Is routine gastroscopy/colonoscopy reasonable in patients with suspected ovarian cancer: A retrospective study

Guochen Liu  
Sun Yat-sen University Cancer Center

Junping Yan  
Guangdong Second Provincial General Hospital

Shanshan Long  
Sun Yat-sen University Cancer Center

Zhimin Liu  
Sun Yat-sen University Cancer Center

Haifeng Gu  
Sun Yat-sen University Affiliated Tumor Hospital: Sun Yat-sen University Cancer Center

Hua Tu (✉ tuhua@sysucc.org.cn)  
Sun Yat-sen University Affiliated Tumor Hospital: Sun Yat-sen University Cancer Center
https://orcid.org/0000-0003-4361-4961

Jundong Li  
Sun Yat-sen University Affiliated Tumor Hospital: Sun Yat-sen University Cancer Center

Research

Keywords: ovarian cancer, gastroscopy, colonoscopy, ovarian metastasis, differential diagnosis

DOI: https://doi.org/10.21203/rs.3.rs-329104/v1

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Abstract

Objective To evaluate the value of routine preoperative gastroscopy/colonoscopy in patients with suspected ovarian cancer for differential diagnosis and judgment of bowel resection.

Methods All women diagnosed with suspected ovarian cancer who underwent gastroscopy/colonoscopy before surgery in our center were retrospectively identified. Gastroscopy/colonoscopy results and clinical pathology, imaging, and surgical findings were analyzed.

Results 389 patients were included. Among them, 40 were ovarian metastasis. Compared with imaging, gastrointestinal endoscopy showed no statistical advantage in the specificity and sensitivity (99.4% vs. 99.7%, \( P = 1.0; \) 55.0% vs. 45.2%, \( P = 0.057; \) respectively). All patients with gastric/colonic cancer metastasize except for one had indicative imaging or tumor marker abnormalities. Three patients with colonic cancer metastases underwent optimal surgery and alive with no recurrence, the other 19 patients experienced palliative chemotherapy. There is no significant difference in the sensitivity of colonoscopy and imaging in predicting intestinal incision (61.5% vs. 43.8%, \( P = 0.804; \)) whereas, the latter had higher specificity (87.8% vs. 74.3%, \( P = 0.001; \)).

Conclusions For patients with suspected ovarian cancer, the incidence of gastrointestinal metastases is low, routine gastroscopy/colonoscopy before treatment is less efficient. Gastroscopy/colonoscopy has limited power to predict the need for gastrointestinal resection before ovarian cancer surgery.

Introduction

Early screening for ovarian cancer lacks effective methods, and many patients will resort to medical care because of symptomatic pelvic masses. According to the National Comprehensive Cancer Network guideline, patients with newly diagnosed pelvic masses suspected of ovarian cancer should be considered as candidates for gastroscopy/colonoscopy. This strategy is mainly due to the following considerations: First, it has been reported that about 3.2%~7.0% of ovarian tumors are metastasized from the stomach or colon, namely the Krukenberg tumor. [1–3]. For these patients, therapeutic decisions should be made by surgical oncologists rather than gynecologists. Second, ovarian cancer is prone to disseminate in the abdominal cavity, in which the digestive tract is most vulnerable, and preoperative evaluation is quite important. According to the literature, approximately 20% of patients underwent gastrointestinal procedures during cytoreductive surgery for ovarian cancer [4–7]. Findings from gastrointestinal endoscopy will allow more sufficient preoperative preparation for these patients. Mucosa involvement and loss of elasticity are signs of tumor invasion that may require bowel resection.

However, gastrointestinal endoscopy also has several disadvantages, including causing discomfort to the patients, increased medical costs, delay in treatments, and the risks of gastrointestinal perforation, bleeding, and cardiovascular and cerebrovascular accidents, etc. Moreover, employing patient symptoms, physical examination, tumor marker examination, preoperative imaging evaluation, and puncture pathology, it is also possible to indirectly determine the source of the tumor or whether there is
gastrointestinal involvement. The small intestine is also frequently involved in ovarian cancer, for which colonoscopy has limited detection capability[6]. In clinical practice, we found that few patients had changed their diagnoses or established treatment strategies due to the findings from gastrointestinal endoscopy. Therefore, it is questionable whether routine gastrointestinal endoscopy is a rational strategy for patients with suspected ovarian cancer at initial diagnosis.

In our center, gastroscopy/colonoscopy has been routinely performed in the majority of patients who were suspected to have ovarian cancer. Based on a large number of screened patients, we conducted a retrospective study to evaluate the rationality of routine gastroscopy/colonoscopy before treatment for patients suspected of ovarian cancer.

Methods

The study was approved by the Research Ethics Committee of the Sun Yat-sen University Cancer Center. We retrospectively collected the information of patients diagnosed with suspected ovarian cancer by imaging examinations who underwent gastroscopy/colonoscopy before treatment in our center from November 1, 2016, to October 29, 2019. Pelvic mass biopsy or surgical pathology results in our hospital were required. Patients with a history of gastrointestinal cancer or ovarian cancer, or who had a definite pathological diagnosis before the gastrointestinal examination, or did not undergo imaging (BUS/CT/MRI/PET-CT) examination were excluded. The endoscopy system was searched to obtain the gastroscopy/colonoscopy results. Clinicopathological information includes age, symptoms, physical examination, tumor marker values, preoperative imaging results, surgical records, and postoperative pathological results were obtained from the hospital information system (HIS) system. We reviewed the patients’ imaging reports in the PACS system to determine the possible source of pelvic masses and whether it involves the intestine. If the source of the tumor cannot be determined, it was recorded as unknown.

We used SPSS software (SPSS Inc., Chicago, IL, version 19) for data analysis. The Mann-Whitney U rank-sum test and the Wald chi-square (χ²) test was used for comparison of two sets of quantitative data and categorical parameters, respectively. The McNemar test was used to compare the difference between specificity and sensitivity. All P values were two-sided, and P values of less than .05 were considered statistically significant.

Results

After screening 1851 patients (Fig. 1), a total of 389 eligible patients were included in the study, of which 302 patients had gastroscopy and colonoscopy at the same time, 46 had gastroscopy only, and 41 had colonoscopy only. The patients' basic information is shown in Table 1.
Table 1
Demographic and clinical-pathological characteristics

| Variables             | Group                                      |          |          |          |          |
|-----------------------|--------------------------------------------|----------|----------|----------|----------|
|                       | Primary ovarian tumor (n = 339)            |          |          |          |          |
|                       | No.(%)                                     |          |          |          |          |
| Age, y                | 0.337†                                     |          |          |          |          |
| Median                | 51                                         | 48       |          |          |          |
| Range                 | 18 ~ 85                                    | 21 ~ 78  |          |          |          |
| Side                  | 0.002*                                     |          |          |          |          |
| Unilateral            | 219(64.6)                                  | 16(40.0) |          |          |          |
| Bilateral             | 120(35.4)                                  | 24(60.0) |          |          |          |
| Digestive symptoms    | 0.074*                                     |          |          |          |          |
| Negative              | 143(42.2)                                  | 11(27.5) |          |          |          |
| Positive              | 196(57.8)                                  | 29(72.5) |          |          |          |
| CA125                 |                                            |          |          |          |          |
| <=35 U/ml             | 63(18.6)                                   | 15(37.5) | 0.003*   |          |          |
| >35 U/ml              | 271(79.9 )                                 | 23(57.5) |          |          |          |
| Unknown               | 5(1.5)                                     | 2(5.0)   |          |          |          |
| CEA                   |                                            |          |          |          |          |
| <=5 ng/ml             | 284(83.8)                                  | 17(42.5) | < 0.001* |          |          |
| >5 ng/ml              | 43(12.7)                                   | 23(57.5) |          |          |          |
| Unknown               | 12(3.5)                                    | 0(0)     |          |          |          |
| CA199                 |                                            |          |          |          |          |
| <=35 U/ml             | 241(71.1)                                  | 21(52.5) | 0.020*   |          |          |
| >35 U/ml              | 88(26.0)                                   | 17(42.5) |          |          |          |
| Unknown               | 10(2.9)                                    | 2(5.0)   |          |          |          |

†P values were calculated using a two-sided Mann-Whitney U rank-sum test.

*P values were calculated using a two-sided Wald χ² test.
### Variables

|                  | Group                                      |         |         |      |
|------------------|--------------------------------------------|---------|---------|------|
|                  | Primary ovarian tumor (n = 339)            | Ovarian metastatic tumor (n = 40) |      |
|                  | No.(%)                                     | No.(%)  |         |      |
| HE4 <=90 pmol/L  | 103(30.4)                                  | 23(57.5)| < 0.001*|
| HE4 >90 pmol/L   | 193(56.9)                                  | 11(27.5)|         |
| HE4 Unknown      | 43(12.7)                                   | 6(15.0) |         |
| CA125/CEA <=25   | 81(23.9)                                   | 24(60.0)| < 0.001*|
| CA125/CEA >25    | 246(72.6)                                  | 14(35.0)|         |
| CA125/CEA Unknown| 12(3.5)                                    | 2(5.0)  |         |

† *P* values were calculated using a two-sided Mann-Whitney U rank-sum test.

* *P* values were calculated using a two-sided Wald $\chi^2$ test.

Of the 348 patients who had a gastroscopy, 13 had pathologically confirmed primary malignant tumors of the stomach (11 biopsies confirmed gastric cancer, 2 cases were gastric lymphoma), 45 were normal, 37 had polypus confirmed by biopsy, 6 had external pressure lesions, 7 cases had gastric inflammation with gastric ulcer, 10 cases had gastritis and polyps at the same time, 230 cases had stomach inflammation (of which 220 cases had chronic non-atrophic gastritis, accounting for 95.7%).

Of the 343 patients who underwent colonoscopy, 10 cases had biopsy-confirmed bowel cancer, 162 had no abnormalities, 78 had polyps, 21 had inflammation and polyps, 72 had extrinsic compression or infiltration (of which 2 had failed colonoscopy due to external pressure of the tumor). It is worth noting that of the 2 patients with multiple intestinal polyps, 1 patient was diagnosed with FAP (familial adenomatous polyposis) and the other was considered P-J (Peutz-Jeghers) syndrome.

As confirmed by biopsy or postoperative pathology (Fig. 1), among all patients with the initial diagnosis of suspected ovarian cancer, 277 cases were ovarian primary malignant tumors, 25 cases were ovarian primary borderline tumors and 37 cases were ovarian primary benign tumors, 40 cases were ovarian metastatic tumors (11 gastric cancer, 9 colonic cancer, 8 appendix mucinous tumor, 2 gastric lymphomas, 2 endometrial cancer, 2 pancreatic cancer, 1 cholangiocarcinoma, 1 peritoneum Malignant mesothelioma, 1 small intestinal stromal tumor, 1 liver cancer, 1 cervical cancer, 1 unclear primary pathology), 4 cases were multiple primary tumors (1 ovarian cancer with galbladder cancer, 1 ovarian cancer sarcoma with appendix mucinous tumor, 1 sigmoid colon cancer with ovarian cancer, and 1 lymphoma with ovarian
teratoma), 6 cases were ovarian non-neoplastic lesions (2 inflammatory lesions, 2 subuterine fibroids, 1 small intestinal stromal tumor, and 1 ovarian tuberculosis).

Compared with primary ovarian tumors, patients with ovarian metastases are more bilateral lesions (60% vs. 35.4%, \( P = 0.002 \)), ovarian cancer indicators CA125 (37.5% vs. 18.6%, \( P = 0.003 \)) and HE4 are mostly normal (57.5% vs. 30.4%, \( P < 0.001 \)), gastrointestinal cancer indicators CEA (57.5% vs. 12.7%, \( P < 0.001 \)) and CA199 (42.5% vs. 26%, \( P = 0.020 \)) are more abnormal, and the ratio of CA125/CEA less than 25 is higher (60.0% vs. 23.9%, \( P < 0.001 \)). Besides, patients with ovarian metastases are more likely to have gastrointestinal symptoms, although there is no statistical difference (72.5% vs. 57.8, \( P = 0.074 \)). Among all the symptoms, abdominal distension and abdominal pain are the most common (24/40). (Table 1)

We analyzed the diagnostic efficacy of gastrointestinal symptoms, tumor markers, unilateral and bilateral accessory lesions, imaging examination, and gastroscopy/colonoscopy for metastatic ovarian tumors. (Table 2) Gastrointestinal endoscopy and imaging had a high diagnostic efficacy (94.9% vs. 94.4%, respectively) for ovarian metastases, there was no statistical difference between the specificity and sensitivity of the two methods (99.4% vs. 99.7, \( P = 1.0 \); 55.0% vs. 45.2%, \( P = 0.057 \)). We used the ROC curve to separately analyze the discrimination of different tumor markers for ovarian metastases (Fig. 2). The area under the curve (AUC) in descending order were: HE4(0.756), CA125/CEA(0.730), CEA(0.642), CA125(0.629), CA199(0.602). Different from previous research[8], we found that when CA125/CEA = 10.57, the Youden Index value was the largest, and the specificity and sensitivity were 87.83% and 55.26% respectively.

Figure S1 displays the examination results of 13 patients with gastric metastases and 9 colonic metastases. Except for one patient with gastric cancer who has no corresponding imaging or tumor marker abnormalities (Case 2), all the remaining patients had an indication tumor maker or imaging.

| Table 2. The diagnostic value of ovarian metastatic carcinoma by different criterias |
|---------------------------------|------------|----------|--------|--------|--------|
| Criterion                        | Specificity (%) | Sensitivity (%) | PPV (%) | NPV (%) | DE (%)  |
| Gastroscopy / colonoscopy        | 99.4        | 55.0      | 91.7   | 95.1   | 94.9    |
| Imaging scan                     | 99.7        | 45.2      | 93.3   | 94.5   | 94.4    |
| Digestive symptoms              | 42.2        | 72.5      | 12.9   | 92.9   | 45.4    |
| CA125                           | 80.6        | 35.0      | 17.5   | 91.3   | 75.8    |
| CEA                             | 87.0        | 51.3      | 30.8   | 94.0   | 83.3    |
| CA199                           | 75.7        | 63.2      | 22.2   | 94.9   | 74.4    |
| HE4                             | 62.8        | 60.5      | 17.0   | 92.6   | 62.5    |
| Bilateral adnexal lesion        | 54.3        | 66.7      | 15.3   | 93.0   | 55.7    |
| CA125/CEA                       | 74.9        | 63.2      | 22.0   | 94.8   | 73.7    |

PPV=positive predictive value, NPV=negative predictive value, DE=diagnostic efficiency. Digestive symptoms include: abdominal distension, abdominal pain, diarrhea, constipation, vomiting, hematochezia, melena.

Imaging scan include: color ultrasound, CT, MRI, PET-CT.

Patients meet one of the following criterions were considered as ovarian metastatic carcinoma: positive gastrointestinal symptoms, CA125<35U/ml, CEA>5ng/ml, CA199>35U/ml,
HE4<90pmol/L, bilateral adnexal lesion or CA125/CEA<25.

Thirteen patients with malignant tumors of the primary stomach, including 2 cases with gastric lymphoma, all received corresponding chemotherapy after obtaining pathological evidence for metastasis by ultrasound-guided accessory tumors puncture. Three patients with colon metastases underwent surgical treatment after ruled out other sites of metastasis and received chemotherapy for colon cancer subsequently, fortunately, all of them survived without tumors at the last follow-up. The other 6 cases of colon cancer underwent ultrasound-guided puncture of the adnexal mass, after the metastasis was confirmed, they received chemotherapy according to the corresponding chemotherapy regimen. The treatment and prognosis of the patient are shown in Table 3.
Table 3
The treatment and prognosis of the 22 ovarian metastases from the stomach or colon

| Case no. | Gastroscopy/colonoscopy | Correct diagnosis | Initial primary treatment plan | Actual primary treatment | Outcome at last Surveillance | OS (months) |
|----------|-------------------------|-------------------|-------------------------------|--------------------------|-----------------------------|-------------|
| 1        | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | DOD                         | 16.7        |
| 2        | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | DOD                         | 16.3        |
| 3        | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | AWD                         | 16.4        |
| 4        | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | DOD                         | 5.2         |
| 5        | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | AWD                         | 18.5        |
| 6        | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | NED                         | 35.0        |
| 7        | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | AWD                         | 10.0        |
| 8        | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | DOD                         | 21.9        |
| 9        | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | DOD                         | 12.5        |
| 10       | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | NED                         | 19.8        |
| 11       | Gastric cancer          | Surgery           | Surgery                       | Chemotherapy              | Unknown                      | Unknown     |
| 12       | Gastric lymphoma        | Surgery           | Surgery                       | Chemotherapy              | NED                         | 3.2         |
| 13       | Gastric lymphoma        | Surgery           | Surgery                       | Chemotherapy              | AWD                         | 24.2        |
| 14       | Sigmoid colon cancer    | Surgery           | Surgery                       | Chemotherapy              | NED                         | 10.7        |
| 15       | FAP, ascending colon cancer | Surgery      | Surgery                       | Chemotherapy              | DOD                         | 28.3        |
| 16       | Ascending colon cancer  | Surgery           | Surgery                       | Surgery                   | NED                         | 5.03        |
| 17       | Sigmoid colon cancer    | Surgery           | Surgery                       | Chemotherapy              | NED                         | 10.0        |
| 18       | Ascending colon cancer  | Surgery           | Surgery                       | Surgery                   | NED                         | 17.9        |
| 19       | Sigmoid colon cancer    | Surgery           | Surgery                       | Surgery                   | NED                         | 31.9        |
| 20       | Sigmoid colon cancer    | Surgery           | Surgery                       | Chemotherapy              | NED                         | 1.3         |
| 21       | Sigmoid colon cancer    | Surgery           | Surgery                       | Chemotherapy              | Unknown                      | Unknown     |
Of the 277 patients with ovarian cancer who underwent surgery, 32 underwent colon surgery (2 right hemicolectomies, 9 sigmoidectomies, and 21 Dixon), no patient underwent gastric surgery at the same time. Table S1 shows the predictive value of gastroscopy/colonoscopy and imaging in bowel resection. If the intestinal pressure and invasion were considered as signs of intestinal resection, there was no significant difference in the sensitivity of colonoscopy and imaging to the prediction of intestinal incision (61.5% vs. 43.8%, P = 0.804), however, the imaging’s specificity is higher (87.8% vs. 74.3%, P = 0.001). It is noteworthy that the above gastrointestinal surgery mentioned does not include appendix and small bowel surgery, one patient each had preoperative imaging indicated invasion of the appendix and small intestine underwent surgery on the corresponding part.

**Discussion**

There is no clear recommendation for the application of gastroscopy/colonoscopy for patients with suspected ovarian cancer in the current guidelines. The choice of gynecologists can be very varied when treating these patients.

In our study, only 5.7% (22/389) of the patients had gastrointestinal metastases. It is worth noting that, except for one patient, the remaining patients all had imaging or tumor marker indicators. If the gastrointestinal examination is performed only on patients with imaging or tumor marker indicators, 62.2% (242/389) of patients can be spared from an unnecessary gastrointestinal examination, which means great medical cost savings. Therefore, risk-adapted gastrointestinal endoscopy may be a more reasonable strategy for patients suspected of ovarian cancer.

Although one patient got a false-negative results in a gastrointestinal examination, neither her treatment nor prognosis had been impaired. After pathological confirmation of gastrointestinal cancer, salvage treatment of gastrointestinal tumors were given. In the present study, most patients showing positivity in gastrointestinal endoscopy changed their treatment strategies and abandoned the planned surgeries, according to the current recommendations for metastatic gastrointestinal cancer. However, surgery can yet be regarded as another choice for these patients. Many studies had revealed that for patients with primary bowel cancer and ovarian metastasis, optimal cytoreductive surgery could also bring a favorable prognosis[9, 10]. Three patients in our study with bowel cancer achieved long-term survival without recurrence after surgery, which supports the above findings. Similarly, for patients with gastric cancer and

| Case no. | Gastroscopy/colonoscopy | Initial primary treatment plan | Actual primary treatment | Outcome at last Surveillance | OS (months) |
|----------|-------------------------|--------------------------------|--------------------------|-----------------------------|-------------|
| 22       | Descending colon cancer | Surgery                        | Chemotherapy             | DOD                         | 25.9        |

OS, overall survival; DOD, die of disease; AWD, alive with disease; NED, no evidence of disease

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| Case no. | Correct diagnosis | Initial primary treatment plan | Actual primary treatment | Outcome at last Surveillance | OS (months) |
|----------|-------------------|--------------------------------|--------------------------|-----------------------------|-------------|
| 22       | Descending colon cancer | Surgery                        | Chemotherapy             | DOD                         | 25.9        |
ovarian metastasis, Cheong et al. found that the removal of metastases can improve the prognosis of patients [11]. Furthermore, a Korean study validated that the removal of ovarian metastases plus palliative chemotherapy provided a better prognosis than palliative chemotherapy alone [12]. In light of the above evidences, if the surgery can achieve satisfactory resection, it will not delay the treatment of the patients or affect their prognosis. For patients who cannot obtain optimal resection, the removal of ovarian masses can also be beneficial, since these metastases were usually chemotherapy-resistant [9, 13–17]. Moreover, ovarian metastatic tumors are relatively large, with the average sizes of 9–12 cm in previous studies [2, 18]. Palliative resection of these tumors could relieve symptoms such as abdominal distension and pain [10]. In short, it is still of diagnostic and therapeutic value to perform surgery for patients with primary gastrointestinal cancer and ovarian metastases. Accordingly, the value of gastrointestinal endoscopy as a routine procedure to distinguish these patients from those with ovarian cancer is very limited.

Moreover, unlike imaging examinations, gastrointestinal endoscopy cannot find primary lesions beyond the stomach and large intestine, such as appendiceal cancer. However, studies have shown that ovarian metastases more commonly arise from the appendix than the stomach [2]. This further reduces the value of routine gastrointestinal endoscopy. In the present study, metastases originating in the stomach were more common. This discrepancy may be accredited to a higher incidence of gastric cancers in China.

When there is a pelvic mass whose origin cannot be defined, patients usually need to receive fine-needle aspiration of pelvic masses to obtain histopathological evidence. Due to the risks of needle path metastasis and the difficulty of pathological diagnosis based on small-volume tissue, surgical exploration may be a better choice, which is also necessary for patients with ovarian cancer requiring neoadjuvant chemotherapy. When a pathological diagnosis is in doubt, then gastrointestinal examination is performed to exclude gastrointestinal metastasis, which can also avoid a considerable number of patients receiving gastrointestinal endoscopy. Following the recommendations of the guidelines, obtaining accurate histological evidence before chemotherapy in such patients will prevent improper chemotherapy.

Furthermore, similar to the conclusions of previous studies [4, 5], in patients receiving cytoreductive surgery for ovarian cancer, compared with imaging, gastroscopy/colonoscopy cannot predict gastric or large intestine resection well before surgery, nor can it predict the possibility of small intestine and appendectomy. The proportion of these patients in ovarian cancer reduction surgery is not low [6].

This study is a retrospective one, which has several limitations. Firstly, the study included a small number of patients with benign and non-ovarian tumors, for whom the gastroscopy/colonoscopy is not generally necessary. Compared with patients with primary ovarian cancer, patients with metastatic ovarian tumors accounted for a significantly lower proportion in this study, which might have impaired the statistical power of the results. This is mainly due to the inclusion of consecutive unselected patients, which was designed to reflect the actual clinical practice as much as possible. Secondly, the physical examination was not included as an indicator because the results of medical examinations by different doctors are
usually inconsistent and subjective. Besides, considering that rectal cancer accounts for a higher proportion of colorectal cancer and the convenience of physical examination to determine the possibility of rectal resection, the importance of colonoscopy may be further weakened. Thirdly, whether neoadjuvant chemotherapy or satisfactory reduction during surgery had affected the proportion of bowel resection and thereby affected our evaluation on gastrointestinal endoscopy remains unclear. Due to the retrospective nature and the relatively small sample size, we did not discuss it, which requires further investigation by future studies.

Conclusion

Among patients with suspected ovarian cancer, the proportion of gastrointestinal metastases is low, and the efficiency of routine gastroscopy/colonoscopy before treatment is quite lacking. It seems more reasonable to adopt a risk-adapted gastroscopy/colonoscopy strategy based on imaging examination and tumor markers. Before ovarian cancer surgery, gastrointestinal endoscopy has limited power to predict the need for gastrointestinal resection.

Declarations

Authors' contributions:

Guochen Liu: Data curation, Investigation, Writing - original draft, Writing - review & editing. Junping Yan: Data curation, Investigation, Writing – original draft, Formal analysis. Shanshan Long: Data collection, Investigation, Validation, Methodology. Zhimin Liu: Data curation, Formal analysis. Haifeng Gu: Data curation, Formal analysis. Hua Tu: Investigation, Methodology, Project administration, Supervision, Validation, Writing - original draft, Writing - review & editing. Jundong Li: Investigation, Methodology, Project administration, Supervision, Validation, Writing - original draft, Writing - review & editing.

Funding:

This work was supported by funds from the Nature Science Foundation of China (No. 81802615). The authors declare no conflicts of interest.

Availability of data and materials:

All data generated or analyzed during this study are included in this published article.

Ethics approval and consent to participate:

This study was approved by the Ethics Committee of the Sun Yat-sen University Cancer Center ([2018]2-206 No.16). Informed consent was obtained from all participants included in the study.

Consent for publication:
Not applicable

Competing interests:

The authors declare no conflicts of interest.

Acknowledgements:

None.

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Figures
Figure 1

Flow diagram of patient selection and final diagnosis
Figure 2

The ROC curves of different tumor markers

Supplementary Files

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