Surgical safety and personal costs in morbidly obese, multimorbid patients diagnosed with early-stage endometrial cancer having a hysterectomy

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Abstract

Background: Many women who develop endometrial cancer (EC) or endometrial hyperplasia with atypia are obese and therefore at high risk of surgical complications. Recently clinical trials have been initiated offering non-surgical treatment to these women, but not all may agree to participate in such trials. This paper aims to describe the patient characteristics, and surgical outcomes of women with suspected early stage endometrial cancer and body mass index (BMI) of 30 or greater, who declined enrolment in the feMMe trial, which offers non-surgical hormonal treatment, hormonal plus metformin or hormonal plus weight loss as primary treatment.

Methods: Consecutive case series from a tertiary gynaecological oncology unit. Over the course of the first 2 years of the feMMe trial, 27 patients met the initial eligibility screening, but declined enrolment in the feMMe trial and opted for upfront surgery. The main surgical outcome measures were type of surgical approach, need for conversion from laparoscopic to open approach, length of stay in hospital and adverse events.

Results: Patients' median age was 63 years (range 40 to 86); median BMI was 37.3 kg/m2 (range 30.7 to 54.7); median medical co-morbidities were six (range 3–10). Of the 26/27 surgeries planned to be undertaken laparoscopically, 2/26 patients had to be converted (7 %). Overall, the average hospital stay was 4.5 days, and 11/27 (41 %) of the patients developed one or more adverse events grade 2+ rated according to the Common Toxicity Criteria Version 3.

Conclusions: Adverse surgical outcomes are common in multi-morbid, obese or morbidly obese patients diagnosed with early stage EC or endometrial hyperplasia with atypia and who have a hysterectomy.

Keywords: Endometrial cancer, Endometrial hyperplasia with atypia, Obesity, Surgery, Adverse event

Background

Obesity is a massive health issue in many countries around the world and is also the major risk factor for Endometrial Cancer (EC) [1–4]. It has previously been reported that obesity causes at least 39 % of cases of EC [5]. Obesity is also associated with increased risk of medical co-morbidities (e.g., diabetes, cardiovascular) [6–8]; the need for intense preoperative assessments; perioperative complications [9]; conversion from laparoscopic to open surgery [10]; intensive postoperative care [11, 12]; treatment costs [11, 13]; and reduced recurrence-free survival [9, 14, 15].

While surgical treatment of EC is generally effective, it does not address the specific needs of the steadily growing group of morbidly obese and multi-morbid patients as well as young obese patients still desiring fertility [16, 17]. For these growing groups of patients treatment often comes at a high personal cost (long hospital stay, protracted recovery from surgery, high incidence of postoperative complications) and subsequent high healthcare cost. We previously estimated hospital costs of $12,872 vs...
The challenges of pre- and postoperative care for multi-morbid and morbidly obese EC patients impact on a variety of resources. Thus, as highlighted by the Royal College of Obstetricians and Gynaecologists, treatment for EC needs to be reassessed in this complex and increasingly common situation [18]. The search for treatment alternatives that are safe, effective and less harmful than surgery is warranted.

Recently, the Gynecologic Cancer InterGroup identified conservative treatment for fertility sparing purposes and to treat morbidly obese women as a most pressing research priority at their EC Clinical Trials Planning Meeting in the Netherlands [19].

To address this need the feMMe trial was initiated in 2013 [20]. It is an open-label, randomised clinical trial exploring conservative, non-surgical treatment options to achieve a pathological complete response in patients diagnosed with early-stage EC (ANZGOG #1301, NCT01686126).

The aim of the present study was to describe the safety and clinical outcomes of consecutive patients who would have fulfilled the eligibility criteria for the feMMe trial and who were offered participation in the feMMe trial, but declined enrolment and opted for hysterectomy at two institutions instead.

**Methods**

Approval for this study was received from the Royal Brisbane & Women’s Hospital Human Research Ethics Committee (HREC/15/QRBW/113). All patients reported here have been identified through gynaecological oncology services at Royal Brisbane and Women’s Hospital and Greenslopes Private Hospital. These patients would have been considered potentially eligible to be enrolled in the feMMe trial.

The feMMe trial is an open label, randomised phase II trial with three treatment arms and is recruiting patients at present [20]. The three arms consist of Intrauterine Progestin (IUP) placed into the uterine cavity (45 patients); IUP plus Metformin 1000 mg daily (75 patients); or IUP plus weight loss through Weight Watchers (45 patients). Weight Watchers is a standardised, evidence-based and formally tested weight loss intervention including diet, physical activity, social networking and support via a network of lifestyle centres, one-on-one support and an online program [21, 22]. It has been shown to be the most cost-effective among a range of currently available weight loss programs [22].

In this phase II trial randomisation aims to eliminate selection bias rather than allow a formal comparison of groups. Trial methodology, in-/exclusion criteria, randomisation /stratification and study assessments were published recently [20]. Human Research and Ethics Committee and site-specific approvals are underway in various Australian States but only sites in the state of Queensland are fully approved and enrolling at present. All patients are followed for 6 months and a central pathology review will be conducted once all patients are enrolled.

Eligibility criteria: Only patients with histologically confirmed innocuous EC or endometrial hyperplasia with atypia, Body Mass Index (BMI) >30 kg/m² who wish to retain fertility or who suffer from medical impairments and are considered suboptimal candidates for hysterectomy are eligible [20].

Patients are excluded from enrolment if they had a histological type other than endometrioid adenocarcinoma of the endometrium, clinically advanced disease, involvement of the uterine cervix or enlarged retroperitoneal lymph nodes.

To be eligible for this study, patients had to have a CT scan of the abdomen and pelvis as well as imaging (CT or X-Ray to the chest) suggesting the absence of extrauterine disease. Patients are also only considered eligible if their baseline serum CA-125 reading was 30 U/ml or less.

Patients who agree to proceed with enrolment into the feMMe trial receive a pelvic MRI to ensure that the depth of invasion is not greater than 50 % of myometrium and to re-confirm the absence of extrauterine dissemination. Patients who decline participation in feMMe (including the patients reported herein) however do not receive an MRI as it is not part of the standard imaging workup in our institutions.

In addition to the established criteria for low-risk disease (CT and MRI scan showing the absence of extrauterine disease, FIGO grade = 1) we offer enrolment only to patients with serum CA125 of 30 U/ml or less [23]. Considering the strict criteria above, we expect that the risk of enrolling patients with advanced or aggressive disease is minimal.

All patients reported here did not qualify for feMMe because they preferred hysterectomy and as a consequence declined enrolment into the feMMe trial. Hence, following standard protocols, a pelvic MRI was not offered (as has been explained above).

In all 27 patients, pre-existing medical co-morbidities were recorded as well as any intra- or post-operative Adverse Events (AEs) up to 30 days post-operatively. We coded AEs according to the post-operative Common Toxicity Criteria (CTC) Version 3 and report any AEs grade 2+ (moderate to severe AEs). Analyses were restricted to women who completed 30 days of follow-up after surgery.

**Results**

The clinical outcomes of 27 patients who fulfilled the eligibility criteria but declined participation in the feMMe
trial and have chosen primary surgical treatment instead are reported here.

Patients’ median age was 63 years (range 40 to 86 years) and the median BMI was 37.3 kg/m² (range 30.7 to 54.7 kg/m²), median ASA was 3 (range 2–3). At baseline a total of 167 medical co-morbidities were recorded among the 27 patients including hypertension, hyperlipidemia, hypercholesterolemia, diabetes, obstructive sleep apnoea, fatty liver and many other lifestyle-related ailments.

Twenty-six patients had a total laparoscopic procedure of which two patients (7 %) had to be converted to a laparotomy. One patient required an abdominal hysterectomy through a midline incision (Table 1).

The reasons for conversion to open surgery in two patients included an inadvertent gastrotomy through a trocar at primary port entry, which required primary surgical closure. The second patient sustained an enterotomy to the small bowel during adhesiolysis. The

| Patient # | Age (years) | D&C histology | BMI | Surgical approach | Node dissection | LOS | Stage | FIGO grade | Depth of invasion (%) | Intra-, Postoperative complications |
|-----------|-------------|---------------|-----|-------------------|----------------|-----|-------|------------|-----------------------|-----------------------------------|
| 1         | 40          | EHA           | 31.0| TLH               | 0              | 2   | 1a    | 1          | 15                    |                                   |
| 2         | 56          | G1 EAC        | 49.2| TLH               | 0              | 2   | 1a    | 1          | 0                     | Post-operative bleeding           |
| 3         | 62          | G1 EAC        | 35.1| TLH               | 0              | 2   | 1a    | 1          | 11                    | Vault haematoma, abdominal cramping, urinary frequency |
| 4         | 69          | G1 EAC        | 54.7| TLH               | 0              | 2   | 1a    | 2          | 38                    |                                   |
| 5         | 75          | G1 EAC        | 39.6| TLH               | 0              | 2   | 1a    | 1          | 0                     |                                   |
| 6         | 65          | G1 EAC        | 40.9| TLH               | 0              | 2   | 1b    | 2          | 60                    |                                   |
| 7         | 56          | G1 EAC        | 42.7| TLH               | 0              | 5   | 1a    | 1          | 0                     |                                   |
| 8         | 74          | G1 EAC        | 46.8| TLH               | 0              | 1   | 1b    | 2          | 52                    |                                   |
| 9         | 73          | G1 EAC        | 44.8| TLH               | 0              | 8   | 2     | 2          | 18                    | Post-operative bleeding, anaemia, retroperitoneal haematoma, rise in Troponin |
| 10        | 68          | EHA           | 35.4| TLH               | 1              | 2   | 1b    | 2          | 60                    | Unplanned gastrostomy, unplanned stay in ICU |
| 11        | 75          | G1 EAC        | 38.9| TLH converted to TAH | 0          | 14  | 1b    | 2          | 57                    |                                   |
| 12        | 69          | EHA           | 45.5| TLH               | 0              | 2   | 1a    | 1          | 0                     |                                   |
| 13        | 67          | G1 EAC        | 32.5| TLH               | 0              | 3   | 1a    | 2          | 8                     | Vault haematoma, hypokalaemia, sinus tachycardia |
| 14        | 57          | G1 EAC        | 30.7| TLH               | 0              | 2   | 1a    | 1          | 50                    | Vault haematoma, pain             |
| 15        | 73          | G1 EAC        | 34.3| TLH               | 0              | 5   | 1a    | 1          | 10                    |                                   |
| 16        | 69          | EHA           | 29.4| TLH               | 1              | 2   | 1a    | 1          | 0                     | Vault haematoma, hypotension, hypokalaemia |
| 17        | 68          | G1 EAC        | 43.9| TLH               | 0              | 2   | 1a    | 2          | 45                    |                                   |
| 18        | 78          | G1 EAC        | 32.0| TLH               | 0              | 2   | 1a    | 1          | 45                    |                                   |
| 19        | 63          | EHA           | 33.5| TLH               | 0              | 2   | 0     | 0          | 0                     | Hypertension                       |
| 20        | 66          | EHA           | 40.6| TLH               | 0              | 2   | 1a    | 1          | 1                     | Atrial fibrillation               |
| 21        | 60          | EHA           | 37.1| TLH               | 0              | 7   | 1a    | 1          | 37                    |                                   |
| 22        | 70          | EHA           | 43.1| TLH               | 0              | 5   | 1b    | 2          | 75                    | Chest infection, wound dehiscence, vault haematoma, pain |
| 23        | 72          | G1 EAC        | 42.7| TLH               | 1              | 3   | 3b    | 1          | 100                   |                                   |
| 24        | 50          | EHA           | 35.6| TLH               | 0              | 9   | 1a    | 1          | 39                    | Fluid overload, pulmonary oedema   |
| 25        | 55          | G1 EAC        | 33.3| TAH               | 0              | 4   | 2     | 1          | 7                     |                                   |
| 26        | 73          | G1 EAC        | 47.0| TLH               | 0              | 2   | 1a    | 1          | 0                     |                                   |
| 27        | 86          | G1 EAC        | 43.6| TLH converted to TAH | 1          | 28  | 1a    | 1          | 31                    | Unplanned enterotomy, wound infection, Atrial fibrillation, renal failure |

**Abbreviations**: BMI body mass index, EAC endometriod adenocarcinoma, EHA endometrial hyperplasia with atypia, TLH total laparoscopic hysterectomy, TAH total abdominal hysterectomy, LOS length of stay, FIGO the international federation of gynecology and obstetrics
adhesions could not be dissected from the anterior abdominal wall laparoscopically. The enterotomy was recognised at surgery and the operation was completed through open surgery. The median percentage of invasion into the endometrium was 31 % (range 0–100 %).

The average postoperative hospital stay was 4.5 days (median 2 days), ranging from 1 to 28 days. The patient with a 28-day hospital stay was a patient with a body mass index of 43.6 kg/m2 who required conversion from laparoscopic to open surgery. She developed a wound infection (limited to the subcutaneous adipose tissue), atrial fibrillation resulting in a stay at the Cardiac Care Unit followed by acute renal failure. The patient was discharged into rehab on day 28 post surgery.

Within 30 days from surgery, 12 patients developed at total of 30 AEs. One of 27 patients developed an AE CTC grade 1 and 11/12 patients developed one or more AEs CTC grade 2+ (41 %). All but 5 AEs were surgery related (Table 1).

Nine patients were enrolled to treat endometrial hyperplasia with atypia based on a pre-hysterectomy endometrial biopsy or curette and 18 of 27 patients had surgery for histologically proven endometrioid endometrial adenocarcinoma on endometrial biopsy or curette.

Of those nine patients who were treated for endometrial hyperplasia with atypia, seven patients were found to have endometrial adenocarcinoma in the final histopathology specimen of the uterus.

In patients with the final histopathological outcomes confirming endometrial adenocarcinoma, all patients were diagnosed with endometrioid cell type. In those patients FIGO grade was grade 1 in 14/25 patients and grade 2 in 10/25 patients. In one patient there was no residual disease at hysterectomy. The depth of invasion was limited to inner half in all but five patients.

Two patients had extension of disease into the endocervix (stage 2) and one patient had full thickness myometrial invasion of a grade 1 adenocarcinoma and focal involvement of a fallopian tube (FIGO stage 3b).

Discussion

Main findings

Adverse surgical outcomes are common in multi-morbid and morbidly obese patients diagnosed with early stage EC who have a hysterectomy. Obesity is an independent risk factor for AEs, regardless of the surgical approach [12]. Obese women will have a higher risk of conversion to open surgery [10] and their risk of surgical AEs is higher [24].

For comparison we quote data from the prospective randomised and multi-institutional LACE trial below [12, 25]. The LACE trial compared open with laparoscopic surgery for early stage EC or endometrial hyperplasia with atypia. It was an international trial but the vast majority of patients were treated in Australian institutions.

In the case series reported here, all but one operations were planned to be performed laparoscopically; two of the 27 patients required a conversion from laparoscopy to laparotomy (7 %) and one patient required a primary laparotomy, implying that 10 % of patients required a laparotomy to accomplish the surgical task of a hysterectomy. By contrast, the conversion rate from laparoscopic to open in the prospective randomised and multi-institutional LACE trial was only 3.8 %, most likely due to omitting the requirements for a comprehensive pelvic and aortic retroperitoneal node dissection in these patients and a smaller proportion of patients with a BMI of 30 or greater.

By contrast, pelvic and aortic lymph node dissection was mandatory in the LAP-2 trial corresponding to a 25.8 % conversion rate. Patients with a high BMI had an up to 60 % risk of conversion to open surgery [24].

In the context of morbidly obese and multi-morbid patients we typically aim to minimise the risk of conversion, which may attract further intraoperative and postoperative morbidity. In those instances patients’ adjuvant treatment may be guided by their general medical health and histopathological features available from the primary tumour. At present only low-level evidence is available on the feasibility and safety of robotic surgery in morbidly obese patients requiring a retroperitoneal node dissection. Deaths due to complications of robotic surgery have also been reported [26].

In this sample, the mean length of hospital stay (LOS) was 4.5 days. Length of stay was largely associated with the development of postoperative complications. However, in some patients an uneventful postoperative recovery still required a longer than expected LOS due to slow recovery. In the LACE trial, reflecting the Australian health care situation the LOS was 2.4 days for patients assigned to have a laparoscopic hysterectomy and 5 days for patients who were randomised to have a laparotomy.

In this series of patients the per-patient incidence of surgical AEs CTC grade 2+ was high at 41 %.

Strength and limitations

Innovatively this case series details the outcomes of patients who were offered enrolment in a non-surgical clinical trial, but preferred surgery. These results again highlight the increased risk of obese patients to develop complications as previously shown in other international series. This group of morbidly obese and multi-morbid patients carries a high risk of conversion to laparotomy, a longer hospital stay and a three to four times higher risk of surgical AEs compared to previous series with a wider range of BMI. LOS and AEs are significant contributors to health care costs and funders of health care services.
must therefore expect high costs among obese patients treated surgically for EC [13]. Limitations of this study include the non-random assignment to surgery, which was based on patients’ preference, as well as the relatively small number of patients available for data collection.

Conclusions
While hysterectomy for EC offers excellent survival outcomes, it also comes at a price: slow recovery from surgery, surgical AEs, loss of fertility, financial and societal treatment costs [27, 28]. Importantly these results indicate that current risk estimations do not take populations at high surgical risk (e.g. obese and multi-morbid patients) into sufficiently account. Thus, the efficacy of treatment alternatives need to be assessed in the complex and increasingly common situation of obesity and EC or endometrial hyperplasia with atypia [18]. We envisage that for obese and multimorbid patients less invasive treatments will achieve equivalent survival outcomes at a lower personal and financial cost to patients and society [13].

Historically intracavitary brachytherapy has been used to treat patients at advanced age and severe medical illnesses and with the advent of IMRT [29], radiotherapy may be well positioned to be evaluated in clinical trials as an alternative to major surgery.

By contrast, others units currently investigate the effectiveness of a levonorgestrel containing intrauterine device for the treatment of endometrial cancer. The feMMe trial (ANZGOG #1301, NCT01686126) is an international, phase II, 3-arm randomised clinical trial exploring conservative, non-surgical treatment options to achieve a pathological complete response in early-stage endometrial cancer patients who are suboptimal candidates for hysterectomy [20].

In addition to the eligibility criteria for this case series, patients have to have an MRI of the pelvis to determine the depth of myometrial invasion. In the context of the results reported here this is well warranted, and the pelvic MRI will also be critical to exclude involvement of the cervix and/or adnexae.

The trial is recruiting at present (Fig. 1 shows the recruitment chart) and also includes a molecular component investigating the mechanisms of change associated with response or non-response to feMMe intervention (tumour polymorphisms; molecular phenotype of tumours; circulating cytokines, such as adipokines, hormones and growth factors). Phase II results are expected for 2017, and early discussion about the optimal Phase III trial design to follow have been initiated in 2015.

Ethics approval
Royal Brisbane & Women’s Hospital Human Research Ethics Committee (HREC/15/QRBW/113).

Competing interests
AO acts as a consultant for Covidien and receives research support from Ethicon, a Johnson and Johnson company. AO has received travel support from Gate healthcare. AO is the founder and director of SurgicalPerformance.com, an online audit software. All other authors have declared no conflicts of interest.

Authors’ contributions
The authors conceived and designed the feMMe trial and contributed to the conception of the present analyses of patients who prefered surgical treatment. All authors contributed to the writing of this article or revised it critically. All authors gave final approval of the version to be published and will be accountable for all aspects of the work.

Fig. 1 Recruitment chart for the FeMMe trial
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References
1. Reneneh AG, Tyson M, Egger M, Heller RF, Zvahalen M. Body mass index and incidence of cancer: a systematic review and meta-analysis of prospective observational studies. Lancet. 2008;371(9612):569–78.
2. Kaaks R, Lukanova A, Kurzer MS. Obesity, endogenous hormones, and endometrial cancer risk: a synthetic review. Cancer Epidemiol Biomarkers Prev. 2002;11(12):1531–43.
3. Nagle CM, Marquart L, Bain CJ, O’Brien S, Lahrann PH, Quinn M, et al. Impact of weight change and weight cycling on risk of different subtypes of endometrial cancer. Eur J Cancer. 2013;49(12):2717–26.
4. Park SL, Goodman MT, Zhang ZF, Kolonel LN, Henderson BE, Setiawan VW. Body size, adult BMI gain and endometrial cancer risk: the multicentric cohort. Int J Cancer. 2010;126(26):490–9.
5. Bergstrom A, Pisani P, Tenet V, Wolk A, Adami HO. Overweight as an avoidable cause of cancer in Europe. Int J Cancer. 2001;91(3):421–9.
6. Bhakot KJ, Doss-Santos Silva I, Leon DA, Speak L. Body mass index and risk of 22 specific cancers: a population-based cohort study of 5.2 million UK adults. Lancet. 2014;384(9945):755–65.
7. Callie EE, Kaaks R. Overweight, obesity and cancer: epidemiological evidence and proposed mechanisms. Nat Rev Cancer. 2004;4(8):579–91.
8. Lucenteforte E, Bosetti C, Talamini R, Montella M, Zucchetto A, Pelucchi C, et al. Diabetes and endometrial cancer: effect modification by body weight, physical activity and hypertension. Br J Cancer. 2007;97(7):995–8.
9. Gunderson CC, Java J, Moore RN, Walker JL. The impact of obesity on surgical staging, complications, and survival with uterine cancer. a Gynecologic Oncology Group LAP2 ancillary data study. Gynecol Oncol. 2014;133(1):23–7.
10. Mourtis MJ, Bijen CB, Arts HU, ter Brugge HG, van der Spade R, Paulsen L, et al. Safety of laparoscopy versus laparotomy in early-stage endometrial cancer: a randomised trial. Lancet Oncol. 2010;11(8):763–71.
11. Kondalsamy-Chennakesavan S, Gordon LG, Sanday K, Bouman C, De Jong S, Nicklin J, et al. Hospital costs associated with adverse events in gynecological oncology. Gynecol Oncol. 2011;121(1):70–5.
12. Kondalsamy-Chennakesavan S, Janda M, Gebski V, Baker J, Brand A, Hogg R, et al. Risk factors to predict the incidence of surgical adverse events following open or laparoscopic surgery for apparent early stage endometrial cancer: results from a randomised controlled trial. Eur J Cancer. 2012;48(14):2155–62.
13. Graves N, Janda M, Merollini K, Gebski V, Obermair A, committee Lt. The cost-effectiveness of total laparoscopic hysterectomy compared to total abdominal hysterectomy for the treatment of early stage endometrial cancer. BMJ Open. 2013; 3(4): e001884. doi:10.1136/bmjopen-2012-001884.
14. Canlorbe G, Bendifallah S, Raimond E, Graesslin O, Hudy D, Coutant C, et al. Severe Obesity Impacts Recurrence-Free Survival of Women with High-Risk Endometrial Cancer: Results of a French Multicenter Study. Ann Surg Oncol. 2012;22(4):714–24.
15. Mclennan J, Irwin M, Vongruenigen V. Weight, physical activity, diet, and prognosis in breast and gynecologic cancers. J Clin Oncol. 2010;28(26):4074–80.
16. Linkov F, Elshafei E, Gloyeske N, Edwards R, Athouse AD, Geller MA, et al. Bariatric surgery-induced weight loss changes immune markers in the endometrium of morbidly obese women. Surg Obes Relat Dis. 2014;10(5):921–6.
17. Nevadunsky NS, Van Arsdaile A, Strickler HD, Moadel A, Kaur G, Levitt J, et al. Obesity and age at diagnosis of endometrial cancer. Obstet Gynecol. 2014;124(2 Pt 1):300–6.
18. Endometrial Cancer in Obese Women. Scientific Impact Paper No. 32. London, United Kingdom: Royal College of Obstetricians and Gynaecologists; 2012.
19. Creutzberg CL, Kitchener HC, Birrer MJ, Landoni F, Liu KH, Powell M, et al. Gynecologic Cancer InterGroup (GCIG) endometrial cancer clinical trials planning meeting: taking endometrial cancer trials into the translational era. Int J Gynecol Cancer. 2013;23(8):1528–34.
20. Hawkes AL, Quinn M, Gebski V, Armes J, Brennan D, Janda M, et al. Improving treatment for obese women with early stage cancer of the uterus: rationale and design of the levonorgestrel intrauterine device +/- metformin +/- weight loss in endometrial cancer (feMMe) trial. Contemp Clin Trials. 2014;39(1):14–21.
21. Jebb SA, Ahern AL, Olson AD, Aston LM, Holzapfel C, Stoll J, et al. Primary care referral to a commercial provider for weight loss treatment versus standard care: a randomised controlled trial. Lancet. 2011;378(9780):1485–92.
22. Finkelstein EA, Kruger E. Meta- and cost-effectiveness analysis of commercial weight loss strategies. Obesity (Silver Spring). 2014;22(9):1942–51.
23. Nicklin J, Janda M, Gebski V, Jobling T, Land R, Manolitsas T, et al. The utility of serum CA-125 in predicting extra-uterine disease in apparent early-stage endometrial cancer. Int J Cancer. 2012;131(4):885–90.
24. Walker JL, Piedmonte MR, Sipitos NM, Eisenkop SM, Schlaerth JB, Mannell RS, et al. Laparoscopy compared with laparotomy for comprehensive surgical staging of uterine cancer: Gynecologic Oncology Group Study LAP2. J Clin Oncol. 2009;27(32):5331–6.
25. Obermair A, Janda M, Baker J. Kondalsamy-Chennakesavan S, Brand A, Hogg R, et al. Improved surgical safety after laparoscopic compared to open surgery for apparent early stage endometrial cancer: results from a randomised controlled trial. Eur J Cancer. 2012;48(8):1147–53.
26. Shields K, Minion L, Willmott L, Sumner D, Monk B. Ten-Year Food and Drug Administration Reporting on Robotic Complications in Gynecologic Surgery. J Gynecol Surg. 2015;31(6):331–5.
27. Tangjitgamol S, Anderson BO, See HT, Lertbutsayanukul C, Siriabaya N, Manchana T, et al. Management of endometrial cancer in Asia: consensus statement from the Asian Oncology Summit 2009. Lancet Oncol. 2009;10(11):1119–27.
28. Zullo F, Falbo A, Palomba S. Safety of laparoscopic vs laparotomy in the surgical staging of endometrial cancer: a systematic review and metaanalysis of randomized controlled trials. Am J Obstet Gynecol. 2012;207(2):194–100.
29. Paganiat H. Changes in tumor cell response due to prolonged dose delivery times in fractionated radiation therapy. Int J Radiat Oncol Biol Phys. 2005;63(3):892–900.

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