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CRITICAL CARE MANAGEMENT FOLLOWING CYTOREDUCTIVE SURGERY WITH HYPERTHERMIC INTRAPERITONEAL CHEMOTHERAPY: NOT ROUTINELY INDICATED

¹SL Aronson, ²RM van Stein, ³FJ Hendriks, ⁴K Sikorska, ⁵API Houwink, ⁵PFE Schutte, ³CD de Kroon, ⁶GS Sonke, ²WJ van Driel. ¹Gynaecological Oncology, Medical Oncology, Netherlands Cancer Institute – Antoni van Leeuwenhoek, Amsterdam, Netherlands; ²Gynaecological Oncology, Netherlands Cancer Institute – Antoni van Leeuwenhoek, Amsterdam, Netherlands; ³Gynecology, Leiden University Medical Center, Leiden, Netherlands; ⁴Biometrics, Netherlands Cancer Institute – Antoni van Leeuwenhoek, Amsterdam, Netherlands; ⁵Anesthesiology and Intensive Care, Netherlands Cancer Institute – Antoni van Leeuwenhoek, Amsterdam, Netherlands; ⁶Medical Oncology, Netherlands Cancer Institute – Antoni van Leeuwenhoek, Amsterdam, Netherlands

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Introduction/Background Hyperthermic intraperitoneal chemotherapy (HIPEC) is increasingly used for patients with stage III ovarian cancer undergoing interval cytoreductive surgery (CRS). It is uncertain whether routine postoperative admission to an intensive care setting following CRS-HIPEC for ovarian cancer is necessary. We estimated the incidence of patients requiring critical care support and tried to identify patients in whom admission to an intensive care setting can be safely omitted.

Methodology We analyzed 154 patients with primary ovarian cancer, who underwent CRS-HIPEC between 2007–2021 in two Dutch HIPEC-centers. Patients were routinely transferred to an Intensive Care Unit (ICU) or Post Anesthesia Care Unit (PACU). Patients requiring critical care support were identified by predefined criteria based on respiratory, circulatory, and metabolic parameters. Logistic regression analyses with backward selection were used to predict the need for critical care support in individual patients and the are-under-the-ROC-curve (AUC) of the model was estimated.

Results Median ICU/PACU length of stay was 21 hours (IQR 19–29) and 38% of patients received postoperative critical care support, mainly consisting of hemodynamic interventions (37%). Independent predictors for critical care support are age, blood loss, norepinephrine dose during surgery, and peritonectomy extent (table 1). AUC of the model is 0.81 (95% CI 0.73–0.88). Using a 20% cut-off to define low-risk of critical care support, 37% of patients would be eligible to forego ICU/PACU admission.

Abstract 2022-RA-1451-ESGO Table 1 Multivariable logistic regression analysis for probability of critical care support (N=154, events=58)

| Independent predictors | OR | 95% CI | P-value | B |
|-------------------------------------------------|------|------------|---------|------|
| Age > 70 years | 4.79 | 1.93-11.91 | 0.00 | 1.57 |
| Blood loss (liter) | 2.16 | 1.08-4.32 | 0.03 | 0.77 |
| Norepinephrine dose during surgery (µg/kg/hour) | 1.49 | 1.22-1.82 | 0.00 | 0.40 |
| Extensive peritonectomy (in ≥ 2 regions) | 3.74 | 1.25-11.20 | 0.02 | 1.32 |

OR: Odds ratio, CI: confidence interval

Conclusion Postoperative admission to an intensive care setting is not routinely required for ovarian cancer patients undergoing CRS-HIPEC. Following prospective validation, a decision tool based on pre- and intra-operative parameters can help to identify low-risk patients.

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CHEMOTHERAPY RESPONSE SCORE AS A PREDICTOR OF SURVIVAL AMONG PATIENTS UNDERGOING INTERVAL DEBULKING SURGERY FOR OVARIAN CANCER

¹Ioannis Rodolakis, ²Michalis Liotos, ¹Konstantina Papadatou, ¹Anastasia Prodromidou, ¹Dimitrios Haidopoulos, ¹Alexandros Rodolakis, ²Aristotelis Bamias, ¹Nikolaos Thomakos. ¹First department of obstetrics and gynaecology, National and Kapodistrian University of Athens, ATHENS, Greece; ²National and Kapodistrian University of Athens, ATHENS, Greece

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Introduction/Background Neo-adjuvant chemotherapy has been adopted as an alternative mode of therapy for surgically irresectable ovarian cancer in cases of diffuse dissemination, where primary debulking surgery is not feasible or when patient status does not allow extensive procedures. The response to chemotherapy can be evaluated objectively with the use of standard pathology. In the present study we evaluated the prognostic significance of chemotherapy response score in predicting survival rates of patients undergoing interval debulking surgery.

Methodology The study is based in a retrospective cohort of patients. We collected data from 48 ovarian cancer patients that received at least 3 cycles of neo-adjuvant chemotherapy. The evaluation of chemotherapy response score was based on pathology sections of the omentum and ovaries. Following interval debulking surgery chemotherapy was continued until the completion of 6 cycles of perioperative treatment. Twenty two patients received maintenance therapy with bevacizumab following completion of chemotherapy.

Results Median follow-up was 52.5 months ranging between 38.5 and 70.1 months. Agreement rates of chemotherapy rates among omental and ovarian biopsies were moderate (CRS 1 22.9% vs 37.5% respectively, CRS 2 37.5% vs 35.4% and CRS 3 33.3% vs 16.7%). Progression free survival rates gradually declined among patients with omental CRS 3 and those with CRS 1 (18.7 vs 14 vs 10.3 months respectively, $p=.003$). Similar results were observed for overall survival rates, however, the results were not statistically significant (42.3 vs 32 vs 29.3 months respectively, $p=.182$).

Conclusion Evaluation of the chemotherapy response score from omental biopsies is an accurate predictor of survival rates of ovarian cancer patients undergoing interval debulking surgery, irrespective of the use of maintenance therapy. Further studies are needed to support our findings.

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OVARIAN CANCER METASTASES IN THE LIVER AREA: A RETROSPECTIVE ANALYSIS OF SURGICAL, INTRAOPERATIVE AND POSTOPERATIVE OUTCOMES ACCORDING TO A STANDARDIZE ANATOMO-SURGICAL CLASSIFICATION

¹Andrea Rosati, ²Matteo Pavone, ²Antonella de Palma, ¹Carmine Conte, ¹Valentina Ghirardi, ³Agostino Maria de Rose, ^{3,2}Felice Giuliante, ^{1,2}Giovanni Scambia, ^{1,2}Anna Fagotti. ¹Department Woman and Child Health and Public Health, Fondazione Policlinico Universitario A. Gemelli IRCCS, Rome, Italy; ²Università Cattolica del Sacro Cuore, Rome, Italy; ³Hepatobiliary Surgery Unit, Fondazione Policlinico Universitario A. Gemelli IRCCS, Rome, Italy

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