discrimination between benign and malignant adnexal mass, because of its high sensitivity and specificity levels. Subsequently, RMI 4 was introduced by Yamamoto et al in 2009, which included tumor size as an additional parameter.

The objective of the present study was to evaluate the performance of RMI 3 in preoperative discrimination of benign and malignant adnexal masses and its applicability in daily clinical practice.

METHODS

The study was a prospective study, carried out in all patients admitted with adnexal masses to the obstetrics and gynecology department of M.K.C.G Medical College, Berhampur, India between September 2017 to August 2019.

A total of 130 patients were studied and data related to age, parity, menstrual history, family history of cancer, symptoms at diagnosis were abstracted.

Inclusion criteria

- Only women who underwent surgical treatment were included in this histopathological examination was taken as gold standard to calculate the accuracy of RMI.

Exclusion criteria

- Women receiving chemotherapy due to ovarian cancers, masses arising from urinary tract and gastrointestinal tract, ectopic pregnancy were excluded from the study.

Detailed history, pelvic and physical examination, serum CA 125 levels, abdominal ultrasound findings and menopausal status of all cases were recorded. Ultrasound scan was performed by expert radiologists.

The modified RMI (RMI 3) for each woman was calculated using the formula:

\[ \text{RMI} = U \times M \times \text{serum CA 125} \]

The ultrasound findings were evaluated and one point was given for each: multilocularity, bilaterality, presence of solid areas, presence of ascitis or presence of intra-abdominal metastases. A zero or one point gives U = 1 and total of 2 or more points gives U = 3. Patients with amenorrhea more than a year or who had hysterectomy and older than 50 years were described as postmenopausal women and they scored M = 3. Other patients scored M = 1. The absolute values of serum CA 125 (U/mL) was entered directly in the formula. The levels of < 35 U/mL were considered to be normal. Histopathologic diagnosis regarded as a gold standard for evaluation of results.

Statistical analysis

All statistical analysis were done using SPSS version 25 (IBM) and Microsoft Excel 2016 for windows. The t-test for the means and Chi square test was used to compare the differences in distribution of age, menopausal status, ultrasonographic score and other discrete variables. A probability value of p < 0.05 was considered to be statistically significant. The sensitivity, specificity, positive (PPV), and negative (NPV) predictive values with reference to the presence of malignant and benign disease were calculated. Receiver operating characteristics (ROC) curves of RMI were plotted to determine the appropriate cut-off value for discriminating between benign and malignant adnexal masses.

RESULTS

According to the histopathological examination of surgical specimens, 85 (65.4%) were benign and 45 (34.6%) were malignant. The most frequent benign conditions included serous cystadenoma (n = 25) and dermoid cyst (n = 85). Majority of malignant tumors were epithelial origin with predominant types being papillary serous cystadenocarcinoma (n = 15) (Table 1). Most of the women presented with age group of 40-59 years i.e. 57 (43.8%) (Table 2).

The distribution of age, menopausal status, ultrasound score, CA125 levels and RMI are summarized in Table 3. Mean age of patients with malignant adnexal mass (47.89±14.12 years) was significantly higher than mean...
age of patients with benign adnexal mass (37.41±12.66 years) with p value < 0.001.

Table 1: Distribution of adnexal masses according to histopathology.

| Benign masses             | Number | %    |
|---------------------------|--------|------|
| Serous cystadenoma        | 25     | 19.2%|
| Papillary serous cystadenoma | 8       | 6.2% |
| Serous cystadenofibroma   | 1      | 0.8% |
| Mucinous cystadenoma      | 12     | 9.2% |
| Papillary mucinous cystadenoma | 5   | 3.8% |
| Dermoid cyst              | 17     | 13.1%|
| Chocolate cyst            | 8      | 6.2% |
| Simple serous cyst        | 5      | 3.8% |
| Corpus luteal cyst        | 1      | 0.8% |
| Fibroma                   | 2      | 1.5% |
| Fibrothecoma              | 1      | 0.8% |
| Total                     | 85     | 65.4%|

Malignant masses

| Malignant masses               | Number | %    |
|--------------------------------|--------|------|
| Serous cystadenocarcinoma      | 6      | 4.6% |
| Papillary serous cystadenocarcinoma | 15   | 11.5%|
| Mucinous cystadenocarcinoma    | 7      | 5.4% |
| Papillary mucinous cystadenocarcinom | 4   | 3.1% |
| Dysgerminoma                  | 3      | 2.3% |
| Granulosa cell tumor          | 3      | 2.3% |
| Yolk sac tumor                | 1      | 0.8% |
| Krukenberg tumor              | 3      | 2.3% |
| Endometroid adenocarcinoma    | 1      | 0.8% |
| Mmmt (carcinosarcoma)         | 2      | 1.5% |
| Total                         | 45     | 34.6%|

Table 2: Age distribution of patients with adnexal masses.

| Age (mean±SD) | Benign (85) | Malignant (45) | Total (130) | p value |
|---------------|-------------|----------------|-------------|---------|
| < 20          | 3 (3.5%)    | 5 (11.1%)      | 8 (6.2%)    |         |
| 20-39         | 47 (55.3%)  | 4 (8.9%)       | 51 (39.2%)  |         |
| 40-59         | 30 (35.3%)  | 27 (60%)       | 57 (43.8%)  |         |
| ≥ 60          | 5 (5.8%)    | 9 (31.1%)      | 14 (10.8%)  |         |

Premenopausal patients predominate in our study with 84 (64.6%) cases, out of which 68 had benign and 16 had malignant diseases. 46 (35.4%) were in postmenopausal group, of which 17 had benign and 29 had malignant diseases. Significantly more postmenopausal women had malignant disease than premenopausal women (p < 0.001).

Majority of women presented with pain abdomen (55.4%) followed by mass abdomen with pain abdomen (46.2%). Analysis of 130 patients with ultrasound features revealed that presence of solid areas, presence of ascitis and metastasis showed significant correlation with p < 0.05. Bilaterality and multicollarity in predicting malignancy failed to be proved significant in our study with p value > 0.05. 62.3% (81) cases had an ultrasound score of 1, of which 72 had benign, while 9 had malignant diseases. Ultrasound score of 3 was statistically significant for malignant masses (p < 0.001).

Table 3: Distribution of cases according to age, USG score, menopausal status, serum CA125 levels and RMI.

| Variables                        | Benign (85) | Malignant (45) | Total (130) | p value |
|----------------------------------|-------------|----------------|-------------|---------|
| Age (mean±SD)                    | 37.4±12.66  | 47.9±14.12     | 41.0±14.05  | p < 0.001|
| Menopausal status                |             |                |             |         |
| Premenopausal                     | 68 (80%)    | 16 (35.6%)     | 84 (64.6%)  | p < 0.001|
| Postmenopausal                    | 17 (20%)    | 29 (64.4%)     | 46 (35.4%)  |         |
| Sonographic morphology            |             |                |             |         |
| Bilateral                        | 7 (53.8%)   | 6 (46.2%)      | 13 (10%)    | p = 0.357|
| Multilocularity                   | 34 (58.6%)  | 24 (41.4%)     | 58 (44.6%)  | p = 0.146|
| Solid Areas                       | 19 (30.2%)  | 44 (69.8%)     | 63 (48.4%)  | p < 0.001|
| Ascitis                           | 5 (16.1%)   | 26 (83.9%)     | 31 (23.8%)  | p < 0.001|
| Metastasis                        | 0 (0%)      | 4 (100%)       | 4 (3.1%)    | p < 0.005|
| USG score (U)                     |             |                |             |         |
| 1                                | 72 (84.7%)  | 9 (20%)        | 81 (62.3%)  | p < 0.001|
| 3                                | 13 (15.3%)  | 36 (80%)       | 49 (37.7%)  |         |
| CA 125                           |             |                |             |         |
| < 35 U/mL                         | 41 (48.2%)  | 11 (24.4%)     | 52 (40%)    | p = 0.008|
| ≥ 35 U/mL                         | 44 (51.8%)  | 34 (75.6%)     | 78 (60%)    |         |
| Serum CA-125 (U/mL) (Mean±SD)     | 60.9±127.45 | 277.6±427.57   | 135.9±289.20| p < 0.001|
| RMI (mean±SD)                     | 111.9±375.27| 1641.0±1933.38| 633.5±1496.04| p < 0.001|
| RMI ≥ 225                         | 1 (1.2%)    | 35 (26.9%)     | 36 (27.7%)  | p < 0.001|
| RMI < 225                         | 84 (98.8%)  | 11 (24.4%)     | 95 (73.1%)  |         |
Table 4: Sensitivity, specificity, PPV, NPV, odds ratio and AUC for rmi-3 at different cut off points.

| Cut off | Sensitivity (%) | Specificity (%) | PPV (%) | NPV (%) | LR | OR | AUC | Youden index | Accuracy (%) |
|---------|-----------------|-----------------|---------|---------|----|----|-----|--------------|---------------|
| 150     | 77.77           | 87.05           | 76.08   | 88.09   | 55.778 | 23.545 | 0.824 | 0.64 | 88.46        |
| 200     | 75.5            | 96.4            | 91.89   | 88.17   | 79.27    | 84.485 | 0.86  | 0.79 | 86.92        |
| 225.33  | 75.55           | 98.82           | 97.14   | 88.42   | 90.521    | 259.63 | 0.87  | 0.85 | 90.76        |
| 250     | 68.8            | 98.8            | 96.875  | 85.71   | 78.426    | 186.0 | 0.839 | 0.81 | 88.46        |
| 300     | 64.44           | 98.82           | 96.66   | 84.0    | 71.006    | 152.25 | 0.816 | 0.80 | 88.46        |

Table 5: Diagnostic performance of criteria evaluated.

| Criteria                  | Sensitivity (%) | Specificity (%) | PPV (%) | NPV (%) | Accuracy (%) |
|---------------------------|-----------------|-----------------|---------|---------|---------------|
| RMI ≥ 225                 | 75.55           | 98.82           | 97.14   | 88.42   | 90.76         |
| CA-125 ≥ 35               | 75.55           | 48.23           | 43.58   | 78.84   | 57.69         |
| USG score 3               | 80              | 84.7            | 73.5    | 88.9    | 83.1          |
| Menopause score 3         | 64.4            | 80              | 63.04   | 81      | 74.6          |

Figure 1: ROC curve for RMI-3 in differentiating between benign and malignant adnexal masses.

The mean value of CA 125 was 277.63±427.57 U/mL for malignant adnexal masses compared to 60.99±127.45 U/mL for benign masses (p < 0.001). CA 125 at cut off level of 35 U/mL gave sensitivity of 75.6%, specificity of 48.2%, positive predictive value of 43.6% and negative predictive value of 78.8%.

The best performance in our study obtained for RMI-3 was at the cut-off point 225 with highest area under the ROC curve i.e., AUC = 87%, sensitivity of 75.55%, specificity of 98.82%, PPV of 97.14%, NPV of 88.42% and an accuracy of 90.76% (Table 4). Taking into account the best obtained cut-off point for RMI-3, 1 case was false positive (fibroma) and 34 cases were true positive (RMI ≥ 225 malignant tumors) while 84 cases were true negative and 11 cases were false negative (RMI < 225 malignant tumor); (3 cases were dysgerminoma, 3 cases were granulose cell tumor, 1 yolk sack tumor, 2 cases were papillary mucinous cystadenocarcinoma, 1 mucinous cystadenocarcinoma and 1 serous cystadenocarcinoma).

The diagnostic performance of RMI-3 > 225, against CA-125 level > 35, ultrasound score of 3 and menopausal score of 3 is compared in Table 5. Among the criteria RMI score > 225 has highest sensitivity, specificity, PPV, NPV and diagnostic accuracy, when compared with individual parameters. Among the individual parameters, USG score of 3 has the highest sensitivity, specificity, PPV, NPV and diagnostic accuracy (80%, 84.7%, 73.5%, 88.9% and 83.1% respectively).

DISCUSSION

The present study has established the effectiveness of RMI in assessment of women with adnexal masses, particularly in low-resource settings. Some of the potential advantages of RMI include rapid triage of patients and timely referral to gynaec oncologists, thereby avoiding suboptimal primary surgeries at local hospitals or peripheral centres. In the present study commonest age group was 40-59 years which was consistent with previous studies, which showed that the disease was more prevalent in this age group (mean 50 years). In our study mean age of benign group was 37.41±12.66 years and mean age of malignant group was 47.89±14.12 years which coincides with study of Ashrafgangooei et al and Simsek et al who revealed mean age in benign as 37.0±8.79, 35.23±10.87 and 50.8±12.9, 50.78±13.39 in malignant group respectively. In our study mean age of benign group was 37.41±12.66 years and mean age of malignant group was 47.89±14.12 years which coincides with study of Ashrafgangooei et al and Simsek et al who revealed mean age in benign as 37.0±8.79, 35.23±10.87 and 50.8±12.9, 50.78±13.39 in malignant group respectively. In our study mean age of benign group was 37.41±12.66 years and mean age of malignant group was 47.89±14.12 years which coincides with study of Ashrafgangooei et al and Simsek et al who revealed mean age in benign as 37.0±8.79, 35.23±10.87 and 50.8±12.9, 50.78±13.39 in malignant group respectively. In our study mean age of benign group was 37.41±12.66 years and mean age of malignant group was 47.89±14.12 years which coincides with study of Ashrafgangooei et al and Simsek et al who revealed mean age in benign as 37.0±8.79, 35.23±10.87 and 50.8±12.9, 50.78±13.39 in malignant group respectively. The chances of ovarian malignancy increases with the increasing age.

A total 64.4% of malignancies occurred in postmenopausal women and 35.6% among premenopausal women. This was in agreement with Rao JH et al, and Kumari N et al, showing similar incidence rates and preponderance in postmenopausal patients.
Table 6: Comparison of RMI from the various previous studies with the present study.

| Study               | Year | Number | Sensitivity | Specificity | PPV  | NPV  |
|---------------------|------|--------|-------------|-------------|------|------|
| Jacob et al         | 1990 | 143    | 85.4        | 96.6        | -    | -    |
| Davies et al        | 1993 | 124    | 87          | 89          | -    | -    |
| Tingulstad et al    | 1996 | 173    | 71          | 96          | 89   | 88   |
| Tingulstad et al    | 1999 | 365    | 71          | 92          | 69   | 92   |
| Morgante et al      | 1999 | 124    | 58          | 95          | 78   | 87   |
| Manjunath et al     | 2000 | 152    | 73          | 91          | 93   | 67   |
| Ma et al            | 2003 | 140    | 87.3        | 84.4        | 82.1 | 89   |
| Torres et al        | 2003 | 158    | 73          | 86          | -    | -    |
| Anderson et al      | 2004 | 180    | 70.6        | 87.7        | 66.1 | 89.8 |
| Obeidat et al       | 2005 | 100    | 90          | 89          | 96   | 78   |
| Leelahakorn et al  | 2007 | 175    | 88.6        | 90.7        | 70.5 | 97   |
| Ulusoy et al        | 2010 | 296    | 71.7        | 80.5        | 67.3 | 83.6 |
| Van Den Akker et al| 2011 | 548    | 81          | 85          | 48   | 96   |
| Bouzari et al       | 2014 | 182    | 91.3        | 88          | 52   | 98.58|
| JH Rao et al        | 2015 | 90     | 84          | 89          | 93   | 71   |
| M Zarchi            | 2017 | 200    | 78.95       | 58.44       | 90.08| 78.93|
| SK Dora et al       | 2017 | 126    | 72.5        | 98.2        | 98.1 | 74.7 |
| Present study (225) | 2019 | 130    | 75.55       | 98.82       | 97.14| 88.42|

Ultrasonography has been widely used for evaluation of adnexal masses. In our study, an ultrasound score of 3 had the sensitivity 80%, specificity 84.7%, positive predictive value 73.5% and negative predictive value (88.9%) which is in agreement with Vasudevan et al. We got higher sensitivity and specificity than others because USG were done by expert radiologist in majority of our cases.

Serum CA 125 level is universally used as a tumor marker for diagnosing ovarian cancer, though other gynecological pathology can also increase its levels. Simsek et al reported a sensitivity of 78.6% and specificity of 63.5% for CA-125 > 35 U/ml. Recent study by 2018 Singh S et al gave a sensitivity of 75% and specificity of 90% for CA-125 levels > 35U/ml. In our study, CA125 levels ≥ 35 U/ml had a sensitivity of 75.6%, the specificity of only 48.2%, PPV 43.6% and NPV of 78.8%. A higher prevalence of pelvic inflammatory diseases and endometriosis might have contributed to elevated CA125 levels in the majority of our patients along with fluctuation of CA125 during the menstrual cycles in premenopausal patients with adnexal masses and its more specificity for non-mucinous epithelial ovarian tumor might be the reason for its low diagnostic performance in the detection of malignant ovarian disease.

The RMI cut-offs in many studies ranged from 25 to 250 (reviewed in Geomini et al). Most studies reported an increased diagnostic accuracy and performance with an RMI cut-off of 200.6-8,13-16 Jacobs et al, studying 143 patients, reported a sensitivity of 85.4% and specificity of 96.9% for this method, with a cut-off of 200.
The present study demonstrated that RMI was a better estimate in diagnosing adnexal masses with high risk of malignancy and subsequently guiding the patients to gynecological oncology centers for suitable and effective surgical interventions compared with individual parameters of Ultrasound score CA 125 or menopausal score. RMI seems to be simple, easily applicable and available method which can be directly applied in clinical practice in non-specialized gynecologic departments.

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