Mini-laparotomy in advanced ovarian cancer

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Abstract Optimal cytoreduction (OCR) remains the gold standard treatment of ovarian cancer. Current radiological imaging has limited sensitivity and specificity in prediction of achieving OCR prior to surgery. This prospective pilot study included 50 patients with advanced ovarian carcinoma. Prior to the main laparotomy, a mini-laparotomy—just large enough to allow a hand in-was performed. A decision was then made as to whether achieving OCR is "possible", "not possible" or lastly "unsure". Formal laparotomy then followed. At the end of the formal laparotomy, cytoreduction was regarded as either "optimal" or "suboptimal" based on residual disease. Out of 45 cases where results were deemed suitable for analysis, 27 were regarded as "OCR possible", out of which OCR was achieved in 24 cases following full laparotomy. Ten were commented upon as "unsure" and only in three cases OCR was feasible. Eight were classed as "OCR not possible" and in none of these OCR was obtained. The only noted complication associated with minilaparotomy was bleeding in just three cases (6%). The sensitivity, specificity, PPV and NPV of mini-laparotomy were 100%, 73%, 89% and 100%, respectively. However, when "unsure", only in 30% OCR was achieved. We concluded that mini-laparotomy is a safe, simple and effective technique for predicting feasibility of OCR. This simple technique could obviate the need for full laparotomy in patients who may benefit from neo-adjuvant chemotherapy.

Keywords Mini-laparotomy · Optimal · Resectability · Advanced ovarian malignancy · Ovarian cancer

Background

Ovarian cancer is the most common gynaecological malignancy in the United Kingdom with nearly 7,000 reported cases per year [1]. The mainstay of treatment remains cytoreductive surgery followed by adjuvant chemotherapy. Most patients present with omental and peritoneal metastasis, hence making complete surgical resection a challenge [2]. Overall survival greatly depends on the possibility of resecting most of the diseased areas with little residual disease (optimal cytoreduction [OCR]). In early stages (I and II), when the disease is confined to the pelvis, the overall survival exceeds 70% whilst in advanced stages (III and IV) the 5-year survival is less than 40% [3]. In Scotland alone, there are over 600 reported cases per annum, of which over 400 succumb to the disease, hence an overall reported survival of 30% [4].

Achievement of OCR is the single most important factor for survival in patients with ovarian cancer [5]. Currently the preoperative selection of patients for primary surgery is mainly based on detailed imaging [6, 7]. The commonest imaging modality is computer tomography (CT) in most centres. CT accuracy varies greatly in assessment of resectability and prediction of OCR in advanced stage ovarian cancer [8]. Other preoperative assessment methods such as laparoscopy have not as yet been established in most centres.

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This pilot study aims to evaluate the role of minilaparotomy as an alternative or complementary (to CT imaging) means of determining tumour resectability and the chances of achieving OCR in advanced stage ovarian cancer.

Methods

A total of 50 consecutive patients with stage III/IV ovarian cancer were included in this prospective study. The study was conducted in the department of gynaecological oncology at Glasgow Royal Infirmary, over a period of 18 months. They had all undergone CT imaging and clinical examination by the three gynaecological oncologists who performed the operations (FAS, JD and JK). Their CT images were reviewed in our regular multidisciplinary team meetings (MDT), involving our lead radiologist and all the above operating surgeons. Following this review, only selected patients in whom OCR was deemed possible on CT imaging were then admitted for surgery. All subjects had also been previously examined and were regarded as fit and potentially suitable for up-front surgical intervention. These were therefore regarded as the inclusion criteria. Patients with advanced metastatic stage (stage IV), unfit or in cases OCR was deemed unfeasible, were excluded from the study.

Data gathering and documentation were pro-forma based, recording the operative findings from both minilaparotomy and formal laparotomy. The details of minilaparotomy included the site, length of minilaparotomy incision, time taken to enter the peritoneal cavity, time taken to finish the procedure and any operative or post operative complication.

Prior to operation, an examination under anaesthesia was performed in all of the cases. A mini-laparotomy incision was made through a midline incision. The incision was sited below, over or above the umbilicus, depending on where the surgeon deemed it most appropriate for full intra-

abdominal assessment. The length of the incision was kept to the minimum, allowing only the insertion of one hand after some negotiation. This length usually corresponded to the surgeon's glove size or width of the palm at metacarpophalangeal joint (Fig. 1). To keep the incision length to the minimum, the surgeon placed his hand over the preferred site of incision and marked either side of his knuckles with a marker pen. The length between the two marked points was measured prior to incision and starting time noted.

The palpation findings were charted as main tumour mass, other pelvic and abdominal organs, peritoneal and omental disease. This was followed by visual inspection, usually using a Morris retractor to allow lifting of the anterior abdominal wall. The time taken to enter abdomen, and to complete both palpation and inspection was recorded.

The criteria for OCR were regarded as residual tumour less than 2 cm in volume. This criteria still remains an acceptable standard in most UK centres. Following inspection and palpation through the mini-laparotomy, the surgeon's impression of achieving OCR was recorded as "possible" (meaning that it may be possible to reduce the volume of disease to less than 2 cm), "not possible" or "unsure". The impression was mainly based on palpation findings, as the visual inspection in most cases was limited.

Formal laparotomy was then carried out by the same operator, extending the mini-laparotomy incision, both caudally and cranially. On completion of the surgery, the operating surgeon documented whether "optimal" or "suboptimal" cytoreduction had been achieved.

Findings and analysis

Data was collected for a total of 50 consecutive patients with suspected stage III/IV ovarian cancer, over a span of 18 months. Data from five cases was regarded as either

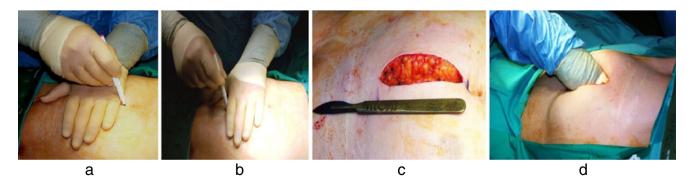


Fig. 1 The steps involved in minilaparotomy: demarcation of a suitable midline site and length corresponding to the width of the palm at meta-carpo-phalygeal joint (a and b), formation of

minilaparotomy was limited to the site marked (c), systemic intraabdominal palpation, examination and assessment of resectability (d)



inadequate or retrospective and hence excluded. Hence results from 45 cases were analysed. The median age of the patients was 64 years (mean 62.5 years, range 34–80, SD 11.71). The median BMI was 29.1 kg/m² (mean 29.9 kg/m², range 18.7–67.5, SD 7.55). The median abdominal girth was 95 cm (mean 95.87 cm, range 42–148, SD 16.79). Fifteen patients (33%) had previously undergone abdominal surgery for various gynaecological and non gynaecological reasons. Based on clinical and radiological imaging, stage III disease was suspected in thirty eight (84.44%) and stage IV in seven cases (5.56%). The majority of these patients (n=40, 89%) underwent upfront surgery prior to adjuvant chemotherapy. Only five patients had received neo-adjuvant chemotherapy (11%) prior to their delayed primary surgery.

The median length of the mini-laparotomy incision was 9 cm (mean 8.59 cm, range 5–12, SD 1.16). The median time to enter the peritoneal cavity, starting from knife to skin, was 4 min (mean 5.10 min, range 1–15, SD 3.04) while the median time taken to complete the palpation and visual exploration was 6.5 min (mean 7.52 min, range 2–30, SD 4.83). The total median time for making the incision and completion of examinations was 10 min (with mean of 12.62 min, range 5–45, SD 7.23).

Satisfactory palpation of pelvis and upper abdomen, through the mini-laparotomy incision, was proven to be possible in all cases (100%) but visual inspection was limited, particularly for upper abdominal organs.

Following formation of mini-laparotomy, 27 out of the 45 cases were deemed optimally cyto-reducible or "OCR possible". From this "possible" group, OCR was achieved in 24 cases. In eight cases, OCR was regarded as "not possible" and in none of these cases was OCR achieved. In the remaining ten patients, the surgeon was "unsure" and in only three of these cases was OCR achieved (Table 1).

The specificity of mini-laparotomy to predict OCR was noted to be 73% and sensitivity was 100%. Positive predictive value (PPV) of mini-laparotomy in assessment of OCR was calculated to be 89% and negative predicative value (NPV) of 100%.

In three cases, OCR was not possible despite initial assessment through mini-laparotomy. In one patient, part of ovarian capsule could not be fully resected away from pelvic side wall. In the second case, failure to achieve OCR was due to tumour plaques in Pouch of Douglas (POD) and over rectum. The third patient had previous total abdominal

hysterectomy and residual tumour plaques could not be fully resected from vaginal vault and bladder base.

Out of ten cases commented as "unsure" at minilaparotomy, in only three cases (30%) was OCR achieved at laparotomy. Each of these three cases required radical surgery, including bowel resection in two patients.

In eight cases OCR was regarded as not possible after mini-laparotomy examinations and in none of these was OCR achieved after full laparotomy. In four of these patients (50%) the main reason was fixed central pelvic mass densely adherent to the adjacent pelvic and abdominal structures. In the other four patients (50%), extensive upper abdominal disease involving diaphragmatic surfaces, spleen, stomach and porta hepatis were the main reasons precluding OCR. In addition to the above eight cases, there were ten patients that OCR was not achieved (3 falsely predicted as "OCR possible" and in seven cases the operator was "unsure"). This was due to following reasons: fixed and densely adherent pelvic mass, extensive supracolic omental mass, significant peritoneal disease and involvement of upper abdominal organs.

There were no complications associated with minilaparotomy. However, excess bleeding according to the surgeon's judgment was reported in three cases (6%). The bleeding in all three cases was noted to be from central tumour mass that was adherent to the peritoneum over the site of mini-laparotomy incision.

Discussion

OCR remains the gold standard treatment for all stages of ovarian cancer. In advanced ovarian malignancy, achievement of OCR has been shown to have a direct impact on patient's disease-free interval and overall survival [9–11]. The definition of OCR has greatly changed over the years: from residual disease of less than 2 cm to resection or ablation of all macroscopic disease in recent years [12]. However, in most UK centres, a residual disease of less than 2 cm remains an acceptable goal. Currently the mainstay of preoperative assessment of such patients with regards to resectability and likelihood of achieving OCR are based on clinical examinations, tumour marker levels and CT findings [8–11, 13–16]. It is also common practice in UK to administer neo-adjuvant chemotherapy prior to

Table 1 Comparison of outcome of mini-laparotomy and laparotomy

	OCR achieved after full laparotomy	OCR not achieved after full laparotomy	Total
OCR predicted possible on mini-laparotomy	24 (true positive)	3 (false positive)	27
OCR predicted not possible on mini-laparotomy	0 (false negative)	8 (true negative)	8
Unsure following mini-laparotomy	3	7	10



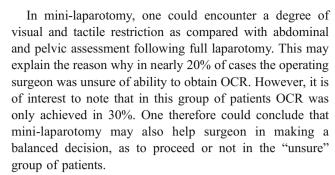
delayed primary surgery in patients that OCR may not appear feasible on initial assessment.

Current preoperative assessments have limited predictability in selection of patients suitable for OCR. The sensitivity of CT in identifying patients that OCR can be achieved is around 60–70%, with varying reported specificity between 70% and 100% [17, 18]. The main drawbacks of CT imaging is the inability to reliably assess the extent of disease in the supra colic compartment, porta hepatis, miliary disease over bowel mesentery and peritoneum, especially in absence of ascites [8–11, 19].

In order to overcome the limitations associated with CT, a few centres routinely perform laparoscopic staging in all their ovarian cancer patients prior to formal laparotomy. This modality of assessment, in experienced hands, has been shown to have PPV of 87% and NPV of 100% [20-22]. However, laparoscopic assessment requires an experienced surgical team and laparoscopic resources. In addition, it is associated with increased theatre time and possible increased morbidity. Furthermore, there are reported technical difficulties with the laparoscopic procedure, particularly when intra abdominal adhesions are present in advanced stage disease [23–26]. More importantly, there is a lack of tactile sensation which is of great importance in assessment of resectability and achievement of OCR [27-29]. We believe such shortcomings can be avoided by using a simple minilaparotomy as described in this paper. Furthermore, time required to perform mini-laparotomy would be much less than laparoscopic method due to its simplicity and greater familiarity with the technique. There is no need for a learning curve in mini-laparotomy unlike laparoscopic techniques and is clearly more cost effective.

In our group of patients, all 45 patients were regarded as suitable candidates for OCR following MDT review. However, OCR was achieved in only 27 out of the 45 patients (60%). This finding is consistent with reported suboptimal CT accuracy in predicting OCR in the literature [30]. In contrast, following intraabdominal palpation through mini-laparotomy, the accuracy for predicting OCR was much higher at 89% (24 of 27cases). More importantly, OCR was not achieved in any of the eight patients where it was predicted that OCR would not be possible at mini-laparotomy, reflecting an NPV of 100%. This has great clinical importance in avoiding unnecessary major laparotomy. In addition, considering small size and lack of morbidity associated with mini-laparotomy, it should result in much quicker recovery and hence shorten delay in receiving neo-adjuvant chemotherapy.

We acknowledge the principle shortcoming of the study to be the small sample size, hence potential for type II statistical error. However, despite this limitation, we believe the findings to be of clinical importance.



In order to overcome the visual limitation of minilaparotomy, one may consider a combination of the techniques of laparoscopy and mini-laparotomy, better known as hand-assisted laparoscopy. However such technique will be inevitably associated with increased theatre time and operative complexity. In future, it is anticipated that integrated FDG-PET/CT may have a role in assessment of disease spread and decision with regards to best therapeutic strategy in ovarian cancer. FDG-PET/CT has been shown to have a higher sensitivity (91%) than PET, CT or MRI in assessment of disease spread [31, 32]. Hence, it may further assist in early identification of patients who are not suitable for up front surgery.

Conclusions

Despite our small sample size, this study has shown minilaparotomy to be a safe, simple and effective technique in assessment of tumour resectability and chances of obtaining OCR. More importantly it is very accurate in identifying patients unsuitable for primary cytoreduction. By avoiding an unnecessary full laparotomy, patients' recovery will be expedited and chemotherapy could be initiated earlier.

Conflicts of interest The authors report no conflicts of interest. The authors alone are responsible for the content and writing of the paper.

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