Research Report

Neoadjuvant chemotherapy for advanced stage endometrial cancer: A systematic review

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ABSTRACT

Objective: While primary cytoreductive surgery (PCS) is considered the standard of care for women who present with stage IV endometrial cancer, neoadjuvant chemotherapy (NACT) followed by interval cytoreductive surgery (ICS) has emerged as an alternative treatment strategy. We summarized the literature and compared outcomes of PCS compared to NACT and ICS.

Methods: We conducted a systematic search on PubMed, Embase, Web of Science, and Scopus for articles published from January 1, 1990 to December 31, 2020. Key search terms included multiple descriptors of advanced disease status in combination with "endometrial cancer" and "neoadjuvant chemotherapy". Our review included studies that examined survival and surgical outcomes of patients with stage III or IV endometrial cancer treated with neoadjuvant chemotherapy followed by interval cytoreductive surgery versus those who received primary cytoreductive surgery. We excluded studies examining only patients with leiomyosarcomas, carcinosarcomas, and stromal sarcomas due to the biologic heterogeneity of these malignancies.

Results: The nine included studies encompassed 5,844 patients, of which 1,317 (22.5%) received NACT and 4,527 received PCS (77.5%). With the exception of a single study, all were retrospective observational studies or case series. Use of NACT in patients with stage IV EC increased from 16.0% in 2010 to 23.9% in 2015. Five studies analyzed median overall survival and all but one reported no significant difference between NACT + ICS vs. PCS. Optimal cytoreduction (<1 cm of residual disease) rates were similar across both treatment groups in three separate analyses, however pooled data suggest improved rates of optimal cytoreduction for NACT + ICS vs. PCS patients (81.9% vs. 51.5% respectively). Patients receiving NACT experienced significantly shorter hospital admissions and lower operative times compared to PCS counterparts.

Conclusions: NACT followed by ICS reduces perioperative morbidity while offering similar overall survival.

1. Introduction

Endometrial cancer is the most common gynecologic cancer in developed nations and the fourth most common cancer in women (Lee et al., 2017; Rabinovich, 2016). Over 70% of endometrial cancer patients present with stage I disease which is associated with a greater than 90% five year survival rate (Galaal et al., 2016). Mortality markedly increases with advanced stages of disease; women with stages III and IV endometrial cancer experience five year survival rates of 56% and 17%, respectively (Siegel et al., 2019). Women with advanced endometrial cancer represent only 10–15% of all endometrial cancer cases, yet account for over half of all endometrial cancer deaths (Rabinovich, 2016; Palisoul and Mutch, 2016). Patients with inoperable endometrial cancer experience a median survival of only 2–8 months (Palisoul and Mutch, 2016).

Primary cytoreduction, followed by adjuvant chemotherapy and/or
radiotherapy, represents the current mainstay of treatment for advanced stage endometrial cancer (Wright et al., 2012). Cytoreduction has emerged as the most important component of treatment (Burke et al., 2014). In multiple analyses, patients with stage III or IV endometrial cancer who underwent optimal cytoreduction to <1 cm of residual disease experienced an overall survival benefit compared to women left with bulky residual disease after surgery (Bristow et al., 2001; Shih et al., 2011; Lambrou et al., 2004). Despite the survival benefit associated with optimal surgical cytoreduction, the procedure is associated with significant morbidity (Rabinovich, 2016). Over the last two decades the use of cytotoxic chemotherapy has progressively expanded (Cain et al., 2017; Deng et al., 2017; Wright et al., 2016; Blumenhhal et al., 2018; Grossman et al., 2003; Dehal et al., 2018). However, to date, there have been few reports of the use of primary, or neoadjuvant chemotherapy (NACT), for women with stage IV endometrial cancer (Tobias et al., 2020).

Over the same time period, use of NACT followed by interval debulking surgery has become an accepted treatment modality for advanced ovarian cancer (Wright et al., 2016). Data from four randomized controlled trials demonstrated the non-inferiority of NACT followed by interval cytoreduction compared to primary cytoreduction for stage IIB and IV ovarian cancer (Onda et al., 2008; Vergote et al., 2010; Keboe et al., 2015; Fagotti et al., 2016).

In light of these findings and the similarities in presentation between advanced ovarian and endometrial cancer, a re-evaluation of NACT for endometrial cancer is warranted. We performed a systematic review to examine the use and outcomes associated with NACT for metastatic endometrial cancer. Specifically, we examined surgical outcomes, use of chemotherapy, and survival to inform management of endometrial cancer patients who are precluded from primary cytoreduction.

2. Methods

2.1. Search strategy and information sources

This systematic review was conducted in accordance with the Preferred Reporting Items for Systematic reviews and Meta-analyses (PRISMA) guidelines (Liberati et al., 2009). A systematic search for articles published from January 1, 1990 to December 31, 2020 was performed in PubMed, Embase, Web of Science, and Scopus. Key search terms included descriptors of advanced disease status (“advanced stage”, “unresectable”, “metastatic”, “stage III”, “stage IV” and others) in combination with “endometrial cancer” and “neoadjuvant chemotherapy” (Supplemental Table 1). Where applicable, we searched using Medical Subject Heading (MeSH) terms “Neoadjuvant therapy”, “Endometrial neoplasms”, and “Uterine neoplasms” or other related exploding search terms. All literature searches were executed between March 3, 2020 and March 11, 2020.

2.2. Eligibility criteria

We sought to identify all reports that examined survival and surgical outcomes of patients with advanced endometrial cancer treated with neoadjuvant chemotherapy followed by interval cytoreductive surgery versus those who received primary cytoreductive surgery. Inclusion criteria were as follows: (1) patients with international FIGO stage III or IV endometrial cancer, (2) patients with endometrioid, clear cell, serous, and mixed histologic subtypes, (3) reporting of relevant survival outcomes, including overall survival, and (4) English language reports. We did not formulate selection criteria based on neoadjuvant chemotherapy regimens or cycle numbers as these parameters vary widely across practices. Reviews, case reports, and letters were excluded for systematic review. We excluded studies examining only patients with uterine leiomyosarcomas, carcinosarcomas, and stromal sarcomas due to the biological heterogeneity found in these malignancies.

2.3. Study selection and data extraction

All studies were screened for relevance by title and abstract by two independent reviewers (ABH and JW). Any disparities in opinion were resolved after both reviewers conferred and achieved consensus. The full texts of relevant studies were further screened by two independent reviewers (ABH and JW) according to the inclusion and exclusion criteria. Disparities in opinion were once again resolved after conference and consensus to produce a final group of included studies. The following data and descriptors were extracted from included studies: publication year, years of analysis, study type, endometrial cancer FIGO stages included, number of patients, histology, response to NACT, extent of debulking, progression to IDS or adjuvant chemotherapy, median overall survival (OS), and any other relevant additional findings.

2.4. Quality assessment

The NIH Study Quality Assessment Tools were used to evaluate each included study for the quality and risk of bias due to study design flaws or implementation (Study Quality Assessment Tools, 2020). Two independent reviewers (ABH and JW) scored each study according to NIH Study Quality Assessment Tool criteria and rated study quality as “good” (valid results with least risk of bias), “fair” (not enough bias to invalidate results), and “poor” (significant risk of bias). Disparities in ratings were resolved through discussion between the two reviewers.

2.5. Data synthesis

After literature search and review it was noted that there was significant heterogeneity among the identified studies. The decision was therefore made to review the studies descriptively rather than to perform a formal meta-analysis.

3. Results

3.1. Study selection

After systematic literature search, 1,213 unique articles were identified, of which 1,178 irrelevant articles were excluded during title and abstract screening. Of the thirty-five articles that underwent full-text screening, nine were selected for patient populations, study design, and outcomes appropriate to this systematic review. Twenty-six studies were excluded: thirteen for irrelevant study designs (review article, case report, or letters), five for lack of survival outcomes, four for non-English language formats, two for patients receiving salvage chemotherapy rather than NACT, one duplicate study, and one for including only patients with uterine carcinosarcomas (Fig. 1).

3.2. Study characteristics

The nine included studies encompassed 5,844 patients, of whom 1,317 received NACT and 4,527 received PDS (Table 1) (Tobias et al., 2020; Bogani et al., 2019; de Lange et al., 2019; Eto et al., 2013; Holman et al., 2017; Khouri et al., 2019; Rajkumar et al., 2019; Vandenput et al., 2009; Wilkinson-Ryan et al., 2015). Tobias et al. study demonstrated that use of NACT in patients with stage IV EC increased from 16.0% in 2010 to 23.9% in 2015. This study accounted for the largest sample size, including 4,890 patients (Tobias et al., 2020). With the exception of the study by Vandenput et al., all included studies were retrospective observational studies or case series. No randomized controlled trials examining NACT in advanced endometrial cancer were found. In the only prospective study included, Vandenput et al. enrolled thirty patients to receive three to four cycles of NACT following laparoscopic confirmation of intraabdominal spread of disease (stage IV). (Vandenput et al., 2009)

Nearly all of the studies examined only patients with FIGO stage III or...
IV EC. The majority of NACT patients received three to six cycles (range: 1–8) of a taxane and platinum agent, although NACT regimens were highly variable between and within studies (Table 1). Three of the nine studies exclusively examined patients with serous carcinomas. Six of the studies included a comparator PDS arm, while the remaining three were case-series lacking a PCS arm. Bogani et al. and Tobias et al. selected a comparator arm via propensity score adjustment to account for allocation bias (Tobias et al., 2020; Bogani et al., 2019).

3.3. Quality assessment

Quality assessment and critical appraisal of included studies was conducted according to the NIH Study Quality Assessment Tools (Appendix 1). Separate assessment criteria were applied according to study design. Five studies were judged to be at low risk of bias, three studies at some risk of bias, and one study at high risk of bias. Major sources of bias include widely varying NACT regimens across studies (inconsistently implemented exposure measure) and lack of sample size or power description among studies.

3.4. Chemotherapeutic and surgical outcomes

Major chemotherapeutic and surgical outcomes of interest included response to NACT, performance of ICS, and extent of cytoreductive surgery are summarized in Table 2. Five studies reported a total of 257 patient responses to NACT, of which 178 (69%) experienced at least a partial response, 25 (10%) experienced stable disease, and 46 (18%) experienced progression of disease. Notably, Eto et al. reported NACT responses only for those patients that underwent ICS. Holman et al. found that patients receiving NACT experienced a significantly lower complete response rate than those receiving primary surgery with adjuvant chemotherapy and/or radiotherapy, which likely reflects selection bias inherent in nonrandomized, retrospective observational studies. (Holman et al., 2017) Wilkinson-Ryan et al. examined changes in CA-125 as proxies for responses to therapy, and observed no significant difference in treatment response between NACT and PDS cohorts at the conclusion of therapy (mean CA-125 decline of 80% and 92.5% respectively, \( P = 0.43 \)). (Wilkinson-Ryan et al., 2015)

All nine studies reported the proportion of NACT patients receiving ICS (Table 2). In the studies by Bogani et al., Rajkumar et al., and Wilkinson-Ryan et al., receipt of IDS was an inclusion criterion for patients in the NACT arm. Excluding patients from the aforementioned studies, 58.9% of NACT patients overall received ICS, driven largely by the sample size of Tobias et al. The most commonly reported contraindications to ICS included disease progression and/or surgically unresectable tumor burden, although reasons for exclusion from IDS were infrequently reported. (de Lange et al., 2019; Khouri et al., 2019; Vandenput et al., 2009)

The extent of cytoreductive surgery was classified as complete (no gross residual disease), optimal (<1cm residual disease), and suboptimal (greater than1cm residual disease). Six studies reported surgical outcomes for patients receiving NACT followed by ICS, encompassing 204 patients in total. Of these patients, 167 (81.9%) received complete/optimal cytoreduction and 37 (18.1%) received suboptimal cytoreduction. Only three studies compared surgical outcomes between NACT with ICS and PCS arms, encompassing 328 PCS patients in total. Of these PCS patients, 169 (51.5%) received complete/optimal cytoreduction and 159 (48.5%) received suboptimal cytoreduction.

Vandenput et al. reported the highest rate of complete cytoreduction (92%) among patients receiving NACT with ICS. Cytoreduction rates were similar between NACT + ICS and PCS patients in the analysis by Bogani et al. (\( P = 0.96 \)). Eto et al. compared rates of optimal/complete cytoreduction between NACT with ICS and PCS arms and found no significant difference (57% vs. 45%, \( P = 0.087 \)). Wilkinson-Ryan et al. similarly concluded that rates of complete (70% NACT + ICS vs. 32% PCS) and optimal (30% NACT with ICS vs. 50% PCS) cytoreduction were
3.5. Survival outcomes

Several studies conducted novel subgroup analyses, highlighted in Table 3. Both de Lange et al. and Rajkumar et al. demonstrated increasing survival in women who received more extensive cytoreductive surgeries (de Lange et al., 2019; Rajkumar et al., 2019). De Lange et al. reported a median OS of 41, 16 and 13 months for women who received complete/optimal, suboptimal, and no cytoreductive surgery, respectively. Receipt of ICS improved median OS from 7 months for women receiving NACT without ICS to 21 months for those receiving NACT with ICS in the analysis by Eto et al.

4. Discussion

These data suggest that use of a strategy of neoadjuvant chemotherapy with interval cytoreductive surgery is increasingly common among women with advanced stage endometrial cancer. Compared to

| Reference          | Study Design       | Years Analyzed | Stages Included | NACT Regimen* | NACT Histologies NACT | PDS Histologies PDS |
|--------------------|--------------------|----------------|-----------------|---------------|------------------------|----------------------|
| Bogani et al., 2019| Retrospective cohort, propensity-matched | 2005–2016 | IVb             | 3-6 cycles paclitaxel/cisplatin or doxorubicin ± carboplatin | 15 15 (100) serous | 15 15 (100) serous |
| de Lange et al., 2019| Retrospective case-series | 2005–2014 | III, IV         | 1-6 cycles paclitaxel/cisplatin ± doxorubicin | 102 44 (43) endometrioid | – – |
| Eto et al., 2013  | Retrospective cohort | 1996–2005 | IVb             | 63% taxane + platinum 30% doxorubicin + platinum ± other 7% other | 125 68 (54) endometrioid 57 (46) non-endometrioid | 279 163 (58) endometrioid 116 (42) non-endometrioid |
| Holman et al., 2017 | Retrospective cohort | 1993–2012 | III, IV | NACT details not reported | 27 19 (70) serous | 233 79 (34) serous |
| Khouri et al., 2019 | Retrospective case-series | 2002–2016 | NR             | 85% paclitaxel/cisplatin 10% paclitaxel only 5% paclitaxel only | 39 11 (28) endometrioid 17 (44) serous | – – |
| Rajkumar et al., 2019 | Retrospective cohort | 2010–2016 | IIIC, IV       | 3-6 cycles taxol/carboplatin or cisplatin/doxorubicin | 17 NR | 28 NR |
| Tobias et al., 2020 | Retrospective cohort | 2010–2015 | IV               | NR                | 952 223 (23) endometrioid 249 (26) serous 34 (4) clear cell 115 (12) carcinosarcoma 90 (9) sarcoma 248 (21) serous 117 (3) clear cell 612 (16) carcinosarcoma 554 (14) sarcoma | 3938 1114 (28) endometrioid 628 (21) serous |
| Vandenput et al., 2009 | Prospective cohort | 1999–2007 | IV             | 3-4 cycles paclitaxel/carboplatin or doxorubicin/cisplatin | 30 2 (7) endometrioid 27 (90) serous 1 (3) clear cell | – – |
| Wilkinson-Ryan et al., 2015 | Retrospective cohort | 2000–2013 | IV             | 3-8 cycles taxane/carboplatin | 10 10 (100) serous or mixed histology with >50% serous component | 34 34 (100) serous or mixed histology with >50% serous component |

NACT = neoadjuvant chemotherapy, IDS = interval debulking surgery, PD = primary debulking surgery.
EM NOS = endometrioid not otherwise specified.
NR = not reported.

* Summaries of the most common NACT regimens are listed, as studies reported highly variable NACT regimens.

similar between treatment groups (P = 0.10).

Patients who received NACT experienced significantly shorter hospital admissions compared to PCS counterparts in the studies of Bogani et al. (4 vs. 6 days, P = 0.011) and Wilkinson-Ryan et al. (3 vs. 5 days, P = 0.0021). Wilkinson-Ryan et al. additionally reported significantly lower operative times in patients who received NACT with ICS compared to those who underwent PCS (136.9 vs. 202.6 min, P = 0.025), Bogani et al. noted similar trends (127 vs. 177.6 min, P = 0.072).
primary cytoreduction, NACT with ICS appears to be associated with improved rates of optimal cytoreduction, decreased perioperative morbidity, and decreased operative times. While the relationships between NACT and PCS are complex, patients undergoing NACT followed by ICS experience similar overall survival compared to those receiving PCS.

Neoadjuvant therapy has been employed successfully in a spectrum of solid tumors, often as a technique for reducing surgical morbidity and improving the extent of cytoreduction, or as a method of downstaging advanced disease. In a large retrospective study of patients in the National Cancer Database (NCDB), Dehal et al. demonstrated improved 3-year survival for patients with T4b colon cancer (invasion of extracolonic organs), but no benefit for patients with T3 or T4a disease (Dehal et al., 2018). A RCT comparing NACT vs. PCS in 317 patients with muscle-invasive bladder cancer found a 33% mortality reduction in the NACT arm, attributed largely to the ability of NACT to downstage disease (Grossman et al., 2003). In a meta-analysis of seven RCTs by Fu et al., patients with esophageal adenocarcinoma experienced improved overall survival (HR 0.74; 95% CI, 0.63–0.88) with neoadjuvant chemoradiation therapy (Fu et al., 2015). In the setting of breast cancer, NACT is most commonly used in early, rather than advanced, disease, allowing for the possibility of breast-conserving surgery (Cain et al., 2017). Collectively, these studies highlight the importance of careful patient selection in optimizing survival benefits from NACT. Despite limited data, it appears that use of NACT is increasing for women with stage IV endometrial cancer (Tobias et al., 2020).

Gynecologic oncologists have increasingly employed NACT followed by ICS in advanced ovarian cancer. Pooled analysis of the EORTC 55,971 and CHORUS trials demonstrated similar overall survival in women with stages IIIA-IV ovarian cancer treated with NACT vs. PCS. (27.6 vs. 26.9 months, respectively) (Vergote et al., 2010, 2018). The SCORPION and JCOG0602 trials further demonstrated decreased perioperative morbidity, operative times, and length of hospital stay with NACT + ICS (Fagotti et al., 2016; Onda et al., 2016). Given these findings, the Society of Gynecologic Oncology and the American Society of Clinical Oncology endorse NACT for patients with advanced ovarian cancer who are deemed at high risk of perioperative morbidity or have low probability of cytoreduction to <1 cm of residual disease with PCS (Wright et al., 2016). Endometrial cancer resembles ovarian cancer in its propensity for peritoneal spread, and has therefore drawn focused interest as a disease site candidate for NACT (Bogani et al., 2019; Holman et al., 2017; Wilkinson-Ryan et al., 2015). Given the poor overall outcomes of women with metastatic endometrial it is unclear why NACT has not received more attention in this setting. Further, many women with endometrial cancer are elderly with significant comorbidities, a population that may derive the most benefit from decreased perioperative

### Table 2

| Study               | NACT n | PDS n | Response to NACT | NACT patients receiving IDS n (%) | Extent of Debuling Surgery NACT+IDS, n (%) | Additional Findings |
|---------------------|--------|-------|-------------------|----------------------------------|------------------------------------------|---------------------|
| Bogani et al., 2019| 15     | 15    | NR                | 15 (100)                         | 14 (93) complete 1 (7) optimal           | p = 0.96 for debulking outcomes |
|                     |        |       |                   |                                  | 13 (87) complete 2 (13) optimal         | • NACT arm with significantly shorter hospital stays (4 vs. 6d, p = 0.011) |
|                     |        |       |                   |                                  |                                          | • NACT arm trend towards shorter op. time (127 vs. 177.6, p = 0.072) and trend towards lower transfusion rates (6.6% vs. 33.3%, p = 0.067) |
| de Lange et al., 2019| 102   | –     | 4 (4) CR 73 (72) PR 9 (9) SD 11 (11) PD 2 (2) died during treatment 3 (3) no imaging available | 80 (78) | 48 (60) complete 23 (29) optimal 9 (11) suboptimal | p = 0.087 for optimal cytoreduction |
| Eto et al., 2013    | 125    | 279   | 40 (68) CR/PR 9 (15) SD 7 (12) PD | 59 (47) | 19 (32) complete 15 (25) optimal 25 (43) suboptimal | p = 0.087 for optimal cytoreduction |
| Holman et al., 2017 | 27     | 233   | 8 (30) CR 9 (33) PR/SD 10 (37) PD | 17 (63) | 17 (63) NR | p = 0.010 for debulking outcomes |
| Khouri et al., 2019 | 39     | –     | 22 (56) PR 1 (3) SD 16 (41) PD | 16 (41) | 16 (41) NR | • NACT arm with significantly shorter mean operative time (136.9 vs. 202.6 min., p = 0.025) and length of hospital stay (3 vs. 5 d., p = 0.0021) |
| Rajkumar et al., 2019 | 17   | 28    | NR                | 17 (100)                         | 7 (70) complete 3 (36) optimal           | • NACT arm with trend towards less blood loss (410ml vs. 781.8ml, p = 0.063) |
| Tobias et al., 2020 | 952    | 3938  | NR                | 555 (58)                         | 11 (32) complete 6 (18) suboptimal       | – |
| Vandenput et al., 2009 | 30   | –     | 2 (7) CR 20 (67) PR 6 (20) SD 2 (7) PD | 24 (80) | 22 (92) complete 2 (8) optimal | – |
| Wilkinson-Ryan et al., 2015 | 10  | 34    | Average decline in CA 125 post-NACT ~ 91 ± 11% | 10 (100) | 7 (70) complete 3 (36) optimal | – |

CR = complete response, PR = partial response, SD = stable disease, PD = progressive disease, NR = not reported.
Table 3
Survival outcomes and additional findings of included studies.

| Reference              | Median OS (months) | Additional findings                                                                 |
|------------------------|--------------------|--------------------------------------------------------------------------------------|
|                        | NACT Overall       | NACT + IDS subgroup                                                                 |
|                        |                    |                                                                                      |
|                        |                    | Median OS (months)                                                                 |
|                        |                    | arms cross after 8 months                                                             |
| Bogani et al., 2019    | –                  | 16.7                                                                                 |
| de Lange et al., 2019  | 27                 | 21                                                                                   |
|                        | 18                 | p = 0.35                                                                              |
| Eto et al., 2013       | 12                 | 21                                                                                    |
| Holman et al., 2017    | 20.4               | 16                                                                                   |
|                        | NR                 | p = 0.84 for NACT + IDS vs. PDS                                                       |
| Khouri et al., 2019    | NR                 | 16                                                                                    |
|                        | 22.5               | p = 0.57, HR = 1.26, 95% CI (0.56-2.85)                                               |
| Rajkumar et al., 2019  | –                  | 29                                                                                    |
| Tobias et al., 2020    | 19                 | 25                                                                                    |
|                        | (95% CI: 17-21)    | (95% CI: 24-27)                                                                      |

Table 3 (continued)

| Reference              | Median OS (months) | Additional findings                                                                 |
|------------------------|--------------------|--------------------------------------------------------------------------------------|
|                        | NACT Overall       | NACT + IDS subgroup                                                                 |
|                        |                    |                                                                                      |
|                        |                    | Median OS (months)                                                                 |
|                        |                    | arms cross after 8 months                                                             |
| Vandenput et al., 2009 | 23                 | NR                                                                                   |
|                        | 17.3               | 20.7                                                                                 |
| Wilkinson-Ryan et al., 2015 | –          | 17.3                                                                                 |

NACT = neoadjuvant chemotherapy, PDS = primary debulking surgery, adj. CT = adjuvant chemotherapy, adj. RT = adjuvant radiotherapy, NR = not reported.

Our findings suggest that NACT followed by ICS reduces perioperative morbidity while offering similar overall survival. The findings of the large analysis by Tobias et al. are intriguing as survival was initially superior for patients who received NACT but among patients who survived longer, overall survival was more favorable for those who underwent PCS. These findings indicate that PCS may be associated with significant perioperative morbidity and mortality or delay in receipt of chemotherapy. However, for those patients who can undergo primary cytoreduction without undue morbidity and go on to receive chemotherapy, long term outcomes may be more favorable. NACT may also be useful in identifying chemotherapy resistant disease, allowing early optimization of end-of-life care. Developing algorithms or clinical trials to differentiate patients who will benefit most from NACT vs. PCS will be critical in supporting clinical decision-making for advanced EC.

We acknowledge several important limitations in this systematic review. The majority of studies included were retrospective studies with relatively small cohorts. No randomized controlled trials have been performed examining NACT in EC, thus limiting the strength of the clinical recommendations made on the basis of existing evidence. Second, conclusions about survival drawn from this review are skewed by the large sample size of Tobias et al., which accounts for 72% of NACT patients and 87% of PCS patients included in our pooled cohort. However, surgical outcomes, perioperative morbidity, and chemotherapeutic outcomes were unaffected by Tobias et al., which focused only on time-varying survival outcomes. Third, NACT regimens varied widely between studies in terms of agents used and number of cycles administered, which prevents our study from commenting on the efficacy of different regimens. Similarly, included studies examined outcomes across many histologic subtypes of EC. There are undoubtedly genetic differences that carry implications for treatment across EC subtypes, thus this review can only offer insight into NACT for EC patients as a whole (Chang et al., 2019).

Our review suggests that NACT followed by ICS represents a viable alternative to PCS in select patients with advanced EC. Those patients who receive ICS experience improved cytoreduction and decreased perioperative and postoperative complications. The current evidence rests solely on small retrospective cohort studies and a single, large observational analysis. Based on the findings of our paired meta-analysis, every attempt should be made to perform complete cytoreduction in women with metastatic endometrial cancer chosen for PCS. Further characterization of surgical, survival, and chemotherapeutic outcomes, ideally through a multi-institutional RCT, will be necessary to refine an algorithm for NACT candidate selection. Further, as therapy for endometrial cancer evolves and incorporates targeted therapeutics, further work to characterize outcomes of women treated...
with NACT will be essential.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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Appendix A. Supplementary material

Supplementary data to this article can be found online at https://doi.org/10.1016/j.gyno.2021.100887.

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