

Stratifying by stage the residual tumour (OR=2.4;  $p=0.0001$ ), age (OR=1.9  $P=0.0001$ ), and the performance status (OR=1.2;  $p=0.03$ ) resulted as independent survival prognostic factors according to Cox multivariate analysis.

**Conclusion\*** Our data suggest that patients aged  $\geq 70$  can tolerate radical surgical treatments in the same way as younger patients without a significant increase in morbidity and, obviously, without ignoring the appropriate geriatric precautions. Furthermore, maximal surgical effort with optimal cytoreduction should be considered the gold standard regardless of age.

Therefore, our data underlines the importance of managing these patients within Gynecologic Oncology units equipped with a multidisciplinary team.

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### OVARIAN CANCER AND BRCA1 AND 2 GERMLINE MUTATIONS – THE PORTUGUESE EXPERIENCE

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10.1136/ijgc-2021-ESGO.458

**Introduction/Background\*** Ovarian cancer (OC) is the second most common gynecologic malignancy in developed countries and the third most common in developing countries. Approximately 13 to 15 percent of OC are attributable to heritable mutations in *BRCA1* and 2.

This study aims to: I- assess median overall survival (mOS); II- characterize patients (pts) with OC assessed in a Family Risk Consultation (FRC).

**Methodology** This is a multicentric, descriptive and retrospective study of pts with OC followed at the FRC between 2007-2019 in two Portuguese hospitals. Data was obtained from clinical files. Statistical analysis was performed using SPSS version 24<sup>®</sup> and OS using Kaplan-Meier method.

**Result(s)\*** There were included 70 pts, of which 23% ( $n=16$ ) had *BRCA1/2* mutation: *BRCA1* mutation occurred in 56% ( $n=9$ ) of pts and *BRCA2* in 44% ( $n=7$ ). The Portuguese founder mutation - *BRCA2* c.156-157insAlu was found in 2 pts.

Median age of *BRCA* mutated (mut) pts was 56 years (39-71) and *BRCA* wild type (wt) was 62 years (35-78).

The most frequent histology was Serous Carcinoma, in 86% ( $n = 60$ ) of pts; most frequent stages were IIIC 46% ( $n = 32$ ) and IV 17% ( $n = 12$ ). Neoadjuvant chemotherapy (CT) was performed in 50% of pts ( $n = 35$ ) and in 37% ( $n = 26$ ) surgery was the first therapeutic approach followed by adjuvant CT. Eleven pts (16%) were treated with PARP inhibitors: 6 pts *BRCAmut* and 5 *BRCAwT*.

There was family history of cancer in 56% of *BRCAmut* and in 45% of *BRCAwT*.

mOS of *BRCAmut* was 13.81 years (CI 95% 10.36-17.26) and 5.54 years (CI 95% 4.21-6.88) to *BRCAwT*, with a significant difference between the two groups ( $X^2=4.460$ ;  $P=0.035$ ).

**Conclusion\*** Detection rate of *BRCA1/2* mut was higher than described in literature. *BRCAmut* pts showed a statistically significant longer survival, when compared with *BRCA wt* pts. Characterization of these pts at a national level would be an opportunity to obtain real data from the Portuguese population.

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### OUR EXPERIENCE IN OVARIAN CANCER 2006–2015 . STANDARDS OF QUALITY IN SURGICAL MANAGEMENT

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10.1136/ijgc-2021-ESGO.459

**Introduction/Background\*** It is important to know the survival data of patients with ovarian cancer treated in our unit, the variables associated with the prognosis and the degree of compliance with the standards in the surgical management of ovarian cancer

**Methodology** Retrospective study of patients with ovarian cancer diagnosed and treated in CHUIMI in the period between 2006-2015. We studied epidemiological variables, stage at diagnosis, type of treatment, histopathological study, follow up and current status of patients.

**Result(s)\*** The total number of patients diagnosed with ovarian cancer in the study period was 331, with a mean age of 57.84 years (range 26–85 years). 69.8% were in advanced stages at the time of diagnosis (Stage I 23.9% (79), Stage II 6.3% (21), Stage III 54.1% (179) and Stage IV 15.7% (52).

Regarding the histological type, serous was the most frequent representing 49.8% of the sample, followed by endometrioid with 16.3% and clear cells with 10.9%. We found that endometrioid, clear cells and mucinous types were more frequent in the grupo diagnosed with early stages versus the serous type that were more associated with the advanced disease.

Overall survival (OS) at 5 years is 40.8% for the complete series. 83.3% for stages I, 72.2% for stages II, 29.1% for stages III and there are no patients in stage IV who lived after 5 years. In stages III, the most frequent therapeutic approach is initial surgery in 41.1%, followed by neoadjuvant chemotherapy in 30.3%. Stage III patients receiving surgery + adjuvant chemotherapy showed an OS of 47% at 5 years (median survival 44 months) meanwhile those who received neoadjuvant chemotherapy and get the surgery in second place showed an OS of 27.1% (median survival 35 months).

When we studied the effect of tumor residue after surgery in stage III patients, the OS when the surgery was complete was 52,9% at five years, 15% if there were residual tumour, regardless of the size. Initial surgery was performed in 58% of all stages III-IV (objective > 50%). Complete cytoreduction was achieved in 51% of all stages III-IV (minimum objective > 50%, optimal > 65%)

**Conclusion\*** Our epidemiological and survival data coincide with what has been published in the literature. Having surgeons with experience in the management of peritoneal carcinomatosis will allow to increase the rate of complete cytoreductions

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### ESGO CERTIFICATION FOR ADVANCED OVARIAN CANCER SURGERY: THE EXPERIENCE OF AN ONCOLOGY CENTER TO AIM ACCREDITATION

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10.1136/ijgc-2021-ESGO.460

**Introduction/Background\*** The outcomes of advanced ovarian cancer surgery is related to the size of the largest residual