

卵巣がん進行の緩徐化 (Abstract LBA5503)

経静脈投与に腹腔内化学療法を追加することにより卵巣がんの進行が緩徐になる

Adding intraperitoneal chemotherapy to intravenous therapy slows progression of ovarian cancer

手術が成功し、経静脈投与 (IV) 化学療法に加え腹腔内 (IP) 化学療法を施行された一部の進行卵巣がん女性においては、IV化学療法のみへの施行に比べ効果が高いようである。手術前に最初に化学療法を施行された女性においては、IPおよびIV化学療法を施行された患者の23.3%が9か月後に進行したのに対し、IV化学療法のみ患者におけるその割合は42.2%であった。このスタディ結果は2016年American Society of Clinical Oncology年次集会で発表された。

Full Text

For some women with advanced ovarian cancer that was successfully treated surgically, delivering chemotherapy intraperitoneal (IP) as well as intravenously (IV) appears more effective than IV chemotherapy alone. For women who were initially treated with chemotherapy prior to surgery (e.g., neoadjuvant therapy), the initial results from a randomized phase II trial show that 23.3% of women who received IP and IV chemotherapy had disease progression at nine months, vs. 42.2% of those who received IV chemotherapy alone.

The study was presented at the 2016 American Society of Clinical Oncology (ASCO) Annual Meeting.

According to the authors, the proportion of women with ovarian cancer who receive neoadjuvant chemotherapy prior to surgery is growing. Women who undergo optimal debulking surgery following this approach may now be candidates for IP/IV combination chemotherapy.

IP chemotherapy allows the delivery of higher doses of chemotherapy to the tumor, while sparing other parts of the body from side effects. Several prior randomized clinical trials showed that IP chemotherapy improved outcomes for certain women with ovarian cancer. However, this is the first randomized study to explore the benefit of IP chemotherapy among women who had received neoadjuvant chemotherapy.

"At this early time frame, we already see that women are doing better with IP chemotherapy, without a significant difference in toxicity," said lead study author Helen Mackay, MD, Divisional Head of Medical Oncology and Hematology at the Sunnybrook Odette Cancer Centre in Toronto, Canada. "However, women should consider the side effects of IP and IV chemotherapy, as well as recovery from cancer surgery, when discussing this option with their doctors."

This randomized phase II trial compared the efficacy and side effects of two combination chemotherapy regimens in patients with stage IIB - IV epithelial ovarian cancer. The majority (82%) of women had stage IIIC disease.

In this study, 275 women received neoadjuvant platinum - based chemotherapy, followed by surgery to remove their ovarian cancer. Following debulking surgery, 200 were randomly assigned to treatment with IV chemotherapy or an IV/IP regimen.

At nine months, 42.2% of women who received IV chemotherapy had disease worsening compared to 23.3% of those treated with IP/IV chemotherapy. The median progression-free survival was similar between the two groups - 11.3 months with IV chemotherapy and 12.5 months with the IV/IP regimen. The median overall survival was longer with IV/IP therapy than with IV therapy alone (59.3 months vs. 38.1 months), but the difference was not statistically significant.

"Although this randomized phase II trial was not statistically powered to evaluate survival, our results offer information on how to incorporate IP chemotherapy when women receive neoadjuvant chemotherapy followed by debulking surgery," said Dr. Mackay. "The findings also offer supportive and additional information to the previous published adjuvant randomized trials that showed an improvement in overall survival when IP chemotherapy was given following initial optimal debulking surgery."

The rate of severe side effects was slightly lower among women who receive IP/IV chemotherapy (16% vs. 23%), but this difference was not statistically significant.

Prior research has suggested that some molecular subtypes of ovarian cancer are more sensitive to chemotherapy than others. The researchers plan to assess tissue samples collected during this study to see if certain biologic characteristics were associated with improved outcomes with IP vs. IV chemotherapy. "If we can identify the long term survivors, we hope this will help us better predict who truly benefits from this approach," said Dr. Mackay.

"Intraperitoneal chemotherapy is an effective yet underused treatment for women with newly diagnosed ovarian cancer that has been successfully removed surgically. These data now suggest that IP treatment may have a role in the postoperative setting for women who initially were treated with intravenous chemotherapy," said Don Dizon, MD, FACP, ASCO Expert in ovarian cancer. "Furthermore, this study provides reassurance for patients and providers that the carboplatin-based IP regimen is both effective and well tolerated with maintenance of quality of life. That said, we need to further define those who derive the greatest benefit from this approach and to identify better options for all women with ovarian cancer."

In 2012, 239,000 women were diagnosed with ovarian cancer worldwide. Due to lack of screening and specific symptoms, most women already have late-stage disease at the time of their diagnosis.

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