

ADAM19, Ki-67, and Caspase-3) and chemoresistance test (to carboplatin chemotherapy).

**Results** It was found that the number of spheroids obtained, all gene expression, and number of chemoresistance to carboplatin regimen in CD133+ CSC cultures were higher than the main population and CD133-. CSCs with CD133+ had a higher ability to proliferate with increased Ki-67 gene expression, stronger stemness with higher NANOG gene expression, and greater chemoresistance ability with increased ATM and ATR gene.

**Conclusion/Implications** It can be concluded that ATM and ATR gene expression are positively correlated with the resistance of CSC in ovarian cancer patients.

EP222/#246

### QUANTIFYING PLATINUM SENSITIVITY AS A MARKER FOR RESPONSE TO PARP INHIBITION IN PATIENTS WITH ADVANCED OVARIAN CANCER

<sup>1</sup>Anne-Marie Bergeron\*, <sup>2</sup>Marta Kisiel, <sup>1</sup>Melissa Laverchia, <sup>1</sup>Vanessa Carlson, <sup>1</sup>Clare Reade, <sup>1</sup>Lua Eiriksson, <sup>1</sup>Julie Nguyen, <sup>1</sup>Waldo Jimenez, <sup>2</sup>Alice Lytwyn, <sup>1</sup>Andra Nica. <sup>1</sup>Juravinski Cancer Centre, McMaster University, Hamilton, Gynecologic Oncology, Hamilton, Canada; <sup>2</sup>Juravinski Cancer Center, Pathology and Molecular Medicine, Hamilton, Canada

10.1136/ijgc-2023-IGCS.300

**Introduction** All patients with high grade epithelial ovarian cancer(HGEOC) do not benefit equally from PARP inhibitors, but all are exposed to PARP-associated toxicities. This study aims to assess the correlation between the pathology-based Chemotherapy Response Score(CRS) at the time of interval debulking surgery(IDS) and progression free survival (PFS) in patients who received PARP maintenance, to determine this score's potential as a marker of expected benefit from PARP.

**Methods** This is a retrospective cohort study of patients with HGEOC who underwent IDS between January 2016 and September 2022. Demographic and clinical parameters were collected.  $\chi^2$  test and Student t-test were used to compare descriptive variables and Kaplan-Meier survival analysis with log rank test comparison for PFS.

**Results** On 169 patients, 47 received PARP maintenance and the majority needed dose reduction due to toxicity(53.2%). Patients with CRS 1(No/Minimal response) or CRS 2-3(Moderate/Complete response) were comparable in terms of baseline characteristics. Patients CRS 1 compared to CRS 2-3 had lower PFS regardless of maintenance( $p=0.017$ ). Patients with CRS 2-3 who received PARP showed significantly improved PFS(20 vs 15 months,  $p=0.029$ ) compared to those who did not, while in those with CRS 1 maintenance was not associated with improved PFS( $p=0.27$ ). Results were similar on multivariate analysis, adjusting for BRCA status and surgical outcomes.

**Conclusion/Implications** In HGEOC patients demonstrating response(CRS 2-3) to NACT, PARP maintenance was associated with a significant improvement in PFS. CRS can be a helpful tool in counseling prior to PARP inhibitor initiation, in patients BRCA-intact, and in settings where homologous recombination deficiency testing is not easily available.

EP223/#840

### KAZAKHSTAN NATIONAL EXPERIENCE: GRANULOSA CELL TUMORS OF THE OVARY

<sup>1</sup>Dilyara Kaidarova, <sup>2</sup>Raikhan Bolatbekova\*, <sup>3</sup>Jubilee Brown, <sup>3</sup>Robert Naumann, <sup>2</sup>Aisulu Sarменова, <sup>2</sup>Askar Aidarov, <sup>2</sup>Arailym Akkassova, <sup>2</sup>Gulnur Bagatova, <sup>2</sup>Diana Zhaksylykova, <sup>2</sup>Aisulu Yestayeva. <sup>1</sup>Kazakh Institute of Oncology and Radiology, Chairman of The Board, Almaty, Kazakhstan; <sup>2</sup>Almaty Oncology Center, Gynecologic Oncology, Almaty, Kazakhstan; <sup>3</sup>Levine Cancer Institute at Atrium Health, Division of Gynecologic Oncology, Charlotte, USA

10.1136/ijgc-2023-IGCS.301

**Introduction** Objectives: Ovarian granulosa cell tumors (OGCT) are rare, and Eurasian data have not been published. This study objective was to describe OGCT among the 19 million residents of Kazakhstan.

**Methods** The Kazakhstan Cancer Registry Database was queried for descriptive and outcomes data of all consecutive patients with histologically verified OGCT from 2014-2020. Descriptive statistics and log likelihood ratios were performed using JMP Version 14.0.

**Results** 240 patients with OGCT were included, representing 3.9% of ovarian cancer in Kazakhstan. The median age was 52 years (range, 15-87 years). Nationality of origin was 55% Kazakh, 30.8% Russian, 3.8% Ukrainian, and 11% other. Stage distribution was 53.7% Stage I, 20.4% Stage II, 22% Stage III, 2.5% Stage IV, and 1% unknown. In total, 89 patients (37.6%) received chemotherapy; this did not correlate with stage. Common regimens included paclitaxel/carboplatin; bleomycin, etoposide, and cisplatin (BEP); and EP. Of the 240 patients, 67 patients (28%) recurred; recurrence correlated with stage ( $p<0.001$ ). Treatment for recurrent disease included surgery, chemotherapy, and radiation therapy. After 16-month median follow up (range, 0-90 months), 186 patients (77.5%) were without evidence of disease, 12 patients (5%) were alive with disease, and 42 patients (17.5%) had died. Risk of death increased with advancing stage ( $p<0.001$ ). Stage I OGCT patients who received adjuvant chemotherapy were significantly less likely to die than those who had not ( $p=0.003$ ).

**Conclusion/Implications** This is the first description of OGCT in Kazakhstan. There is a survival advantage to chemotherapy administration in early-stage patients, supporting the importance of access to chemotherapy.

EP224/#880

### BRCA MUTATIONS IN HIGH GRADE SEROUS OVARIAN CANCER IN KAZAKHSTAN

<sup>1</sup>Dilyara Kaidarova, <sup>2</sup>Raikhan Bolatbekova\*, <sup>3</sup>Robert Naumann, <sup>3</sup>Jubilee Brown, <sup>4</sup>Yerlan Kukubassov, <sup>3</sup>Tatyana Goncharova, <sup>4</sup>Madina Orazgaliyeva, <sup>2</sup>Askar Aidarov, <sup>4</sup>Saniya Osikbayeva, <sup>4</sup>Alima Satanova, <sup>4</sup>Dauren Kaldybekov, <sup>4</sup>Sanzhar Ismailov. <sup>1</sup>Kazakh Institute of Oncology and Radiology, Chairman of The Board, Almaty, Kazakhstan; <sup>2</sup>Almaty Oncology Center, Gynecologic Oncology, Almaty, Kazakhstan; <sup>3</sup>Levine Cancer Institute at Atrium Health, Division of Gynecologic Oncology, Charlotte, USA; <sup>4</sup>Kazakh Institute of Oncology and Radiology, Oncogynecology, Almaty, Kazakhstan

10.1136/ijgc-2023-IGCS.302

**Introduction** Objectives: More than 1,000 new cases and 500 deaths from ovarian cancer are detected annually in Kazakhstan. The aim of this study was to examine the BRCA1/2